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medically assisted procreation

Working Paper 65

Canada
MEDICALLY ASSISTED PROCREATION

Working Paper 65

1992
Commission*

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* Mr. Justice Linden had just left the Commission at the time this working paper was approved, but he was involved in all the stages of its preparation. Judge Rivet was still a member of the Commission and Mr. Trevor-Deutsch was Project Co-ordinator. Madam Justice Picard and Professor Frémont joined the Commission after the document was approved.
Editor's Note

In keeping with the proposal advanced in *Equality for All: Report of the Parliamentary Committee on Equality Rights*, we have conscientiously endeavoured to draft this working paper in gender-neutral language. In doing so, we have adhered to the standards and policies set forth in *Toward Equality: The Response to the Report of the Parliamentary Committee on Equality Rights* pertaining to the drafting of laws, since the Commission's mandate is to make proposals for modernizing Canada's federal laws.
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Acknowledgements

The analysis in the following pages is based on information available in December 1990. However, the medical evaluation used as the starting-point for our study was completed in March 1990. Readers should bear these dates in mind.

We would like to thank the members of our Advisory Group of Experts on Health Law, which was officially consulted in November 1989. Needless to say, the views expressed in this paper do not necessarily reflect the positions of the group’s individual members. The following individuals were consulted; they are listed in alphabetical order by field of expertise.

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Introduction

Technological development and the advancement of science, particularly medical science, constantly raise new challenges for our political and legal institutions. Social, moral, political and economic pressures force Parliament to systematically review the law and the practices arising therefrom. The law must take account of and promote scientific development, but must also impose the restrictions dictated by certain human and social values. Medically assisted procreation is perhaps one of the best examples of the challenges posed by the development of medical science and the tensions to which they give rise for the law.

National and world interest in medically assisted procreation reflects the importance ascribed to the risks, consequences and social and legal implications of the technologies being used. Whether the goal is to develop policies on reproductive technologies or reduce legal ambiguity, society must reflect upon the choices to be made in view of existing conflicts and re-examine certain fundamental values and principles of law. Among issues of particular concern are the definition of the family; the filiation of children born as a result of medically assisted procreation; the commercialization of procreation, the human body and its products and substances; and the legal status of gametes and embryos. The potential and actual risks, both physical and psychological, raise concerns about public protection, including the protection of children born as a result of these technologies. We must therefore also consider the adequacy of the controls that apply to medically assisted procreation and the selection and storage of gametes and embryos.

However, developing a consistent national social policy on medically assisted procreation is not an easy task. The diametrically opposed views expressed in the reports of the Ontario Law Reform Commission\(^1\) and the Barreau du Québec\(^3\) on a number of fundamental aspects of the issue illustrate the problems. And the provisions in the Canadian Constitution on the distribution of powers between the federal government and the provinces do nothing to make matters easier.

The complexity and gravity of the issues raised by medically assisted procreation therefore demand careful consideration and serious discussion within Canadian society. It was

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1. Legal uncertainty can give rise to tremendous personal, family and social conflict. The American cases of Baby M, infra, note 302, Davis v. Davis, infra, note 203 and York v. Jones Institute, infra, note 204, as well as the Roe case involving orphaned embryos, infra, note 304, are good illustrations of this.
with this objective in mind that the federal government established the Royal Commission on New Reproductive Technologies. The Commission was created for a specific purpose and is not intended to replace any permanent agency. Its mandate is to inquire into and report on current and foreseeable medical and scientific developments related to new reproductive technologies and their social, ethical, health, research, legal and economic implications. The Commission is also required to recommend policies and safeguards pertaining to a number of related issues. It is to submit its report in October 1992.

The Law Reform Commission of Canada undertook its examination of medically assisted procreation in an effort to advance the public debate and to complete its trilogy of studies in the area of medical law and procreation. In May 1988, the task force set up by the Commission to review the status of the fetus recommended that the Commission inquire into the field of medically assisted procreation. In its working paper Crimes against the Foetus the Commission identified a number of issues that called for further research, among them surrogate motherhood, the need to establish national standards in view of the risk of interprovincial “procreative tourism” and to define the liability of donors who supply false information about their health status (genetic disorders, hereditary diseases, medical history) or who fail to provide pertinent information. The Commission subsequently began a comprehensive study of the various technologies currently in use, and the phenomenon of surrogate motherhood.

The serious concerns that unsettle our society were also a factor in the Commission’s decision to conduct this study. These concerns are often accompanied by demands for state

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5. P.C. 1989-2150. The Commission’s terms of reference include the following main issues:

(a) implications of new reproductive technologies for women’s reproductive health and well-being;

(b) the causes, treatment and prevention of male and female infertility;

(c) reversals of sterilization procedures, artificial insemination, in vitro fertilization, embryo transfers, prenatal screening and diagnostic techniques, genetic manipulation and therapeutic interventions to correct genetic anomalies, sex selection techniques, embryo experimentation and fetal tissue transplants;

(d) social and legal arrangements, such as surrogate child-bearing, judicial interventions during gestation and birth, and “ownership” of ova, sperm, embryos and fetal tissue;

(e) the status and rights of people using or contributing to reproductive services, such as access to procedures, “right” to parenthood, informed consent, status of genetic donors and confidentiality, and the impact of these services on all concerned parties, particularly the children; and

(f) the economic ramifications of these technologies, such as the commercial marketing of ova, sperm and embryos, the application of patent law, and the funding of research and procedures including infertility treatment.


8. Supra, note 7 at 61. Ibid. "Regulation of medical practice falls under provincial jurisdiction. In the absence of uniform, national accreditation procedures and limits of practice for institutions, the possibility of interprovincial 'procreative tourism' cannot be ignored and should be seriously examined."
intervention in the form of limits or controls justified by the scale of the costs involved, the need to impose limits on the development of medicine, the dangers of the marketing of procreation, the protection of the family unit and moral values, and the provision of safeguards against the exploitation of embryos, children, infertile couples and, especially, women.

Such demands are made by individuals and groups whose interests are sometimes at odds with the needs of those who are most directly involved (infertile or sterile individuals, physicians, scientists, and so on), and this can give rise to legal and social instability. In light of this instability and the inadequacy of other social controls, legislative intervention may be needed to define and regulate the relationships between the parties and the social groups concerned and thus to restore social equilibrium. In this area, perhaps more than any other, we must be careful not to act too swiftly. The first step is to establish whether there is in fact a need for reform; only then can the scope of the change be determined.

Moreover, these social demands involve various aspects of the law which, needless to say, are of special interest to a law reform commission: law as an instrument of social change; law as a protector of the fundamental values of society; and law as a regulatory agent.

Our work centred around certain main themes. First, we examined the appropriateness of state intervention for purposes of limiting access to medically assisted procreation technologies or of controlling the use of such technologies. We considered the "problematic" medical aspects of the technologies and the ability of existing social and professional controls to provide adequate public protection (physical and psychological risks, selection, screening and storage standards, success rates, record keeping and access to medical and genetic information). We also looked at the need to review and, where necessary, adapt the law to the specific problems connected with the donation and deposit of gametes and embryos (ownership of genetic material and donor liability), and the filiation of the children involved. Finally, we examined the legal aspects of surrogate motherhood and the commercialization of procreation in general (sale of gametes and payments to surrogates).

Our study is divided into four chapters. The first two deal with the medical and legal aspects of the various technologies. The other two cover the role of the state and the reforms proposed by the Commission.

More specifically, chapter 1 explains the process involved in achieving pregnancy, the various kinds of infertility, and the technologies most often used to treat it. We will also discuss the genetic indications that may encourage the use of some of these technologies, and the risks and consequences associated with them. Particular attention will be paid to

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9. See chap. 3 regarding existing controls on medically assisted procreation. See chap. 2 regarding the shortcomings of positive law. See also Robert L. Kadera, *Connecting Law and Society: An Introduction to Research and Theory* (Englewood Cliffs, N.J.: Prentice-Hall, 1983), regarding the role of social forces.
success rates, the various ways of determining them and their effects on a couple's ability to choose the technology best suited to their situation. Finally, chapter 1 explains the various procedures used in gamete donation and the importance of donor selection.

Chapter 2 sets out the issues raised in Canadian law by the various situations made possible by the use of medically assisted procreation, in terms both of private law and of the rights protected by the Canadian Charter of Rights and Freedoms. The main private law issues have to do with the legal status of gametes and embryos, the parentage of children born by means of new reproductive technologies, gamete donor liability, the liability of gamete and embryo banks and of physicians, the nature of surrogacy contracts, and the application of the principles of law with respect to the commercialization of the body, its products and substances. We will also study the influence of the Charter on state regulation of the use of medically assisted procreation technologies. In this connection, we will consider the question of the existence and limits of a possible constitutional right to procreate, and the influence of equality rights on the regulation of access to reproductive technologies. We will conclude with brief comments on the application of section 1 of the Charter in this context.

In order to determine the need for and the scope of possible state intervention in the area of medically assisted procreation, in chapter 3 we will deal with existing regulatory mechanisms, their scope and the role of the state.

In light of the analyses in the preceding chapters, chapter 4 will set out a series of proposals for reform designed to better define the interests at issue, to specify the rights and obligations of those involved in the use of reproductive technologies, and to create the necessary balance among the various values involved, such as respect for human dignity and respect for the individual rights and liberties of those concerned. We conclude our proposals with suggestions for putting in place a mechanism for implementing our recommendations.

Appendix A provides a description of the various measures recommended or adopted abroad — in particular Australia, the Council of Europe, Denmark, France, Germany, Norway, Spain, Switzerland, the United Kingdom, the United States — and, in Canada, at the federal level and in the common law provinces and in Quebec. Finally, appendix B provides a draft proposal indicating the structure that legislation governing the main aspects of medically assisted procreation might take.

We are aware that many of the questions raised in this study fall under provincial jurisdiction. But since they involve public health, the protection of certain fundamental values of our society, the protection of human life and bodily integrity, interprovincial and international trade, certain rights guaranteed by the Canadian Charter of Rights and Freedoms, and a clear need for uniformization, we believe that the federal government can and must play a major role as a catalyst in research and the development of a Canadian policy in the area of medically assisted procreation.

CHAPTER ONE
Medical Aspects of Medically Assisted Procreation

About 15 percent of couples seek medical assistance concerning fertility problems. Whether this is a reflection of an increasing incidence of infertility over the years is subject to debate. What is not in debate, however, is that the adoption of infants is becoming increasingly difficult and can no longer be considered the alternative to childlessness it once was. As adoption has become more difficult, the use of reproductive technologies and the medical treatment of infertility have become more widespread. In this first chapter, we present an overview of infertility and the reproductive technologies most commonly used to overcome it. These include artificial insemination (AI), in vitro fertilization (IVF) and gamete intrafallopian tube transfer (GIFT).

It is clear that some procedures are of more therapeutic value than others. Because the ability of infertile couples to choose the option most suited to them may be impaired by confusing reports of success, success rates are discussed in detail. Also explained are the procedures of gamete (sperm and egg) donation, and the importance of preventing the transmission of serious infectious and genetic diseases through adequate donor screening.

I. Achieving Pregnancy

Males produce sperm continuously throughout their reproductive lives. An ejaculation often contains more than 100 million sperm per millilitre. This is in sharp contrast to the total of 400 to 500 eggs that will be ovulated, usually one at a time, once a month, from puberty to menopause in a woman’s lifetime.

In the sexually mature woman, several immature eggs (oocytes) begin to develop each month in the ovaries. Usually only one reaches maturity. A structure called the follicle houses and nurtures the maturing egg until it reaches maturity under the cyclic influence of hormones released from the pituitary gland. During ovulation the mature egg is released from the follicle, complete with protective coverings. After ovulation the follicle continues to function, secreting hormones (progesterone and estrogen) that are necessary in preparing the uterine lining (the endometrium) for implantation.

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12. OLRC, supra, note 2 at 16, where the decrease in the number of newborns available for adoption in Ontario is discussed. In 1982, for example, 73 infants were placed for adoption, compared to 961 in 1969.
Pregnancy depends on a series of events that must be successfully synchronized. Sperm must be deposited into the vagina in adequate numbers and must have sufficient motility to travel from the vagina through the uterus and unobstructed fallopian tubes to meet the egg that has been released from the ovary.13

The egg is receptive to fertilization for approximately 24 hours, after which it dies. The average lifespan of sperm is 48 hours (although fertilization may occur up to three or more days after insemination).14 Conception is therefore possible during only a few days of the menstrual cycle.

Fertilization begins with contact between a spermatozoon and the egg, and ends with the union of maternal and paternal chromosomes (the male and female pronuclei). This occurs in the upper third of the fallopian tube. The cell resulting from the union of male and female pronuclei is referred to as a zygote, pre-embryo, or conceptus, and its genetic material is already different from that of its biological parents.

Approximately 30 hours after fertilization, cell division occurs for the first time and continues while the pre-embryo or conceptus is still traversing the fallopian tube. At about three days following fertilization, when the conceptus is at the 16-cell stage, it enters the uterus. Implantation in the uterine wall occurs between the sixth and tenth day after fertilization. The conceptus is now a hollow ball of cells referred to as the "blastocyst."15

Relatively few conceptus result in babies. It has been estimated that more than 60 percent are lost prior to 12 weeks’ gestation, and 90 percent of these losses occur before the first missed menstrual period without the knowledge of the mother.16 The clinical spontaneous abortion rate or rate of miscarriage is about 15 percent in the general population.17 Approximately 50 to 60 percent of these miscarriages are due to chromosomal abnormalities, a result of imperfect gametes or abnormal fertilization. Thus, the number of spontaneous abortions is to a large extent a natural form of protection against the continuation of an abnormal pregnancy.18

Consequently, in any given cycle of 100 ovulatory women who actually conceive, only 25 will become aware of pregnancy; of these, approximately four will have a miscarriage and about 21 go on to produce a live baby. This is why humans are said to be inefficient procreators; and, as we shall see, the fact that they are constitutes one of the most intractable problems of medically assisted procreation.

13. For a complete discussion see Keith L. Moore, Before We Are Born: Basic Embryology and Birth Defects, 3d ed. (Philadelphia: W.B. Saunders, 1989) chaps 2 and 3.
14. The process of swimming through the uterus and into the fallopian tubes induces a process called capacitation, which is the final step in the maturation of sperm. Capacitation is essential for successful fertilization.
17. Ibid. This figure varies according to age. For an explanation of spontaneous abortion, see infra at 14.
18. Moore, supra, note 15 at 34.
II. Infertility

Infertility is the involuntary, significant reduction of reproductive capacity. In North America, the generally recognized threshold of infertility is an inability to become pregnant after one year of unprotected intercourse.\(^{19}\) The World Health Organization’s standard is two years.\(^{20}\)

Although Canadian studies of infertility prevalence are scarce, it has been reported that 15 percent of couples seek medical advice for infertility.\(^{21}\) In the United States the prevalence of infertility has not changed significantly from 13.3 percent in 1965 to about 13.9 percent in 1982, excluding surgically induced sterility.\(^{22}\)

Some of the factors influencing the prevalence of infertility are:\(^{23}\) (1) trends toward childbearing later in life;\(^{24}\) (2) environmental factors, such as infection from sexually transmitted diseases, and occupational exposure; (3) medical treatments such as those used for high blood pressure, stomach ulcer and cancer, as well as non-therapeutic drugs such as narcotics, alcohol and tobacco.\(^{25}\)

A. Evaluation of the Infertile Couple

The infertile couple seeking medical help undergoes a series of procedures to determine the nature and severity of the problem. First a medical history is taken and, if necessary, counselling about timing effective intercourse is given.\(^{26}\)

The woman is tested to detect hormonal dysfunction. There may be a biopsy of the uterine lining, and a hysterosalpingogram, which is an X-ray that reveals blockages of the fallopian tubes. Laparoscopy, which is the introduction of an endoscope into the abdomen, may be used to inspect the outer surfaces of the uterus, fallopian tubes and

24. For statistics on Canadian trends towards childbearing later in life see A. Romanow, Fertility in Canada: From Baby-boom to Baby-burst (Ottawa: Statistics Canada, 1984) at 27-32. Also, the prevalence of infertility has been found to be about 10 percent higher in women over 35: see Infertility: Medical and Social Choices, supra, note 22 at 32.
surrounding structures for any abnormalities. These procedures are often painful, include slight risks of infection, and may result in the puncture of the uterus, although this last is rare. Medical precautions, such as the administration of antibiotics, are therefore taken to minimize risks.27

The man must undergo a semen analysis to evaluate the number and quality of sperm. If the semen is abnormal, blood tests may be performed to detect hormonal abnormalities.28 A post-coital test may also be used to determine if there is incompatibility between the semen and female reproductive factors. This test requires the couple to have sexual intercourse timed to coincide with ovulation; within a few hours, post-coital tests of cervical mucus are performed.29

B. Causes of and Treatments for Infertility

Infertility may be traced to one partner, both partners, or to biochemical or immunological incompatibility between partners. Most female infertility is due to: ovulation disorders, usually because of hormonal abnormality; tubal blockage as a result of infection and other disease processes; endometriosis; and other causes, including abnormalities of the vagina or cervix, and mucous incompatibilities with sperm.30

Treatments for female infertility include hormone or drug therapy, surgery, and medically assisted procreation technologies such as IVF and GIFT.

Infertility due to an ovulation disorder is treated with ovulatory stimulants, which are very successful if infertility is due only to an ovulation disorder.33 Other medical

27. For further discussion see Leon Speroff, Robert H. Glass and Nathan G. Kase, Clinical Gynecologic Endocrinology and Infertility, 4th ed. (Baltimore: Williams & Wilkins, 1989) chap. 17.

28. Serial studies of fertile men have demonstrated great variability in sperm counts of individuals over time, emphasizing that at least two or three sperm counts are often necessary before an accurate count can be assigned. See Richard F. Spark, The Infertile Male: The Clinician’s Guide to Diagnosis and Treatment (New York: Plenum, 1988) at 130. See also David W. Keller, Ronald C. Strickler and James C. Warren, Clinical Infertility (Norwalk, Conn.: Appleton-Century-Crofts, 1984) at 100.

29. Hammond, supra, note 19 at 826.


31. The tissue that lines the uterus, the endometrium, grows abnormally outside the uterus in endometriosis. Approximately 10 percent of pre-menopausal women have endometriosis, and about 30 percent of affected women are infertile. Precisely how this form of infertility occurs is unknown. See Ron Muse, “Clinical Manifestations and Classification of Endometriosis” (1988) 31:4 Clin. Obstet. Gynecol. 813.


treatments include drugs to treat endometriosis, infection, and immune incompatibilities. For fallopian tube blockage, surgery may be used. When other infertility treatments are unsuccessful, artificially assisted procreation may be employed, but as a last resort.

Male infertility typically results from decreased numbers or an absence of sperm in the semen, abnormal motility and structural abnormalities, all of which prevent normal fertilization of the egg. Precise causes of male infertility are often undetectable, but varicocele (varicose veins of the testes) or infection may play a role. The absence of sperm (azoospermia) may be caused by impaired production of sperm or blockage of passageways. Although greatly reduced numbers of sperm (oligospermia) reduce fertility, there is still controversy as to the number of sperm necessary for normal reproductive functioning.

When sperm counts fall below five million, fertility is significantly reduced. Therefore, couples unwilling to wait the several years often necessary to achieve "natural" pregnancy may seek treatment for male factor infertility. These treatments include hormonal therapy and such laboratory techniques as the "swim-up" procedure that aim to improve the concentration of normal sperm available for fertilization. However, the success of these procedures in conjunction with the use of artificial insemination is less than 20 percent.

with the use of menotropin (Pergonal®). The risk of provoking "hyperstimulation syndrome" is greater with the use of Pergonal® than with chorionepine. It is a syndrome of varying severity, where a mildly affected woman (15 percent) will suffer enlarged ovaries and a severely affected woman (one percent) will suffer fluid shifts in the body which may be severe enough to cause death. Her ovaries may enlarge to the extent that rupture is a possibility, requiring removal of the ovaries and, very rarely, death has been reported. Therefore close medical observation is required with the use of some ovulatory stimulants. See also Richard Borenstein et al., "Severe Ovarian Hyperstimulation Syndrome: A Reevaluated Therapeutic Approach" (1989) 51:5 Fertil. Steril., 791.


36. Ibid. Although a sperm count less than 10 million is considered unacceptably low, some pregnancies have occurred in partners of men with sperm counts below 5 million (ibid. at 129).

37. Erik Bostofte, Jorgen Sperup and Heinrich Recke, "Hammed semen Quality Classification and Pregnancies Obtained during a Twenty-Year Follow-up Period" (1981) 36:1 Fertil. Steril. 84.

38. The purpose of the "swim-up" procedure is to obtain the highest concentration of mobile sperm possible by collecting sperm that have been placed in a container and have swum to the top of a special medium. Nancy J. Alexander and Steven Ackerman, "Therapeutic Insemination" (1987) 14:4 Obstet. Gynecol. Clin. N. Am. 905 at 909.

39. Ibid. at 911. A compilation of the literature reporting 812 cases of male factor infertility treated by means of AIH sperm yielded a pregnancy rate of 18 percent.
In theory, one might expect that IVF could be useful in the treatment of male factor infertility. Once the egg is placed directly in a container with the partner's sperm, the normal sperm, even if there are relatively few, should be able to fertilize the egg. This would provide the couple with a child genetically related to both parents. But the ability of the sperm to fertilize the egg appears to be only half as successful as in cases of IVF with non-male factors.\(^{40}\) Nevertheless, there are reports that find IVF for male factor infertility as successful as IVF for other reasons.\(^{41}\) In any event, artificial insemination by donor (AID) is considered a leading remedy for both the infertile and sterile male because it is less costly, less invasive, and statistically much more successful than IVF.\(^{42}\)

III. Genetic Indications for Medically Assisted Procreation

Individuals who are not infertile but risk transmitting serious genetic diseases\(^{43}\) or abnormalities may be considered "reproductively disabled" and are therefore candidates for medically assisted procreation. These individuals may require a donation of one or both gametes if the disease the gene of which they carry is not detectable through prenatal diagnostic techniques or if the couple is not amenable to the therapeutic abortion of an affected fetus. There are three types of transmissible genetic disease: an autosomal recessive disease such as Tay-Sachs disease or thalassemia;\(^{44}\) an X-linked disease such as Duchenne muscular dystrophy or hemophilia;\(^{45}\) or an autosomal dominant disorder such as Huntington's disease.\(^{46}\) The presence of chromosomal (structural or numerical) abnormalities may also present significant risks for the offspring.

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40. Ibid. at 513. See also A. Apra et al., "The Role of In Vitro Fertilization in Male Infertility" (1986) 541 Ann. N.Y. Acad. Sci. 297 at 301; and Irvin Hirsch et al., "In Vitro Fertilization in Couples with Male Factor Infertility" (1986) 45:5 Fertil. Steril. 659 at 662.

41. In selected populations of male factor infertility, IVF success is comparable to other indication for IVF. See Hirsch et al., supra, note 40 at 663.

42. Alexander and Ackerman, supra, note 38 at 918ff.

43. Individually, genetic diseases are usually very rare, but when all genetic diseases are considered together, they account for more than 30% of all pediatric hospital admissions. Judith G. Hall, "Impact of Genetic Disease on Pediatric Health Care" in M.M. Kaback and L.J. Shapiro, eds, *Frontiers in Genetic Medicine: Report on the 22nd Ross Conference on Pediatric Research* (Columbus, Ohio: Ross Laboratories, 1987) 1 at 1.

44. Only when both partners carry the gene for an autosomal recessive disease is the offspring at risk. The disease occurs only if the offspring inherits both disease genes, one from each parent. The risk for disease occurrence is 25 percent for each pregnancy.

45. Also referred to as a sex-linked disease, the disease gene is located on the X chromosome and is most often passed on from an unaffected mother (who has two X chromosomes) to her son (who has one X chromosome along with a Y chromosome). Each son has a 50 percent chance of inheriting the disease, and each daughter has a 50 percent chance of inheriting the disease gene and becoming a carrier, usually without being affected herself.

46. If either parent carries the disease gene, each child will have a 50 percent chance of inheriting the disease.
IV. Medically Assisted Procreation Technologies

A. In Vitro Fertilization (IVF)

In 1878 the first attempts at in vitro fertilization were made using rabbits and guinea pigs. One hundred years after these first animal experiments, the first test-tube baby was born. Since that time more than 3,000 children have been born worldwide as a result of IVF. It is most commonly used when infertility is caused by blocked fallopian tubes, but is also used in some cases of endometriosis, and in male factor and unexplained infertility.

By definition, in vitro fertilization implies the fertilization of an egg outside the future mother's body, or "in glass." The procedure is accomplished by removal of mature eggs from the ovary, fertilization in a container, and transfer of the early-developing concepti back into the uterus.

The IVF procedure involves five steps: (1) administration of superovulation drugs; (2) removal of one or more mature eggs from the ovary; (3) fertilization of one or more of these eggs outside the body; (4) incubation of the fertilized egg(s) until they are ready for transfer; and (5) transfer of early-developing concepti into the uterus. The success of each step is contingent on the preceding one. Failure at any stage results in failure to become pregnant.

1. Superovulation

The probability of achieving pregnancy, and eventually live birth, increases when three or four embryos are simultaneously implanted in the uterus. Therefore, virtually all IVF programs throughout the world use drugs, individually or in combination, that induce the maturation and ovulation of more than one egg per cycle.

49. Ibid. at 828-29.
51. The most common drugs used are clomiphene citrate (Clomid), menotropins (Pergonal), and human chorionic gonadotropins (HCG). IVF-ET Canada, supra, note 50 at 16. Gary D. Hodgen, "Physiology of Follicular Maturation" in Howard W. Jones, Jr., In Vitro Fertilization — Norfolk (Baltimore: Williams & Wilkins, 1986) 8 at 9. See also F.T. Cameron and D.L. Healy, "Patient Management" in Carl Wood and Alan Trounson, eds., Clinical In Vitro Fertilization, 2d ed. (London: Springer-Verlag, 1989) 11 at 15. For the risks involved with the use of these drugs, see supra, note 33.
Ovulation induction is successful approximately 80 percent of the time (an unsuccessful stimulation is referred to as a "cancelled cycle"). During treatment, patients are either admitted to hospital or outpatient facilities so that frequent blood samples can be taken to evaluate hormone levels and ultrasound monitoring of developing follicles may be carried out.

Common side-effects of superovulation drugs include: hot flushes (10 percent), abdominal distention and pain (5.5 percent), breast discomfort (2 percent), visual disturbances (1.5 percent), headache (1.3 percent) and loss of hair (0.3 percent). The "hyperstimulation syndrome" occurs rarely in IVF trials. The occurrence of this syndrome is minimized by medical precautions including the emptying of all the follicles, frequent ultrasound evaluations of ovaries and the control of blood hormonal levels.

2. Egg Retrieval

Egg retrieval is performed with the aid of laparoscopy or ultrasound. In laparoscopy, the abdominal wall is punctured and the abdomen inflated with a gas mixture. This creates a space between the abdominal wall and the intra-abdominal organs. One or more tubes, including an endoscope, are then inserted into the space so that the surgeon can observe and manipulate the ovaries without having to make too large an incision. The mature eggs are then removed from the follicles. The main disadvantage of laparoscopy is that a general anesthetic is required, in addition to the use of several drugs during and after the laparoscopy to maintain the effect of the anesthetic and relieve pain. There has been concern regarding the effects of these drugs on the pregnancy, since it is known that they may enter the follicular fluid. Studies thus far, however, show no apparent effect on the outcome of the pregnancy.

Ultrasound-guided retrieval of ova is often preferable to laparoscopic techniques because it is less invasive, and because a general anesthetic is not necessary. Ultrasound employs high-energy sound waves, beamed from outside the body, to create images of internal organs and structures. The surgeon uses the ultrasonic image of the ovary to guide a needle into the follicles through either the vagina, the urethra, or the abdominal wall. Mature ova are then aspirated out of the ovary.

52. Cameron and Healy, supra, note 51 at 15.
54. Kennedy and Adashi, supra, note 33 at 838. See also Speroff, Glass and Kase, supra, note 27 at 591.
56. J. Webster, "Laparoscopic Oocyte Recovery" in Fishel and Symonds, eds, supra, note 47 at 69. See also J. Leeton, "Oocyte Pick-up" in Wood and Trounson, eds, supra, note 51, 23 at 24.
57. Leeton, supra, note 56 at 24.
59. Procedures that involve puncture of a full bladder to reach the ovary create the most discomfort, and limited relief is achieved from the use of a local anesthetic. Leeton, supra, note 56 at 23.
In terms of numbers and quality of recovered eggs, the laparoscopy and ultrasound methods are comparable. On average, four eggs are retrieved per cycle, but as many as 20 may be produced.60

Medical risks of egg retrieval include those inherent in the administration of a general anesthetic, if used. Also, there are risks of vaginal bleeding, pain, pelvic infection,61 and the inadvertent puncture of blood vessels. There has been at least one death resulting from undetected bleeding following an egg-retrieval procedure.62

3. Evaluation of Eggs, Fertilization and Embryo Transfer

Recovered eggs are examined under the microscope for quality and maturity, so as to increase the likelihood of normal fertilization. A similar evaluation and preparation will already have been carried out on the sperm sample. The sperm is then added to a dish or test-tube containing the egg.63

If fertilization is successful, concepti are incubated. This induces cell division to the four-cell stage within approximately 44 hours. The concepti are transferred into the uterus at between the four- and eight-cell stage of development, after which the woman is discharged from the hospital and is usually encouraged to resume normal activity.64

4. IVF Success Rates

The difficulty in assessing the success of in vitro fertilization is that one clinic’s criteria may not be comparable to another’s. Criteria of success include the number of “chemical,” “clinical,” or “viable” pregnancies. These may be calculated on the basis of “treatment cycles,” “egg retrievals,” “embryo transfers” or number of women treated. The many combinations of “success” criteria can be confusing. To dispel the confusion, the definitions of each possible “numerator” and “denominator” should be understood.65

61. The risk for pelvic infection, which may be severe, is reported to be about 3 percent and is reduced with the administration of antibiotics. See Robert S. Howe et al., “Pelvic Infection after Transvaginal Ultrasound-Guided Ovum Retrieval” (1988) 49:4 Fertil. Steril. 726 at 728.
63. The concentration of sperm added to the egg(s) is much greater than that which would normally meet the egg in the fallopian tubes. See A. Trounson, “Fertilization and Embryo Culture” in Wood and Trounson, eds, supra, note 51, 33 at 33-47.
64. Selb, supra, note 48 at 830-31. See also IVF-ET Canada, supra, note 50 at 17.
5. Numerator Determination

A “chemical pregnancy,” or preclinical miscarriage, is a pregnancy that is detectable by biochemical means (HCG determinations) but does not persist long enough to delay menstruation beyond 14 days. The pregnancy is not detectable clinically, no identifiable tissue is passed, and no medical action is necessary.

A clinical pregnancy is a pregnancy detectable both by biochemical means and ultrasound, maintained until at least 28 days after egg retrieval. These criteria would indicate that the conceptus has implanted and may be considered analogous to pregnancy by natural means. A pregnancy is considered “clinical” until the point of viability. If a clinical pregnancy is lost, it is considered a spontaneous abortion or miscarriage. Medical action may be required.

Finally, although the definition of viability may differ from jurisdiction to jurisdiction, a pregnancy is generally considered to be “viable” beginning 22 weeks after the last menstrual period. The fetus is then considered capable of independent existence outside the mother’s body.

The number of live births is probably the most important criterion of success, and certainly the numerator in which most couples treated are interested. It should be noted, however, that even this “bottom-line” figure can be confusing if the rate of multiple pregnancy is not clearly reported. For example, 100 live births does not necessarily mean that 100 couples will take home one baby each, since a substantial number of these live births will be births of twins, triplets, and so on. Therefore, some authors feel that reporting the proportion of deliveries relative to attempts would be the least confusing reflection of success because the chances of taking home at least one baby could be determined.

6. Denominator Determination

Couples must be made clearly aware of the variations and limits of reporting methods, so they can exercise fully informed consent to IVF. For example, an IVF clinic may indicate that it has achieved a 20-percent pregnancy success rate. As explained above, couples should first be informed what the denominator is: the hormonal treatment cycle? egg retrieval procedure? or embryo transfer (ET) procedure? Second, couples should be aware that a 20-percent pregnancy success rate per embryo transfer still does not necessarily reveal all the statistical and psychological realities of the process.

66. Page, supra, note 65 at 334.
67. Ibid.
68. Crimes against the Foetus, supra, note 7 at 43 n. 93.
Let us assume, for instance, that 200 couples enter an IVF clinic that claims a success rate of 20 percent per embryo transfer. Couples might reasonably conclude that 40 of the women are likely to become pregnant. In fact, many fewer are likely to do so because many couples will not reach the embryo transfer stage. At step one of the IVF process, the treatment cycle, some 40 of the original 200 couples will leave the program because, in general, 20 percent of IVF hormonal treatment cycles are unsuccessful.71 Of the remaining 160 couples, another 24 (15 percent) will likely drop out prior to embryo transfer owing to difficulties at the egg retrieval and fertilization stages.72

This brings about 136 of the original 200 couples to the embryo transfer stage. Applying the clinic’s stated 20-percent pregnancy rate per embryo transfer means that about 27 of the original 200 women will become pregnant, not 40 as one might reasonably expect. Statistically this translates into only a 13-percent pregnancy rate per treatment cycle (if this is the denominator chosen). Clearly, a “success rate of 20 percent,” taken in isolation, is meaningless. The numerator and denominator must be known and must be the same in order for clinics to be compared with each other. Standardized methods of reporting success rates would greatly enhance couples’ ability to make informed choices.

7. International Results

Table 1 (see infra at 35) gives recent statistics of national registries from Australia73 (13 centres), the United States74 (96 centres), and the United Kingdom75 (42 centres). Because different reporting methods are used, it is difficult to establish a common criterion of “success.” However, a crude measure of delivery rate (clinical pregnancies minus miscarriages and ectopic pregnancies) per hormonal treatment cycle may be instructive. Current data yield success rates of between six and nine percent. However, this is an over-estimation of the “take-home baby” rate because stillbirths and early deaths of newborns are not considered.

Further, there is great variation in reporting methods among clinics. For example, in the United Kingdom, the Interim Licensing Authority (ILA) (formerly called Voluntary Licensing Authority) classifies data according to whether the centre is small, medium or large. The six large centres average a clinical pregnancy rate of 14.3, and 10 small centres

71. See supra, note 52 and accompanying text.
72. Between 10 and 20 percent of patients who undergo egg retrieval do not reach the embryo transfer stage of IVF. H.W. Jones, Jr., and P.A.W. Rogers, “Results from In Vitro Fertilization” in Wood and Trounson, eds, supra, note 51, 51 at 57.
average 9.2 percent. In France there is also a disparity among clinics. As of April 1987, 17,000 oocyte recoveries were reported nationwide, resulting in 1,340 deliveries (7.8 percent). Although the number of pregnancies per embryo transfer generally was 14 percent, some clinics reported rates as high as 35 percent. Individually published reports offer better success rates than those using compiled data.

8. Canadian Results

IVF/ET programs have been in operation in Canada since 1982; currently there are 13 active programs. A recent report evaluating data from 11 Canadian centres between 1982 and 1988 showed that of 5,921 treatment cycles begun for 3,277 couples, a total of 460 live children were delivered. This translates into a rate of 7.9 percent live births per treatment cycle and 14.3 percent per couple treated. Since multiple pregnancy rates were not reported, exact delivery rates cannot be calculated. Assuming, however, rates similar to those in other reports in the literature, it can be presumed that the take-home baby rate was about 20 percent less than stated success rates.

Reports from individual clinics in Canada are scarce. One recent report from the University Hospital and the University of Western Ontario summarized rates and outcomes of pregnancies from February 1, 1984 to December 31, 1987 at that centre. A clinical pregnancy rate of 12.3 percent per treatment cycle with a take-home baby rate per treatment cycle of 6.4 percent was reported. The authors consider this to be an underestimation of current success rates, given recent improvements in technology.

A confidential voluntary national registry of pregnancies achieved by IVF or GIFT was begun at the Toronto East General Hospital in 1987. This registry is incomplete because several centres have not submitted results. More recently, the Ontario Medical Association has proposed guidelines (the first in Canada) to ensure the quality of IVF

76. Ibid. at 20.
80. See IVF-ET Canada, supra, note 50 at 15. Two from among the few reports are Patrick J. Tapping et al., "Initial Experience with In Vitro and Embryo Transfer at the University of Calgary/Boothill Hospital" (1985) 2:2 J. In Vitro Fert. Embryo Transfer 112; and Jacques-E. Rioux et al., "Center for In Vitro Fertilization, Québec, Canada" (1984) 1:1 J. In Vitro Fert. Embryo Transfer 89.
81. Yuzpe et al., supra, note 65 at 167.
82. Ibid. at 169-70. The rate of 6.4 percent was established on the basis of delivery and stillbirth rates.
84. Personal communication, J. Tolentino and P. Phillips at the LIFE Program, Toronto.
services, including recommendations that a provincial registry be established and operated by the Ministry of Health. The registry would include details of parentage, success or failure rates, and pregnancy outcome. The guidelines recommend that this registry be confidential and available for peer review.85

9. Other Outcomes of IVF

Although the birth of a child is the first goal, IVF may have other outcomes: spontaneous abortion,86 perinatal mortality and morbidity,87 multiple88 and ectopic89 pregnancies, and Cesarean sections.90 Rates for all of these are substantially higher in IVF pregnancies than in the general population.

(a) Multiple Pregnancies and Perinatal Risks

One in four women who have had successful IVF or GIFT treatment is delivered of twins or higher-order multiple births.91 This is a much higher rate than is seen in the

86. According to the report, miscarriage rates are approximately 25 percent compared to 15 percent for the general population. See table 1, infra at 35.
87. Perinatal mortality is defined as the number of fetal deaths (stillbirths) and neonatal (newborn) deaths, from viability until 28 days after birth. Judith S. Maung and Shira Kramer, Epidemiology -- An Introductory Text, 2nd ed. (Philadelphia: W.B. Saunders, 1985) at 92-93, 104-06. Perinatal morbidity refers to illnesses or defects of live-born infants during the same time span. See discussion in text at note 93, infra.
89. An ectopic pregnancy is a pregnancy that occurs outside the uterus, usually in the fallopian tube, and is usually caused by a blockage of the fallopian tube that prevents the concept from entering the uterus. The pregnancy in the fallopian tube may lead to a potentially life-threatening emergency that requires surgical removal of the affected tube. The ectopic pregnancy rate for IVF ranges between 4.5 and 7.5 percent compared to 1.5 percent in the general population. See table 1, infra at 35 for international rates.
90. Cohen, Mayaux and Guilhard-Moscato, supra, note 88 at 3; and Yuzpe et al., supra, note 65 at 170. Cesarean section rates are 47 and 72 percent for single and multiple pregnancies respectively, compared to general population statistics of 13 and 44 percent.
91. An international analysis of results of IVF and GIFT demonstrates a rate of multiple pregnancy of approximately 24 percent. See tables 1 and 2, infra at 35, 36. These figures represent national averages. It is well known, however, that individual unit rates will vary. For example, Yuzpe et al., supra, note 65, report a multiple pregnancy rate of 16 percent and Prydzin et al., supra, note 78 at 552 and 554 report a multiple pregnancy rate of 12 percent.
general population, where the rate ranges between 1 in 80 to 95 births, although this is
difficult to define precisely in the general population.92

A recent report from Australia concludes that pregnancies conceived through IVF
and GIFT should be considered high-risk procedures because the perinatal mortality rate
is three times that of the general population, and there is a high percentage of low-birth-
weight infants. There are several contributing factors, such as the age of the mother and
the infertility treatment itself. That said, 50 percent of the premature births and 70 percent
of the low-birth-weight infants were associated with multiple pregnancies.93 Of 460 live
births resulting from IVF in Canada between 1982 and 1987, 21 died soon after birth
(neonatal death). The majority of these were a result of premature birth, often associated
with multiple pregnancies.94

Perinatal mortality is consistently higher in twins than in singleton (single baby)
deliveries. In North America there are 14 to 16 deaths per 1,000 births for singletons.
But for twins, perinatal mortality is reported to be four to seven times higher.95 Thus,
twin pregnancies, which represent only one percent of all births, account for 10 percent
of all premature deliveries and 25 percent of pre-term deaths.96 Associated with
prematurity are breathing difficulties,97 a predisposition to hemorrhages in and around
the brain98 and infection.99

92. The actual rate of multiple pregnancies can be difficult to define because reporting methods vary interna-
tionally. See B.J. Botting, I. MacDonald Davies and A.J. MacFarlane. "Recent Trends in the Incidence
the analysis is the number of fetuses per pregnancy is the "vanishing twin" phenomenon, which
describes the spontaneous reduction rate of twin pregnancies resulting in the loss of at least one of
the fetuses. Early pregnancy ultrasound studies have demonstrated spontaneous reduction in
fetuses to be similar to or higher than the spontaneous abortion rate in singleton pregnancies.
Therefore, if IVF units report multiple pregnancy rates detected early in pregnancy, the rates will be
greater than those reported at delivery time. See the discussion in Katharine D. Wentworth and Stanley A. Gall. "Incidence, Mortality and Mortality,

93. Douglas M. Saunders and Paul Lancaster, "The Wider Perinatal Significance of the Australian In Vitro
Fertilization Data Collection Program" (1989) 6:2 Am. J. Perinatol. 252 at 252-53. See also Australian
IVF, supra, note 73 at 433. Of 1138 live-born infants, 438 (38.5 percent) were from multiple births. See
also Stanley, supra, note 69 at 425.


92 at 3, and Botting, MacDonald Davies and MacFarlane, supra, note 92 at 945.

96. See Polin and Frangipane, supra, note 95 at 650. See also Wentworth and Gall, supra, note 92 at 4-5.

97. Respiratory Distress Syndrome (RDS) occurs most often in premature infants who are deficient in surfactant,
a substance that promotes normal expansion of the lungs. This syndrome is especially prevalent (48 percent)
in twins between 25 weeks' gestation and 32 weeks' gestation (between 13 and 8 weeks premature). See
Wentworth and Gall, supra, note 92 at 5.

98. Twins delivered between 25 and 32 weeks' gestation have at least a 20 percent chance of hemorrhaging
into the ventricles of the brain. Ibid.

99. In infants weighing less than 2,500 grams the risk for Group B Streptococcal Disease was found to be
five times greater in twins than in singletons. See Kristine McCallough, "Neonatal Problems in Twins"
In addition to being prone to low birth weight because of prematurity, twins are also at increased risk for growth retardation.100 Distress during labour, reduced oxygen (asphyxia), and ultimately stillbirth are all risks found in association with growth retardation.101

An international collaborative IVF study from 55 centres found the rate of birth defects to be only slightly higher for multiple births than for single births (3.6 compared to 2.5 percent).102 However, studies of the long-term consequences of twin birth reveal a higher rate of learning disability, motor skill deficiency, speech problems, and delayed physical growth.103 Risks to the mother during and following pregnancy include increased risk of pregnancy-induced high blood pressure, anemia, Cesarean birth and excessive bleeding after delivery.104

Approximately 15 percent of all multiple births associated with IVF are triplet or higher-order multiple pregnancies. Risks for both mother and fetus are greatly increased where there are more than two fetuses. For example, the perinatal mortality rate is approximately three times greater for triplets than for twins.105

(b) Lowering the Multiple Pregnancy Rate

Louise Brown was the first test-tube baby. Her birth in 1978 was the result of the in vitro fertilization of a single egg, retrieved during a natural cycle, and the transfer of the single conceptus back into the womb. In those early days, relatively primitive methods of predicting maturation and ovulation of the egg were imprecise. Retrieval of the ovum required the surgical team to be available around the clock, and even then the rate of successful retrieval was less than 50 percent. Failure at the fertilization and implantation stages compounded the problem of low success rates. In order to improve the odds, superovulation techniques were developed. They resulted in the capacity to induce several mature eggs per cycle, and better control of the timing of ovulation. Consequently, higher pregnancy rates were achieved.106 However, as indicated above, the probability of

100. Both terms, intra-uterine growth retardation (IUGR) and small for gestational age (SGA) describe fetuses/infants who are substantially smaller than their gestational age would indicate when compared to general population statistics of singleton fetuses. For example, a baby born after 35 weeks' gestation the size of an average 32-week baby would be considered growth retarded, and therefore at higher risk than one born at a size appropriate to its gestational period. See Polin and Friginalpae, supra, note 95 at 657. See also Richard Bronsteen, Gregory Goyert and Sidney Bottoms, "Classification of Twins and Neonatal Morbidity" (1989) 74:1 Obstet. Gynecol. 98 at 100.
101. Polin and Friginalpae, supra, note 95 at 657, and Bronsteen, Goyert and Bottoms, supra, note 100 at 100.
102. See Cohen, Mayaux and Gharard-Mescate, supra, note 88 at 3.
103. Wenstrom and Gall, supra, note 92 at 5, and Polin and Friginalpae, supra, note 95 at 630-51.
104. Pritchard, MacDonald and Gart, supra, note 88 at 503.
105. Botting, MacDonald Davies and MacFarlane, supra, note 92 at 946.
multiple pregnancies and their attendant risks also increased. How, then, do we balance the maximum benefit derived from multiple transfers while minimizing the risk of multiple pregnancy? In other words, how many concepti should be transferred into the womb per cycle?

This question raises a number of controversial issues, in particular the fact that there is no agreement as to an acceptable rate of multiple pregnancies. Also, the relationship between the number of concepti transferred and the resulting number of multiple pregnancies is unclear, partly because of methods of reporting in the literature. One report, however, demonstrated that when four, five, six and seven embryos were transferred, the chances of delivering at least one baby were 18, 17, 18 and 18 percent, and the chances of multiple pregnancy were 16.7, 31.6, 50 and 50 percent respectively. In other words, although the chance of delivering a baby was not greater with more than four embryos, the chance of multiple pregnancy increased substantially. Further, the same report demonstrated that the transfer of three embryos resulted in a birth rate of 12 percent, but in a multiple pregnancy rate of 21 percent. A similar non-linear trend is seen in other reports (see table 3, infra at 36), demonstrating that pregnancy rate and multiple pregnancy rate are not determined solely by the number of embryos transferred.

Successful pregnancy is determined by the receptivity of the uterus to implantation and the “quality” of the embryos transferred. At present, both uterine receptivity and embryo quality are difficult to quantify clinically. Further research is essential in this area to improve overall success rates in artificial reproduction while reducing the number of embryos that need to be transferred. In the meantime, the most commonly accepted method of reducing the risk of multiple pregnancy is to limit the number of embryos transferred per treatment.

The limitation of embryo transfer has received international attention in the last few years. For example, national professional governing bodies in the U.K. and in Australia have issued guidelines limiting to no more than three or (in extreme cases) four the number of embryos that may be transferred at any one time. Not all practitioners have endorsed these guidelines. One British fertility team has resisted limitations, advocating a flexible approach for the number of embryos transferred in IVF or the number of eggs transferred.

109. Ibid.
when using the GIFT procedure. Professor Craft and his colleagues argued that limiting the number of eggs unfairly limited the chances of pregnancy in some cases; they argued that clinical judgment should prevail over strict enforcement, as is the case with most other medical procedures. For example, the chance of pregnancy and multiple pregnancy declines with advancing age, perhaps warranting higher transfer numbers to achieve pregnancy. On the other hand, previous pregnancy increases the chance of multiple pregnancy. Therefore, individual risk for multiple pregnancy differs depending on such criteria as age and previous pregnancy. In addition, large variations in multiple pregnancy rates have been observed among clinics, even where the number of embryos transferred per cycle was the same.

In Canada, individual clinics have policies regarding the numbers of embryos that may be transferred. Although most agree that the transfer of three or four embryos minimizes the chances of multiple pregnancy, freezing facilities are not always available for storing surplus embryos. Where adequate freezing facilities are available, surplus embryos may be frozen for future use by the couple. If the embryos are not eventually used, they may be donated, used for research or destroyed, depending on the wishes of the couple.

Where freezing facilities are not available, the transfer of more than four embryos to the uterus may result in a higher-order pregnancy. If more than three embryos implant, the couple may be offered the option of selective fetal reduction. Although the procedure has prompted ethical debate that both parallels and differs from the abortion issue, it is generally agreed there is greater risk to all of the fetuses and to maternal health if the pregnancy continues intact than if the number of fetuses is reduced.


115. Ian Craft et al., "Analysis of IVF GIFT Procedures — The Case for a Flexible Approach to Treatment" (1988) 1:8594 Lancet 1094; Professor Craft's observation that multiple pregnancy occurs less frequently with increasing age is consistent with other reports. For example, Corson et al. state that the chance of multiple pregnancy declines by 9 percent per year after the age of 30. See Stephen L. Corson et al., "Outcome in 242 In Vitro Fertilization-Embryo Replacement or Gamete Intralupian Transfer-Induced Pregnancy" (1989) 51:4 Fertil. Steril. 644 at 645. For a discussion of the effects of aging on IVF success, see also Santiago Pizzini and Jairo E. Garcia, "Effect of Maternal Age and Number of In Vitro Fertilization Procedures on Pregnancy Outcome" (1989) 52:2 Fertil. Steril. 270.

116. A 12 percent multiple pregnancy rate was observed in an IVF program in France, compared to 25.4 percent observed in a program in Belgium, both units adhering to a policy that allows a maximum of three embryos to be transferred per attempt. See Friedman et al., supra, note 78 at 553. See also P. Barlow et al., "Early Pregnancy Loss and Obstetric Risk after In-Vitro Fertilization and Embryo Replacement" (1986) 3:5 Human Reprod 671 at 675.

117. For discussion of frozen embryos, see infra at 29; see also A. Trounson, "Embryo Cryopreservation" in Wood and Trounson, eds., supra, note 51, 127 at 138-39.

118. Selective fetocide is referred to in the literature as selective reduction of fetuses, selective abortion, or selective birth. The procedure eliminates one or more fetuses in the pregnancy.

The risks associated with the reduction techniques include spontaneous abortion of the pregnancy, failure to destroy the fetus, risk of infection and, rarely, a clotting disorder that may threaten the remaining fetuses or the mother herself.\textsuperscript{120} The emotional cost of selective fetal reduction to the mother, father, and existing children is unknown; therefore, counselling should perhaps be considered prior to or following the procedure or both.

Clearly, the problem of multiple pregnancies is one that requires more research. However, a promising experimental IVF procedure suggests it may be possible to reduce the multiple pregnancy rate to that of the general population. In one small trial, a 22.5-percent clinical pregnancy rate with an ongoing pregnancy rate of 17.5 percent per cycle was obtained using an unstimulated cycle;\textsuperscript{121} this is as good as or better than results using superovulation techniques. By not using ovulatory stimulation drugs, the procedure eliminates risks associated with drugs.\textsuperscript{122} Moreover, a better quality egg and optimum uterine receptivity are more likely to result from the natural cycle. Because the procedure involves minimal invasiveness, the possibility of more frequent, repeated attempts may provide a greater overall chance of pregnancy. Non-medical advantages include lower cost\textsuperscript{123} and the elimination of ethical and legal issues regarding surplus embryos. It is too early, however, to tell whether these results will be found to be repeatable in other clinics. The main disadvantage of this procedure is failure to retrieve an egg in 10 to 30 percent of cases, and cancellation of cycles because of imprecise hormonal determinations.\textsuperscript{124}

\textit{(c) Birth Defects}

It is difficult to determine if rates of birth defects in infants conceived by means of IVF are higher than would be expected among infants conceived by natural means. Extraneous factors that could lead to a greater incidence of birth defects following IVF may include an older age group (risking chromosomal abnormalities and other hereditary defects), an increased number of multiple births, the underlying causes of infertility, and various clinical procedures including manipulation of gametes and embryos. Among 1,694

\textsuperscript{120} Ronald J. Wapner et al., "Selective Reduction of Multifetal Pregnanacies" (1990) 335:8681 Lancet 90 at 91. For a complete discussion of the history of the procedure and risks, see Fay O. Redwine and Patricia M. Hays, "Selective Birth" (1986) 10:1 Seminars Perinat. 71 at 75-76.

\textsuperscript{121} Fertility drugs are not used in the unstimulated cycle. The egg naturally matures and is retrieved, fertilized, and transferred, thereby eliminating concern regarding multiple pregnancy and surplus embryos. Hervé Foulot et al., "In Vitro Fertilization without Ovarian Stimulation: A Simplified Protocol Applied in 80 Cycles" (1989) 52:4 Fertil. Steril. 617 at 617-21. See also Jairo Garcia, "Return to the Natural Cycle for In Vitro Fertilization (Alleluia! Alleluia!"") (1989) 6:2 J. In Vitro Fert. Embryo Transfer 67 at 67-68.

\textsuperscript{122} For a discussion of risks associated with ovulatory stimulation drugs, see Kennedy and Adashi, supra, note 33; see also supra at 8-9 and 12, and Navot et al., supra, note 55.

\textsuperscript{123} The cost of the procedure is estimated to be reduced from US$6,000 to US$1,000 per attempt. See Garcia, supra, note 121 at 68.

\textsuperscript{124} Personal communication, Dr. E. Hughes, McMaster Unversity, Hamilton, Ontario.
IVF births in Australia and New Zealand between 1979 and 1986, major congenital malformations, including chromosomal abnormalities, were reported at 2.2 percent (compared to 1.5 percent in the rest of the population).\(^{125}\)

It has been suggested that the higher rate of abnormalities may have reflected observer bias, in that infants conceived by means of IVF might have been more carefully evaluated for abnormalities than children conceived naturally.\(^{126}\) Therefore, it is suggested that large, systematic studies be carried out to determine if infants conceived using IVF are in fact at greater risk for malformation.

At the time this working paper was being prepared, Canadian congenital abnormality rates for infants conceived by IVF were not available.\(^{127}\) It has been recommended that registries include this information when reporting outcomes of pregnancies conceived by artificial procreation.\(^{128}\)

\((d)\) Psychological Impact

The psychological impact of IVF procedures on couples is not apparent from a simple description of techniques and outcomes. One author who interviewed 20 Canadian women who underwent IVF found that for most of them the procedure was extremely stressful, both physically and emotionally. A profound fear of "cancellation" at each of the steps leading to embryo transfer prevailed. They also experienced "intense psychological conflict" between hopefulness and realism regarding their chances of achieving pregnancy.\(^{129}\)

The stresses of infertility and IVF are such that, in Australia, legislation requires counselling both prior to and following IVF.\(^{130}\) Indeed, Ontario Medical Association guidelines for IVF state that counselling should be available to all couples.\(^{131}\)

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\(^{125}\) Paul A.L. Lancaster, "Congenital Malformations after In Vitro Fertilization" (1987) 14:3752 Lancet 1392 (letter). Two types of birth defects, spina bifida (incomplete closure of the spine) and a serious heart abnormality (transposition of the great vessels), were significantly increased compared to the general population.

\(^{126}\) Eighty-three children conceived using IVF and 93 naturally conceived children were compared in a single-blind study. No statistically significant increase of abnormalities was seen. Although reassuring about large increases in abnormalities, the authors caution that the sample size was not significant enough to enable detection of small or moderate increases. See Norma C. Morin et al., "Congenital Malformations and Psychosocial Development in Children Conceived by In Vitro Fertilization" (1989) 115:2 J. Pediatr. 222 at 226.

\(^{127}\) Brown, supra, note 79 at 31.

\(^{128}\) See supra, note 75 at 34. See also Ontario Medical Association, supra, note 85.


\(^{130}\) Paul Bravender-Coyle, "In Vitro Fertilization and the Law in Australia" (1986) 6:3 Health L. Cat. 61 at 64.

\(^{131}\) Ontario Medical Association, supra, note 85 at 28.
B. Gamete Intrafallopian Transfer (GIFT)

In gamete intrafallopian transfer (GIFT), eggs and sperm are transferred, unfertilized, directly into the fallopian tubes. Superovulation is practised because increased numbers of eggs improve the chance of successful pregnancy. With the use of a laparoscope, mature eggs are aspirated from the follicles, mixed with sperm, and placed in a syringe. The mixture is then transferred back deep into the fallopian tubes, allowing fertilization to occur naturally. The entire procedure takes approximately 35 to 60 minutes.

This procedure is not indicated for those candidates with fallopian tube disease because of the risk of ectopic pregnancy. Therefore, although indications for GIFT overlap those for IVF (unexplained infertility, endometriosis, male factor, cervical or immunologic reasons), they exclude the largest group of candidates for IVF, those with tubal disease.

The primary advantage of GIFT is that the requirement for laboratory facilities is minimized. The major drawback is that a general anesthetic is necessary because of the laparoscopy. Thus, the invasiveness of the procedure is greater than with IVF as practiced in most centres. This may be a temporary problem, since some reports have shown that egg retrieval and transfer is possible using vaginal ultrasound-guided methods. As technical mastery of this procedure becomes more widespread, the use of laparoscopy for GIFT should decline.

1. Success Rates

An international collaborative study of the first 800 cases of GIFT procedure, using a common GIFT protocol per egg retrieval, yielded a 34.4-percent clinical pregnancy rate and a delivery rate of 25 percent. National reports from the U.S., U.K. and Australia demonstrated rates of clinical pregnancy ranging from 21 to 25 percent per transfer cycle, which is higher than the 17- to 21-percent rate seen in IVF using a similar criterion of pregnancy. Miscarriage rates were not provided in the U.K. data, so an ongoing pregnancy rate could not be calculated. However, an 18-percent delivery rate per egg retrieval was reported in the U.S. and Australian reports. The occurrence of multiple pregnancy is slightly higher than with IVF, ranging from 20 to 28 percent.

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132. Siefel, supra, note 48 at 832.
135. See P.A.L. Lancaster, "Outcome of Pregnancy" in Wood and Trounson, eds., supra, note 51, 81 at 82; Asch et al., supra, note 133 at 723; and Yuzpe et al., supra, note 65 at 168.
137. Asch et al., supra, note 133 at 722-25.
138. See table 2, infra at 36.
139. See table 1, infra at 35. The clinical pregnancy rate is based on the number of embryo transfers.
140. See table 2, infra at 36 for details.
Once again, compiled national data reports demonstrate lower average success rates than those reported from individual centres. 141

2. Complications

The collaborative study of 800 cases demonstrated no immediate major complications due to the procedure or to anesthesia, nor were any cases of induced pelvic inflammatory disease reported, 142 but the usual risks of anesthesia and laparoscopy remain. Although the ectopic pregnancy rate appears to be no higher than that seen in IVF, 143 it has been reported to be as high as 30 percent when undetectable tubal disease is present. 144

C. Other Procedures

Several other variants of IVF and GIFT have been reported. These include peritoneal oocyte and sperm transfer (POST), pronuclear stage transfer (PROST), and tubal embryo stage transfer (TEST).

POST involves the transfer of oocytes and sperm through the posterior vaginal wall, by means of a needle, using the normal fluid in the abdominal cavity as a medium for transfer into the fallopian tube. PROST involves insemination of eggs in vitro and their transfer directly into the fallopian tubes. TEST is the transfer of early-stage embryos (frozen at a four-to-eight-cell stage 145) into the fallopian tubes. None of these procedures has been used on a large scale, and they have not been reported from Canadian centres.

One variant of IVF and GIFT is referred to as IVC or intravaginal culture; this has been reported from a Canadian centre. 146 Eggs and sperm are placed in a small tube and allowed to incubate for about two days in the vagina of the recipient. The tube is held in place by a vaginal diaphragm. Later, the concepti are transferred into the uterus. This technique offers the advantage of simplifying laboratory manipulations and decreasing the cost of the procedure. A preliminary random study showed there was no difference in pregnancy rates between IVC and IVF.


142. Asch et al., supra, note 133 at 724.

143. Ectopic pregnancy is reported to be between 3 and 6 percent, similar to reports of IVF but higher than the general population risk of 1.5 percent. See Asch et al., supra, note 133 at 724; and Borroto et al., supra, note 141 at 228. See also supra, note 89.

144. Jansen, supra, note 136 at 73.


D. Artificial Insemination

The first successful cases of artificial insemination, the simplest and oldest of the reproductive technologies, were first reported in the literature in the 1770s.\textsuperscript{147} It is currently estimated that in North America between 10,000 and 20,000 infants are conceived each year as a result of artificial insemination.\textsuperscript{148} As a solution to male infertility, it is simple, non-invasive and relatively inexpensive, although some new AI procedures are beginning to approach the levels of invasiveness of other reproductive procedures.\textsuperscript{149}

Artificial or therapeutic insemination may be performed using the sperm of the husband/partner (AIH) or of a sperm donor (AID). AID is most often performed after male infertility has been established and medical treatment has been unsuccessful.\textsuperscript{150} Donor sperm is also indicated in some cases where either partner is carrying a genetic disease.\textsuperscript{151}

Artificial insemination by the husband/partner is indicated in fewer than 20 percent of couples where male infertility is present. In some, bypassing the vaginal secretions by the sperm may be sufficient to allow fertilization. Other indications for AIH include anatomic abnormalities of the male, such as hypospadias, where the opening of the urethra is situated in a place other than the end of the penis, or a maternal abnormality such as a malpositioned uterus.\textsuperscript{152}

1. The AI Procedure

It is generally accepted that donor sperm should be frozen prior to use. Frozen sperm is recommended because screening can be done to reduce the risk of transmitting infectious diseases.

\begin{itemize}
\item \textsuperscript{147} For a discussion of the history of AI, see Derek J. Jones, "Artificial Procreation, Societal Reconceptions: Legal Insight from France" (1988) 36 Am. J. Comp. L. 525 at 530-33.
\item \textsuperscript{148} Barbara Eck Meening, "The Psychology of Infertility" in James Aman, ed., Infertility: Diagnosis and Management (New York: Springer-Verlag, 1986) 17 at 23. Also, a 1981 Canadian report stated that about 500 inseminations were done each month at Canadian clinics and more than 1,500 babies had been born at that time as a result of AID. See Report of the Advisory Committee on the Storage and Utilization of Human Sperm (Ottawa: Health and Welfare Canada, 1981) at x-xii [hereafter Report on Human Sperm 1981].
\item \textsuperscript{149} DIPI, or Direct Intraperitoneal Insemination, is a procedure in which sperm is deposited by means of a needle and tube, through the posterior portion of the vagina into a space containing fluid near the ovaries and the fallopian tube, thus bypassing the uterus. See P. Dellenburg et al., "Direct Intraperitoneal Insemination: New Treatment for Cervical and Unexplained Infertility" (1988) 541 Am. N.Y. Acad. Sci. 761. See also Jansen, supra, note 136 at 72.
\item \textsuperscript{150} Alexander and Ackerman, supra, note 38 at 907-08.
\item \textsuperscript{152} Keller, Strickler and Warren, supra, note 28 at 203-04.
\end{itemize}
diseases such as AIDS, availability of specimens from the same donor for repeated inseminations is increased, and possibilities are better for the matching of recipient characteristics with donors.

The storage of sperm for future use by an individual male is another reason that sperm may be frozen. For example, permanent destruction of the capacity to produce sperm can be a side-effect of radiation therapy for some cancers. Therefore, banking of sperm prior to radiation may be desirable.

Insemination is the delivery of sperm through a syringe into the vagina, the cervix, or the uterus. Improved rates of pregnancy have been achieved using an ultrasound-guided tube to deposit sperm directly into the fallopian tube, but this procedure is still experimental.

Timing is essential to the success of artificial insemination. Since sperm survives approximately 48 hours, one insemination one to two days prior to the expected time of ovulation, and another insemination 48 hours later, should provide sufficient coverage of the fertile interval. The timing of ovulation can be determined with a certain accuracy by the charting of body temperature, ultrasound measurement of follicular growth, and measurements of hormone levels.

2. Outcomes

Most studies of clinical pregnancy rates for AID employed fresh (that is, unfrozen) semen samples. These are comparable to results achieved with natural insemination: about 20 percent per cycle, approaching 95 percent by the end of six cycles. Studies have demonstrated that freezing sperm reduces its motility, longevity and fertilizing capacity by half. Cumulative pregnancy rates are approximately half those expected with fresh semen, and several more treatment cycles, on average, are necessary to achieve conception using frozen sperm. The rate of spontaneous abortions, however, is not elevated where frozen sperm is used. As with other artificial reproductive techniques, pregnancy rates are influenced by maternal age and fertility.

153. See discussion infra, note 181.
154. Alexander and Ackerman, supra, note 38 at 919.
156. Jansen, supra, note 136 at 72.
158. Aiman, supra, note 151 at 281-82.
159. The French Federation CECOS (Centres d'études et de conservation du sperme humain) has collected data on about 17,000 pregnancies achieved using frozen sperm. The success rate per cycle is about 8 percent, with a cumulative success rate of 66 percent at 12 months. See D. Le Lannou and J. Lannou, "Artificial Procreation with Frozen Donor Sperm: Experience of the French Federation CECOS" (1989) 47 Human Reprod. 757 at 759.
161. Le Lannou and Lannou, supra, note 159 at 760. See also Christopher L.R. Barratt, Mayoja Chaudhun and Ian D. Cooke, "Donor Insemination — A Look to the Future:" (1990) 52:3 Fertil. Steril. 375 at 382.
The indication for AIH will determine the success rate: AIH is the least effective fertilization technique if the problem is with the husband’s/partner’s sperm.\footnote{Spark, supra, note 28 at 336. See also supra, note 39.} However, couples treated because of male anatomical abnormality will have high success rates.

3. AI Risks

The main risks associated with AID are: infection; transmission of genetic disease; consanguinity, if the same donor is used too many times in a small centre;\footnote{If the same donor is used too many times, theoretically there is a risk that biologically related offspring, without knowledge of paternity, may meet and reproduce, risking genetic abnormality to their offspring. In fact, it has been calculated that the risk of two half-siblings (resulting from AID in a large centre) meeting and reproducing is extremely low (less than 1/1000). Alman, supra, note 151 at 284. The risk of consanguinity would depend on the size of the community served.} administration errors in matching donor with recipient; and risks associated with intra-uterine insemination.

The risk of infection increases when the semen is introduced into the cervix and the uterus. Untreated semen may contain disease-causing organisms such as the gonococcus, chlamydia or HIV. The practice of storing sperm until adequate screening can be done greatly reduces the risk of infectious-disease transmission.\footnote{See “Screening Gamete and Embryo Donations,” infra at 32.}

Women receiving intra-uterine insemination either by donor or husband may experience uterine contractions and, more rarely, low blood pressure, slowed heart rate and weakness. Medical precautions can be taken to minimize such effects.\footnote{These effects can often be reversed with the use of aspirin. See Alman, supra, note 151 at 284.}

The risk of genetic-disease transmission is reduced if donors are adequately screened, enabling the couples to be informed, prior to acceptance, of potential risks for genetic disease in their offspring. This is discussed more fully elsewhere in the text. Consanguinity risks are minimized where donors are “retired” after a number of donations. The smaller the size of the community served by a clinic, the fewer times an individual should be permitted to donate sperm.

4. Donors

The usual practice is that sperm donors are anonymous to recipients. Medical students and other university students often act as sperm donors, perhaps because of their proximity to fertility clinics. However, in some places sperm donors may be sought through advertisements in the media.\footnote{Canadian Fertility and Andrology Society, supra, note 11 at 5.} Canadian centres usually operate their own sperm banks and in some circumstances may also import sperm from such places as New York and California.\footnote{See Report on Human Sperm 1981, supra, note 148 at 13ff.}
E. Ovum and Embryo Donation

Ovum donation is becoming an increasingly popular reproductive option for some sterile individuals, for example those with premature menopause. In 1987, in fact, 17 U.S. centres reported using donated eggs for IVF/GIFT procedures as compared to only one centre in a 1985-86 report.\textsuperscript{168} Ovum donation is most often indicated where the recipient's ovaries are absent or malfunctioning, or where she is carrying a gene for a genetic disorder.\textsuperscript{169}

Embryo donation is the practice of donating an embryo that is genetically unrelated to the recipient couple. Embryos may be available for donation from those undergoing IVF where there is an excess number of embryos for their own purposes.

Embryo freezing both creates and diminishes medical and legal ethical dilemmas. Embryo freezing delays but does not eliminate the burden of disposal of embryos which, if not used, will have to be dealt with eventually. The main advantages of embryo freezing are that: the procedure allows for a limitation of the number of fresh embryos reimplanted in an IVF cycle, therefore reducing the risk of multiple pregnancy; if the first IVF cycle is not successful, it allows for future attempts in natural cycles without further egg retrieval; it allows for simplified development of embryo-donation programs.

Limited cell damage is allowable during the freeze-thaw procedure without detrimental effects on the early conceptus because at this stage all cells have the capacity to develop fully into an embryo. Therefore, a conceptus frozen at the four-cell stage may survive the thawing procedure with three cells remaining and yet retain normal developmental potential.\textsuperscript{170}

\begin{flushleft}
\textsuperscript{170} Trounson, supra, note 117 at 140. An international survey of 24 centres, completed in December 1986, showed that of 3577 frozen embryos, approximately 50 percent were suitable for replacement. A 13-percent pregnancy rate per transfer and a 26-percent spontaneous abortion rate were demonstrated. The delivery rate was unclear, since singletons were not differentiated from multiple pregnancies. André C. Van Steirteghem and Elisurie Van Den Abbeele, “Survey on Cryopreservation” (1988) 541 Ann. N.Y. Acad. Sci. 571. For further information on success rates for pregnancies achieved through the use of frozen embryos, see Jacques Testart, “Results of In Vitro Fertilization with Embryo Cryopreservation and a Recommendation for Uniform Reporting” (1988) 49:1 Fertil. Steril. 156.
\end{flushleft}
1. The Ovum Donation Procedure

General egg retrieval procedures have been described elsewhere in this chapter. Methods of egg retrieval by lavage have been described in the literature, although they are not used in Canada. This involves the donor undergoing superovulation, with or without insemination. Eggs or concepti are flushed from the uterus with a solution. If the recipient has normal ovulatory cycles, the cycles of the donor and recipient must be synchronized to ensure that the uterine lining of the recipient is ready for implantation. If the recipient does not have normal cycles, as in premature menopause, hormones are administered to mimic a normal cycle. At the time of transfer of the concepti or gametes, the recipient’s hormonal levels must be sufficient to allow implantation, and must be maintained until the placenta of the developing embryo takes over the task of producing the hormones that maintain pregnancy, approximately 8 to 12 weeks later.

2. Ovum Donors

The potential pool of ovum donors consists of: women undergoing egg retrieval for their own reproductive purposes; women undergoing sterilization; volunteer, anonymous donors; and known donors (friends or relatives recruited by the recipient).

The process of superovulation and egg retrieval carries with it a small medical risk. Whether this risk is reasonable when donation occurs for purely altruistic reasons is still the subject of debate. Those undergoing the procedure for their own reproductive purposes, where the number of eggs obtained is in excess of individual needs, are probably the most suitable candidates for donation. The availability of eggs through this source becomes limited, however, where freezing facilities are available, because couples undergoing egg retrieval are likely to choose to have excess eggs fertilized, and the concepti frozen for their own future use.

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171. Supra at 12-13.


173. Rogers et al., supra, note 172 at 146-51.

174. Ibid. at 145-46.

175. Freezing of eggs is not an established practice at this time, although four children have been born worldwide with the use of frozen-thawed eggs. The human egg is especially vulnerable to damage during the freeze-thaw process. The risk of abnormality during subsequent cell division, resulting in an abnormal number of chromosomes, also, the protective coverings surrounding the egg may be damaged, allowing more than one sperm to fertilize the egg. More study is needed in this area to determine if there is indeed a future in the freezing of human eggs. Tronson, supra, note 177 at 138-39. See also Christopher Chen, “Pregnancies after Human Oocyte Cryopreservation” (1988) 541 Ann. N.Y. Acad. Sci. 541 at 547.
Known donors are used frequently.\textsuperscript{176} The literature suggests that caution should be exercised in that the future of the psychological relationship between the child, the birth mother and the egg donor is an important consideration.\textsuperscript{177} Further, if egg donation is employed to avoid transmission of genetic disorders, family members may not be appropriate donors. To avoid a substantial risk of genetic abnormality due to consanguinity, the husband’s sister is not a suitable candidate except when donor sperm is used.

Finally, women undergoing sterilization procedures are suitable donors.\textsuperscript{178} In these cases, individuals have already decided to undergo the invasive procedure of sterilization; therefore the only added risk is that of superovulation.

F. Surrogacy

In the context of this working paper, a surrogate is a woman who agrees to gestate a pregnancy with the intention of surrendering the newborn infant to the “social” or “contracting” parents. Leaving aside the important ethical and legal issues to which this phenomenon gives rise, there are three general medical circumstances in which surrogacy may be indicated:\textsuperscript{179} (1) absence or significant abnormality of the uterus (where surrogacy would be the only possible way of producing a child); (2) cases in which there is environmental risk to the developing fetus (such as when the mother, for her own health, must on a continuing basis take medications that may be harmful to a developing fetus); or (3) when pregnancy poses a substantial threat to maternal health, as in the case of severe heart disease. The latter two situations suggest a safer environment for pregnancy, but are not considered absolute indications.

There are several possible combinations of parentage between the surrogate and future parents. The egg may originate from the surrogate or the contracting woman or it may be donated by a third party. The sperm may be the contracting father’s or it may be donated sperm. In all, six combinations of biological parentage are possible.

Methods of fertilization nearly as numerous can be classified under two categories: (1) in vitro fertilization; and (2) in vivo fertilization, including GIFT, artificial insemination, and natural insemination. In vitro fertilization and GIFT are more likely to be used in cases where the egg is not contributed by the surrogate.


\textsuperscript{177} See discussion ILA, supra, note 75 at 15. After a multidisciplinary meeting about egg donation, it was decided that, like sperm donors, egg donors should remain anonymous (Authority’s Guidelines 13(i), ibid. at 47).

\textsuperscript{178} Rogers \textit{et al.}, supra, note 172 at 146. An added advantage of this group as donors is that their fertility has in most cases already been demonstrated.

\textsuperscript{179} For a discussion of pregnancy that poses substantial risk to the mother or the fetus, see Fitchard, MacDonald and Grant, supra, note 88 at 494, 592, 608 and 802.
Surrogate embryo transfer (SET), in which the egg of the donor is fertilized in vivo by artificial insemination, collected by lavage and transferred to the gestational mother, is sometimes included under the category of surrogacy, but in this chapter it is discussed under the heading of ovum and embryo donation.\(^\text{180}\)

V. Screening Gamete and Embryo Donations

Sperm donation is now a well-accepted palliative to infertility, and with the further development of simpler egg-retrieval techniques it is foreseeable that ovum donation will also become widely used. The major risk to the recipient associated with gamete donation is the transmission of infectious diseases. For the resulting offspring, the risks are not only of infectious disease (such as cytomegalovirus), but also of genetic abnormality. Although the merits of screening for both infectious and genetic diseases have been widely discussed and internationally advocated, there is reasonable concern that some clinics may choose not to follow recommendations for screening\(^\text{181}\) established by such professional groups as The American Fertility Society\(^\text{182}\) and the Canadian Fertility and Andrology Society.\(^\text{183}\)

The probability of transmitting the AIDS virus through donated semen, although very rare, has led to firm recommendations that all donor semen in Canada be frozen and stored for at least six months, until the donor is retested for evidence of the virus. This is necessary because evidence of seropositivity in the donor's blood may not be detectable for some time after exposure.\(^\text{184}\)

\(^{180}\) See infra, note 172.

\(^{181}\) A survey of 11,000 physicians participating in AID in the U.S., completed in 1987, found that one-fifth of centres surveyed did not screen donors for sexually transmitted diseases and fewer than half screened for genetic diseases. Further, of those that did screen for genetic diseases, many did not screen appropriately; for example, donors were rejected unnecessarily in some cases and in other cases accepted when there was significant risk. See OTA, Artificial Insemination: Practice in the United States (Washington, D.C.: OTA, 1988) at 8, 33-40. Also, in 1985 an Ontario Law Reform Commission survey of 16 physicians performing AID in Ontario found that donor screening practice varied considerably. See OLRC, supra, note 2 at 22 n. 36-38; see also Barratt, Chavan and Cooke, supra, note 161.

\(^{182}\) The American Fertility Society, "New Guidelines for the Use of Semen Donor Insemination: 1990" (1990) 53:3 (Supp. 1) Fertil. Steril. 18f. see also infra, note 186.

\(^{183}\) Canadian Fertility and Andrology Society, supra, note 11 at 3. Guidelines for Therapeutic Donor Insemination were adopted in October 1988, stating: "Rigorous attention must be paid to all aspects of donor screening and management to reduce the risks of transmitting genetic or other diseases to the recipients to the absolute minimum possible level in accordance with all currently available screening and testing procedures."

\(^{184}\) Six cases of HIV infection have occurred via donated frozen semen (four in Australia and two in Canada). See Supplement to Health and Welfare Canada, Federal Centre for AIDS, "Guidelines for Prevention of HIV Infection in Organ and Tissue Transplantation" (October 1989) 1556 (Supp.) Canadian Diseases Weekly
Other transmissible diseases that should be screened for include hepatitis, cytomegalovirus, herpes, gonorrhea, chlamydia and mycoplasma. This is done by culture of the semen or through blood tests. Careful screening of the donor with regard to lifestyle and medical and sexual history also reduces the risk of transmitting infectious diseases by rejection of high-risk donors.

Ovum donors should be screened in the same manner as sperm donors even though it is not known whether the organisms in question can be transmitted by ova. Since ovum freezing is not common, the time span between donation and acceptance of the ovum by the recipient is limited. Therefore, prompt screening to the extent possible should be carried out to ensure the safety of the recipient and her potential pregnancy. The risk of passing on genetic disease through ovum donation is the same as or even greater than with sperm donation, since there is the added risk of X-linked disorders. Thus, even under time restrictions, it is important that screening guidelines for genetic disease be followed in order that the recipient be fully informed and thus able to make decisions regarding the degree of potential risk.

Report 1 at 2. A more recent report from New York City found that infected semen from six donors was used in 178 women and one woman was recently found to be seropositive. On this issue see Mary Ann Chiacon, Rand L. Stoneburner and Stephen C. Joseph, "Human Immunodeficiency Virus Transmission through Artificial Insemination" (1990) 32 J. Acquired Immune Deficiency Syndrome 69. See also Canadian Fertility and Andrology Society, supra, note 11 at 3, which states that "there is no place for fresh semen in TDI [AID]. Semen cryopreservation must be used in conjunction with repeated AIDS screening of the donors so that only semen which has been quarantined for an absolute minimum period of 6 months (and, wherever possible 12 months) be used." See also Edwin P. Peterson, Nancy J. Alexander and Kamran S. Moghissi, "A.I.D. and AIDS: Too Close for Comfort" (1988) 49:2 Fertil. Steril. 209; Barratt, Chaulan and Cooke, supra, note 161.

185. Canadian Fertility and Andrology Society, supra, note 11 at 5.
188. See supra, note 175.
189. For an extensive discussion of screening, see Jallbert et al., supra, note 151 at 269-75; and F. Clarke Fraser and R. Allan Forne, "On Genetic Screening of Donors for Artificial Insemination" (1981) 10 Am. J. Med. Genet. 399. Canadian Fertility and Andrology Society, supra, note 11 at 10; see also The American Fertility Society, supra, note 182.
190. The Canadian Council of Medical Geneticists is currently developing guidelines for genetic screening of ovum donors (Personal communication, Dr. F.C. Fraser).
Embryo donation carries with it the added (but necessary) burden of genetic screening of both parents. Ideally, the data should be kept in a registry in the event that an abnormality occurs at birth or a genetic disorder develops at a later time. Surrogacy should require the same stringent screening procedures for sexually transmitted and genetic diseases.

To underscore the importance of all this, we might consider the following: the case of a surrogate who, unbeknownst to the contractual parents, was HIV positive. Subsequently, it was discovered that the newborn child was also seropositive. Both the contractual parents and the surrogate decided against keeping the child. Even though the contractual mother and the surrogate were sisters, the surrogate did not reveal that she was a drug addict and therefore at high risk for contracting the AIDS virus. This illustrates that even in the case of known donors appropriate screening should be undertaken; most importantly, it demonstrates the consequences the child may have to bear in the absence of it.

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191. The literature suggests that in the case of gamete and embryo donation, record keeping (using codes to protect the anonymity of both donors and recipients) allowing the donor to be notified in the case of abnormality of the child may be important. One report suggests that notification is warranted when the condition is severe, carries a risk of recurrence (for future offspring) and is preventable. See Jahnert et al., supra, note 151 at 272. It appears that appropriate notification of donors should be subject to further discussion.

<table>
<thead>
<tr>
<th>Country/Country Group</th>
<th>Year</th>
<th>Number of Embryos (IE4)</th>
<th>Number of Co-Cycles (EC)</th>
<th>Number of Clinical (IC)</th>
<th>Number of Miscarriages (MI)</th>
<th>Number of Live Births (LB)</th>
<th>Multiple Births (MB)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Canada</strong></td>
<td>1982-88</td>
<td>5,921</td>
<td>3,277</td>
<td>4,474</td>
<td>11.3%</td>
<td>20.4%</td>
<td>10</td>
</tr>
<tr>
<td><strong>United Kingdom</strong></td>
<td>1988</td>
<td>10,489</td>
<td>7,515</td>
<td>6,553</td>
<td>12.9%</td>
<td>18.9%</td>
<td>159</td>
</tr>
<tr>
<td><strong>Australia and New Zealand</strong></td>
<td>1986</td>
<td>4,967</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td><strong>United States</strong></td>
<td>1979-85</td>
<td>4,867</td>
<td>3,065</td>
<td>2,845</td>
<td>10%</td>
<td>10%</td>
<td>151</td>
</tr>
<tr>
<td><strong>United States</strong></td>
<td>1986</td>
<td>4,867</td>
<td>3,065</td>
<td>2,845</td>
<td>10%</td>
<td>10%</td>
<td>151</td>
</tr>
</tbody>
</table>

**Sources:**

*Note:* IE4 = Insemination embryo transfer; EC = Embryo cryopreservation; IC = In-vitro fertilization cycle; MI = Miscarriage; LB = Live birth; MB = Multiple birth.
### Table II: GIFT International Results

<table>
<thead>
<tr>
<th>Countries</th>
<th>Number of GIFT Procedures (CP)</th>
<th>Number of Clinical Pregnancies (CP)</th>
<th>% of Clinical Pregnancies (CP)</th>
<th>Number of Miscarriages (M)</th>
<th>% of Miscarriages (M)</th>
<th>Number of Ectopic Pregnancies (EP)</th>
<th>% of Ectopic Pregnancies (EP)</th>
<th>Number of Deliveries (D)</th>
<th>% of Deliveries (D)</th>
<th>Number of Live Births (LB)</th>
<th>% of Live Births (LB)</th>
<th>Number of Multiple Births (MB)</th>
<th>% of Multiple Births (MB)</th>
<th>MB:D</th>
</tr>
</thead>
<tbody>
<tr>
<td>United Kingdom</td>
<td>1988</td>
<td>3,392</td>
<td>707</td>
<td>21%</td>
<td></td>
<td>1</td>
<td>9.6</td>
<td>139</td>
<td>20%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Australia</td>
<td>1986</td>
<td>607</td>
<td>136</td>
<td>22%</td>
<td>21</td>
<td>15%</td>
<td>7</td>
<td>108</td>
<td>18%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>United States</td>
<td>1987</td>
<td>1,968</td>
<td>492</td>
<td>25%</td>
<td>116</td>
<td>24%</td>
<td>30</td>
<td>362</td>
<td>18%</td>
<td>489</td>
<td>28%</td>
<td>103</td>
<td>28%</td>
<td></td>
</tr>
</tbody>
</table>


### Table III: Effects of the Number of Embryos Transferred on Pregnancy

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% of clinical pregnancies</td>
<td>% of multiple pregnancies</td>
<td>% of clinical pregnancies</td>
</tr>
<tr>
<td>1</td>
<td>9.6</td>
<td>1</td>
<td>11</td>
</tr>
<tr>
<td>2</td>
<td>14.2</td>
<td>13.1</td>
<td>12</td>
</tr>
<tr>
<td>3</td>
<td>25.2</td>
<td>29.2</td>
<td>20</td>
</tr>
<tr>
<td>4</td>
<td>23.4</td>
<td>24.2</td>
<td>22</td>
</tr>
<tr>
<td>5</td>
<td>24</td>
<td>29</td>
<td></td>
</tr>
</tbody>
</table>

CHAPTER TWO

Issues in Canadian Law

By and large, Canadian law has not adapted easily to the various situations made possible by medically assisted procreation. We will therefore outline in this second chapter the main problems these technologies create for the law. The scope of the task is clearly illustrated by the number of branches of the law and the range of legislation, both federal and provincial, that must be considered. The legal framework within which medically assisted procreation is developing is in fact very broad. It covers such diverse concepts, principles and branches of the law as public policy, parentage, the principle of the non-availability of the human body for commercial purposes,\textsuperscript{193} the right to life, the right to liberty, property law, contract law and tort law. In order to identify the legal problems, we will first examine the main rules of law applicable to medically assisted procreation.

I. Medically Assisted Procreation and Private Law

As we will see, a number of rules of private law affect medically assisted procreation. We will look at the rules that govern consent, parentage, successions, contracts, property and liability and their impact on the use of artificial insemination, in vitro fertilization, gamete intrafallopian tube transfer (GIFT) and egg retrieval by uterine lavage. We will consider these rules from the perspective of potential parents, donors, children and medical personnel, because they are the most affected by the legal problems that can arise when medically assisted procreation technologies are used.

A. Potential Parents

I. Consent

The decision to have a child is a private one, normally made within a marriage by both partners. However, medically assisted procreation makes it possible for a woman to conceive a child without her spouse's knowledge; this raises the problem of attributing

\textsuperscript{193} See \textit{infra} at 41ff.
paternity to a man who is not genetically linked to the child and did not consent to conception.194 Irrespective of the remedies available to the spouse (separation, divorce, disavowal of paternity),195 we may consider the need for specific legislative intervention in this area. Should the law make the consent of both spouses a prerequisite for medically assisted procreation?

The problem with such intervention is obvious. Giving a husband the power to decide whether a child should be conceived in effect violates the wife’s right to control her own body and her reproductive autonomy.196

Currently only Quebec, the Yukon Territory and Newfoundland have provisions dealing with consent to medically assisted procreation, although there is no specific indication of the form of such consent. Article 586 of the Civil Code of Quebec (C.C.Q.) reads as follows:

When a child has been conceived through artificial insemination, either by the father or, with the consent of the spouse, by a third person, no action for disavowal or contestation of paternity is admissible [emphasis added].197

194. We are referring here to cases where gametes from a third person are used, since the spouse’s sperm cannot be used without his consent.


[TRANSLATION]

In all systems of law, spouses accept, through the bond of marriage, the obligation to help, assist and be faithful to one another. It is perfectly logical, therefore, to say that AIH without the husband’s knowledge may be considered a failure to meet that obligation and may become general grounds for divorce or separation as “outrage, illusage or grievous insult,” “mental cruelty” or “irretrievable damage to the will to maintain the bond of marriage.”

196. Requiring the husband’s consent is viewed by some as contradicting health legislation. See Bartha Maria Knoppers, Conception artificielle et responsabilité médicale (Ottawa, Que.: Yvon Blais, 1986) at 96, in the area of consent to medical treatment or access to medical services, respect for the person’s autonomy prevails. Consequently, if one spouse is able to express his or her own wishes, the consent of the other cannot be required before treatment is administered. See also Ellen I. Pickard, Legal Liability of Doctors and Hospitals in Canada, 2nd ed. (Toronto: Carswell, 1984) at 62-63. In Quebec, An Act respecting health services and social services, R.S.Q., c. S-5, s. 156, states that: “The consent of the consent shall not be required for the furnishing of services in an establishment.” See also arts. 19, 19.1 to 19.4 of the Civil Code of Lower Canada (C.C.L.C.) and art. 10ff. of Bill 125, Civil Code of Quebec, 1st sess., 34th Leg., Quebec, 1990 (1st Reading, 18 December 1990) [hereinafter Bill 125]. In Ontario, see the Family Law Act, 1986, S.O. 1986, c. 4, s. 64(2): “A married person has and shall be accorded legal capacity for all purposes and in all respects as if he or she were an unmarried person.”

197. Article 580 of Bill 125, supra, note 196, reiterates these principles:

No person may contest the filiation of a child on grounds relating to his medically assisted procreation, and no claim to another status is admissible from the child.

However, the husband of the mother may disavow the child or contest acknowledgment if he did not give consent to medically assisted procreation or if he proves that the child was not born of such procreation.

For the Yukon, see Children’s Act, R.S.Y.T. 1986, c. 22, s. 133(3). Subsection 133(3) states: “A man who is married to a woman at the time she is artificially inseminated solely with the consent of another man shall be deemed in law to be the father of the resulting child if he consents, in advance, to the insemination.”

For Newfoundland, see The Children’s Law Act, S.N. 1988, c. 61, s. 12(3).
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\textsuperscript{194} We are referring here to cases where gametes from a third person are used, since the spouse's sperm cannot be used without his consent.

\textsuperscript{195} Michèle River, "Quand la médecine intervient dans la genèse de la conception, que fait le droit? Ou le délicat problème de l'insémination artificielle" in \textit{Association Henri Capitant. Le corps humain et le droit: Journées belges, vol. 26} (Paris: Dalloz, 1977) 87 at 95. \textit{Translation} "Artificial insemination, whether AI or AIH, without the husband's consent does not constitute adultery but is a violation of matrimony that in itself warrants sanction."; Jean-Louis Badréau, "Aspects juridiques" in Marcel J. Melancon, ed., \textit{L'insémination artificielle thérapeutique} (Quebec: P.U.L., 1985) 113 at 121.

\textit{Translation} In all systems of law, spouses accept, through the bond of marriage, the obligation to help, assist and be faithful to one another. It is perfectly logical, therefore, to say that AIH without the husband's knowledge may be considered a failure to meet that obligation and may become general grounds for divorce or separation as "outrage, ill-usage or grievous insult," "mental cruelty" or "irremediable damage to the will to maintain the bond of marriage."

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For the Yukon, see \textit{Children's Act}, R.S.Y.T. 1986, c. 22, s. 13(3)-(5). Subsection 13(3) states: "A man who is married to a woman at the time she is artificially inseminated solely with the semen of another man shall be deemed in law to be the father of the resulting child if he consents, in advance, to the insemination."

For Newfoundland, see \textit{The Children's Law Act}, S.N. 1988, c. 61, s. 12(3).
of the producer of the gametes? Three decisions, one in France in 1984,\textsuperscript{202} the other two in the United States in 1988 and 1989 — \textit{Davis}\textsuperscript{203} and \textit{York}\textsuperscript{204} — ruled on such matters. We will turn our attention first to gamete storing in light of the French ruling, and then to the fate of embryos in light of the U.S. rulings.

On the subject of gamete storing, a French court was asked to rule in 1984 in the \textit{Parpalais}\textsuperscript{205} case, which involved a dispute between CECOS, a centre for sperm analysis and storage, and the widow of a man who had stored sperm at the centre. The centre refused to grant the widow's request to retrieve the sperm so that she could be artificially inseminated. The widow argued that her husband and the centre had signed a contract of deposit.

The judge ruled that in the case in question the agreement was not a contract of deposit but rather an innominate contract under which CECOS agreed to store the sperm and return it on request to the producer or, following his death, his heirs. The judge wrote:

\begin{quote}
[\textbf{Translation}]

The rules of the contract of deposit as defined by article 1915 \textit{et seq.} C.C. cannot be applied in the case at bar, which concerns not \textit{objects of commerce}, but rather a secretion that contains the seed of life and is to be used to produce a human being. . . .

It appears that the agreement of 7 December 1981 was a specific contract under which CECOS was obligated to store the sperm and return it to the donor or the woman for whom the sperm was intended [emphasis added].\textsuperscript{206}

The court did not go so far, however, as to determine that the husband owned the gametes. Rather, it ordered that the sperm be returned on the basis of the deceased man's intentions and the absence of a stipulation that CECOS intended to keep the sperm if the producer died.

The reference by the judge in the case to things that are not objects of commerce needs clarification. In the civil law, things that are not objects of commerce are not subject


\textsuperscript{203} \textit{Davis v. Davis} (21 September 1989), Broun Ct B-14496 (Cir. Ct) at 1-2; this decision has since been reversed by 59 U.S.L.W. 2205 (Tenn. App. 1990). See infra, note 216 for more details.

\textsuperscript{204} \textit{York v. Jones Institute}, 717 F. Supp. 421 (E.D. Va 1989) (order of 10 July 1989 denying defendants' motion to dismiss). See infra, note 217, for more details. Note that the United States and Australia have also had to consider this issue in the \textit{Rios} case: \textit{In re Esau of Elsa and Mario Rios} (May 1985), Los Angeles Ct P680582, P680583 (Sup. Ct). The California Superior Court decided not to appoint a guardian for the embryos and ruled that they were neither the heirs nor the property of the Rioses. See George P. Smith, "Australia's Frozen 'Orphan' Embryos: A Medical, Legal and Ethical Dilemma" (1985) 24 J. Fam. L. 27. See also Tamara L. Davis, "Protecting the Cryopreserved Embryo" (1990) 57 Tenn. L. Rev. 307 at 518.

\textsuperscript{205} \textit{Supra}, note 202. For an interesting analysis of this ruling, see Jones, \textit{supra}, note 147.

\textsuperscript{206} \textit{Parpalais}, supra, note 202 at 562.
to human will and cannot be disposed of, even gratuitously.\footnote{207} The word "commerce," therefore, has a very specific meaning here.

\textbf{[Translation]}

a special meaning more general than its usual one. It refers not only to commercial transactions \textit{per se} . . . but to any legal act the purpose of which is to create, modify or extinguish rights. A thing that is not an object of commerce is a thing that cannot be the object of legal acts performed by individuals. "Commerce" evokes the notion of things circulating among persons, but it is not synonymous with the economic term "market."\footnote{208}

This broad definition explains in part why some French authors are sceptical about including the body and its parts and substances among things that are not objects of commerce.\footnote{209}

In any event, in Quebec civil law article 20 C.C.C. permits the \textit{inter vivos} disposal of parts or products of the body,\footnote{210} even in return for payment. It may therefore be concluded that, to the extent that article 20 applies to sperm and ova, gametes could be objects of commerce in Quebec civil law.\footnote{211}

\footnote{207} Marie-Ève Hermitte, "Le corps hors du commerce, hors du marché" (1988) 33 Arch. philo, cbr. 323 at 325.

\footnote{208} Jean-Christophe Galloux, "Réflexions sur la catégorie des choses hors du commerce: l'exemple des éléments et des produits du corps humain en droit français" (1989) 30:4 C. de D. 1011 at 1015-16.

\footnote{209} Hermitte, supra note 207 at 327, holds the view that the body itself is an object of commerce. Commenting on s. 1128 of the French Civil Code (equivalent to art. 1059 C.C.C.), she writes:

\textbf{[Translation]}

It is therefore not the body that is protected in this way, placed beyond the exercise of will by article 1128, but rather the person, a legal abstraction defined by attributes, themselves abstract, that are considered to be the framework of human dignity, . . . This illustrates by reduced as abstruse that the civil law views the body as nothing more than an incidental medium for representations that centre on the person, defined by changing references to morals, dignity and liberty. Violations of the body are not taken into consideration until they engender a violation of these values. See also Galloux, supra, note 208 at 1019, on the limited scope in French law of the notion of "extracommerciality" as it relates to the products and elements of the human body. Citing as examples blood, mother's milk and gametes, he concludes that [Translation] "[H]uman products circulate among private or public individuals; they do not, as the status of extracommerciality would require, remain under the exclusive control of the person from whom they came."

\footnote{210} Jean-Louis Baudoz and Catherine Labrousse-Riou, \textit{Produire l'homme: De quel droit?} (Paris: P.U.F., 1987) at 44: [Translation] "In Quebec law, the provisions so adopted apply not only to organs themselves, but also body substances (blood, sperm, etc.)." See also arts 19 and 24 of Bill 125, supra, note 196. However, art. 25 of the bill drops the distinction between parts of the body that are capable of regeneration and those that are not by requiring that alienation be gratuitous in all cases. Art. 25 reads: "The alienation by a person of a part or product of his body shall be gratuitous; it shall not be repeated if it involves a risk to his health."

\footnote{211} Baudouin and Labrousse-Riou, supra, note 210 at 115: [Translation] "However, such liberalization is possible only from a therapeutic perspective or at least one of scientific experimentation leading to the development of a treatment."
It follows therefore that article 1059 C.C.L.C., which states that "[r]ose things only which are objects of commerce can become the object of an obligation," could not impede the creation of rights and obligations between the bank and persons who deposit their gametes. Gametes could thus be the subject-matter of a contract. It should be noted, however, that such freedom of contract would be subject to the criterion set out in article 20(1) (proportionality of risks) and to articles 13 and 990 C.C.L.C. (public order and good morals).

The agreement between the bank and the person storing his or her gametes could therefore be used to create rights and obligations for the parties. With respect to the couple, it would be sufficient for each partner to enter into a separate contractual relationship with the bank to have independent control over his or her gametes. This would prevent disputes over the use of one partner’s gametes if, for any reason, he or she no longer wished to proceed with the parental plan. However, an additional problem arises with ova. Since freezing of ova seems to pose major difficulties, the normal procedure is to freeze eggs that have been fertilized in vitro (embryos). Once the egg has been fertilized, independent control of the gametes must give way to a form of joint control.

Whether the embryo is produced by the couple or from one or two donated gametes, the question of control is a delicate problem. In case of separation, for example, both spouses may claim the embryos, or one spouse may object to their being used. We must therefore provide for how such disputes can be resolved where no provision has been made in the contract or the consent form signed by the spouses. The U.S. decisions in Davis v. Davis.

212. See also art. 1058 C.C.L.C.
213. In the event of a conflict, the specific prevails over the general provision: Pierre-André Côté, The Interpretation of Legislation in Canada (Cowansville, Que.: Yvon Blais, 1984) at 240.
214. The risk assumed must not be disproportionate to the benefit anticipated. See art. 20 C.C.L.C., infra at 47.
216. Supra, note 203. The case centered on the absence of a consent form or document providing for the disposition of the embryos in the event that Mr. and Mrs. Davis divorced. Despite the divorce, Mrs. Davis wanted to use the frozen embryos in the hope of having a child, but Mr. Davis objected on the grounds that he had no intention of becoming a father. Mrs. Davis claimed that the embryos were living and that as a mother she was entitled to use them to try to conceive. She argued that if she were unable to use them herself, the embryos should be given to an infertile couple so that they could be carried to term. The trial judge, W. Dale Young, ruled as follows (supra, note 203 at 1-2):
and York v. Jones Institute\textsuperscript{217} bear witness to the problems that can arise and the difficulty in resolving them.

It is to be hoped that the experience gained from these cases will result in more suitable consent forms. In any case, is it possible to make provision in a consent form for the fate of an embryo in the event of a dispute, separation, divorce or death?\textsuperscript{218} This leads us

\begin{quote}
The salient findings, conclusions and the judgment are summarized as follows, to-wit: (1) Mr. and Mrs. Davis undertook in vitro procedures for the purpose of producing a human being to be their child. (2) The seven cryogenically preserved embryos are human embryos. \ldots (5) From fertilization, the cells of a human embryo are differentiated, unique and specialized to the highest degree of distinction. (6) Human embryos are not property. (7) Human life begins at conception. (8) Mr. and Mrs. Davis have produced human beings, in vitro, to be known as their child or children. (9) For domestic relations purposes, no public policy prevents the continuing development of the common law as it applies to the seven human beings existing as embryos, in vitro, in this domestic relations case. (10) The common law doctrine of parens patriae controls children, in vitro. (11) It is to the manifest best interests of the child or children, in vitro, for their Mother, Mrs. Davis, to be permitted the opportunity to bring them to term through implantation. \ldots

The temporary custody of the seven cryopreserved human embryos is vested in Mrs. Davis for the purpose of implantation. All issues of support, visitation, final custody and related issues are reserved to the Court for consideration and disposition at such time as one or more of the seven human embryos are the product of live birth. \textsuperscript{219}


Judge Young's conclusion that four-celled preimplantation human embryos are "children" and "human beings" is unprecedented and unwarranted. It has no discernible basis in common law precedents nor in Tennessee law (which recognizes a separate legal interest in prenatal human life only at viability). It is a view rejected by highly respected ethical advisory bodies in the United States, Great Britain, Canada, France, and several other countries. This remarkable conclusion appears to represent the judge's own personal view of the significance of the biological fact that a new human genome exists at or shortly after fertilization.

On appeal from the Davis decision (supra, note 203 at 2206), the judge ruled:

The trial court in its fact finding and legal conclusions, ignored the public policy implicit in the Tennessee Statutes, the cases holdings of the Tennessee Supreme Court and the teachings of the United States Supreme Court. We are required to receive the issue consistent with the existing Tennessee law and the parties' constitutional rights. On the facts of this case, it would be repugnant and offensive to the constitutional principles to order Mary Sue to implant these fertilized ova against her will. It would be equally repugnant to order Junior to bear the psychological, if not the legal consequences of paternity against his will. Jointly, the parties share an interest in the seven fertilized ova.

Accordingly, the cause is remanded to the trial court to enter a judgment vesting Mary Sue and Junior with joint control of the fertilized ova with equal voice over the disposition.
to question the status of the embryo: Does it fall under the law of property or that of persons? Positive law is unable to answer this question, which is a philosophical, theological and ethical problem: At what point does human life begin?219

The question of control over gametes and embryos has been considered from the perspective of property law:

The elements of use, alienation ... disposal and destruction, even when exercised subject to statutory regulation, appear to comprise the power legally contained in the concept of property ownership. According to property principles, it seems that the gamete donors could exercise control over the embryo extra uterum, abandon their respective rights of control to the exclusive exercise of the other (as in ordinary artificial insemination by sperm donor), agree upon its transplantation into another woman without invoking adoption law, and rely on property principles in settling disagreements on disposition. In the same way, gamete donors may delegate to clinics and clinic personnel their own authority to decide, for instance, which women may receive transplantations of spare embryos.220

However, the nature of the "deposited" product makes such an approach difficult.221 Some hold the view that this property right is inconsistent with the traditional

219. *Biomedical Experimentation Involving Human Subjects, supra*, note 7 at 46-47. See also the recent decisions in *Murphy v. Dedd* (1989), 70 O.R. (2d) 681 (H.C.J.); *Tremblay v. Daigle* (7 July 1989), Ahuntsi 170-05-000012-898 (Quebec Sup. Ct., interim interlocutory injunction); 1989 R.J.Q. 980 (Sup. Ct.) (interlocutory injunction), appeal dismissed 1989 R.J.Q. 1735 (C.A.), reversed by unanimous decision of the Supreme Court of Canada, 1989 2 S.C.R. 550. It should be noted, however, that an embryo (artificial in vitro is not covered by the definition of fetus in the working paper *Crimes against the Fetus, supra*, note 7 at 59: "the product of a union in the womb of human sperm cells and egg cells at all stages of its life prior to becoming a person."

220. *Dickens, supra*, note 218 at 62-63. And at 64-65: "Destruction or other misappropriation of an object without the owner's consent may constitute the crime of theft, and/or the tort of trespass to property and conversion."

221. Barbara M. Knoppers, "Reproductive Technology and International Mechanisms of Protection of the Human Person" (1987) 32 McGill L. J. 336 at 346: [A]ll agree that the embryo in vitro constitutes human life worthy of protection ... while the majority deny the possibility of granting the donor a proprietary interest in human gametes or embryos, most would seem to grant the donor at least some possessory interest, and in some cases, a residual right. ... Indeed, there is no area where the need for some common international principles of respect and protection is more imperative, if we are truly to distinguish between human genetic material as property, as a simple product of conception or as human life.

Barbara M. Dickens, "The Ectogeneric Human Being: A Problem Child of Our Time?" (1988) 18 U.W.O. L. Rev. 241 at 245. "Litigation in the United States arising from hospitals losing or incinerating fetuses their mothers wanted to bury has been framed in terms of causing emotional injury rather than of misappropriating property." The author refers to *Brooks v. South Broward Hospital District*, 325 So. 2d 479 (Fla App. 1975) and *Hembree v. Hospital Board of Morgan County*, 300 So. 2d 823 (1978). In *Del Zio v. Manhattan a Columbus Presbyterian Medical Center* (14 November 1978) 74-3538 (D.S.N.Y.), a woman's fertilized egg was destroyed, a U.S. Federal Court judge allowed a claim for damages for loss of property to be heard by a jury. However, “[t]he jury rejected the property claim but awarded plaintiffs damages for the emotional distress. Mrs. Del Zio was awarded $50,000 for emotional distress and Mr. Del Zio was awarded $3.00.” Lori B. Andrews, "My Body, My Property" (1986) 16:5 Hast. Cent. Rep. 28 at 29ff.
legal interpretation of the concept and that it is rather a right of supervision and control.\textsuperscript{222}

In view of this clear gap in the law, several options are possible. One would be to leave it to the courts to adapt current principles of law to gametes and embryos in order to address the problems they create. Alternatively, we could rethink the traditional legal distinctions between persons and things in order to deal specifically with these products of the human body.\textsuperscript{223} Finally, it might also be appropriate to look into the possibility of adapting, with respect to these substances, the notion of things that are not objects of commerce.\textsuperscript{224} This dilemma is not limited to the problem of control over these substances; it also applies to the very legitimacy of donating gametes and embryos and to their commercialization. These issues will be discussed later.

3. Post-Mortem Use of Gametes and Embryos

The issue of the post-mortem use of gametes and embryos brings us back to the question of the right to dispose of them and of their legal status.\textsuperscript{225} Does a widow or widower have any rights over the gametes of his or her deceased spouse or any frozen embryos that they have conceived together?

\textsuperscript{222} According to Baudouin and Labrosse-Riot, supra, note 210 at 45-46.

[TRANSLATION]

the donor must be recognized as having not a true property right, but more a right of supervision and control over the use of his or her gametes, a matter which is part of the broader issue of personality rights. For obvious social reasons, however, the donor must not be permitted to exercise this right in the same way as a true property right. A balance must therefore be struck between respect for the consent and intentions of the donor on the one hand and on the other the exercise of this right in a manner that is compatible with the ethical and social requirements of society as a whole.

The term “donor” seems to be used here to designate both a person who makes a deposit and a person who makes an actual donation, that is, who relinquishes the product donated; \textit{ibid.} [TRANSLATION] “It is difficult to imagine, for example, a donor bequeathing a large fortune to his granddaughter on the condition that she be inseminated after his death with the sperm be deposited in a bank for this purpose!” [emphasis added]. The Ontario Law Reform Commission writes:

It was suggested that the fact that a person does not have absolute beneficial ownership of an object does not mean that he or she has no property or other (for example, possessory) interest in it. [Note 224: For example, under certain circumstances, the \textit{Anatomy Act}, R.S.O. 1980, c. 21, permits the possession of corpses by medical schools for dissection and medical education, even though it has been said that a dead body cannot be the subject of a property right.] Accordingly, even if there are some restrictions on a woman’s rights respecting her own ovum, presumably some type of interest may still be found in her. And, of course, one person’s right or interest in genetic material, however that right or interest is characterized, may permit that person to, for example, donate it to a hospital for experimental purposes, or require it to be destroyed notwithstanding the claims of others to use it for such purposes: that right or interest may well be superior to that of anyone else. [OLRC, supra, note 2 at 88].

\textsuperscript{223} See Hermite, supra, note 207.

\textsuperscript{224} See Galloux, supra, note 208.

\textsuperscript{225} See supra at 39 and 43.
As we have seen in the Purpalaix case, the French courts have held the storing of sperm to be a specific contract and ordered the sperm to be returned to the widow of the donor, thereby permitting post-mortem insemination. This conclusion, however, could only be reached through a very legalistic interpretation of the dispute, since the judge did not rule specifically on the use of the sperm for post-mortem insemination.

On a more practical level, it is easy to anticipate the serious problems that post-mortem use of gametes and embryos will create in parentage law and the law of Successions. In Canada, neither the law of Successions nor parentage law recognizes post-mortem procreation. Presumptions of paternity normally use a test of 300 days between the death of the father and the birth of the child. In Ontario, if no one is presumed to be the father under the 300-day test, any person may apply for a declaration of paternity provided that "both the persons whose relationship is sought to be established are living." We may therefore question the appropriateness of making provision in Canadian law for the impact of the post-mortem conception of a child on parentage and inheritance rights. But we must first decide if we wish to allow or prohibit the post-mortem use of gametes and embryos in Canadian society.

B. Donors

1. The Legality, Legitimacy and Nature of Gamete and Embryo Donation

The legality of such a donation is linked to the principle of the inviolability of the human body.

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227. For example, see the Rin case, supra, note 204.

228. For example, see Children's Law Reform Act, R.S.O. 1980, c. 8(1) 2nd para., which presumes paternity if "... the child was fathered by the mother of the child by a marriage that was terminated by death ... within 300 days before the birth of the child." In Quebec, art. 574 C.C.Q. states that a child born more than 300 days after the death of the biological father is deemed to have been conceived after the father's death; see also art. 523 of Bill 125, supra, note 196.

229. See Children's Law Reform Act, supra, note 228, s. 5(1).

230. Ibid., s. 5(2). Dickens, supra, note 221 at 247, "This would bar, of course, both the dead biological father and the unborn child, and appear to leave the fetus not that of the father for the purposes of the perpetuity rule or other property interests." It should be noted, however, that the Uniform Child Status Act (supra, note 199) provides for an exception to this rule where a presumption of paternity pursuant to s. 9 applies. The presumption of paternity provided for in the event of the death of the child's father indicates no time limit regarding the birth of the child or the death of the husband. See ss 6(6) and 9(a).

231. Legality is used here to mean the compliance of an action or undertaking with the law, while legitimacy refers to the ethical and social criteria that make it desirable to prohibit, allow or simply tolerate such an action or undertaking.

232. In Quebec, this principle is established by art. 19 C.C.L.C.: "The human person is inviolable. No one may cause harm to the person or another without his consent or without being authorized by law to do so." See also arts 19.1-19.4 C.C.L.C. and art. 10ff. of Bill 125, supra, note 196.
Inviolability . . . may have two contents of meaning. It may connote that one is not justified in treating another without his consent, but is justified in doing so with it, in which case it is merely a particular application of the autonomy principle; or it may indicate a principle that protects a person's physical and mental integrity against non-beneficial acts by the person himself, or others, when it is a preservation of life value.  

This second interpretation of the principle of the inviolability of the human body imposes a limit on what a person can consent to. It is this second aspect that seems to have captured the Quebec legislature's interest when the *Civil Code of Lower Canada* was amended in 1971.  

Article 20 *C.C.L.C.* allows competent persons to dispose of parts of their bodies, whether or not the body part is capable of regeneration, subject in both instances to article 19.1 (proportionality of risks) and written consent. Article 20 reads as follows:  

A person of full age may consent in writing to disposal *inter vivos* of a part of his body or submit to an experiment provided that the risk assumed is not disproportionate to the benefit anticipated.  

A minor capable of discernment may do likewise with the authorization of a judge of the Superior Court and with the consent of the person having parental authority, provided that no serious risk to his health results therefrom.  

The alienation must be gratuitous unless its object is a part of the body susceptible of regeneration.  

The consent must be in writing; it may be revoked in the same way.  

To the extent that the conditions set out in article 20 are met and gametes are considered parts of the body within the meaning of the article, there is no doubt in Quebec law as to the legality of gamete donation.  

In the common law provinces, tissue donation is covered by statutes based on the *Uniform Human Tissue Gift Act*, subsection 3(1) of which reads as follows:  

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233. Margaret A. Somerville, *Consent to Medical Care*, study paper prepared for the LRC (Ottawa: Supply and Services Canada, 1980) at 5.  
235. See Baudouin et Labrèce-Riou, *supra*, note 210 at 44. It should be noted that arts 18-22 of *An Act to add the reformed law of persons, Successions and property to the Civil Code of Quebec*, S.Q. 1987, c. 18, essentially restate art 2011 *C.C.L.C.* and add specific requirements and conditions. See Monique Ouimet, "De la justification et de l'existence des droits civils et de certains droits de la personnalité" (1988) 1 C.P. du N. 1 at 20. See also arts 19-25 of Bill 125, *supra*, note 196.  
Any person who has attained the age of majority, is mentally competent to consent, and is able to make a free and informed decision may in a writing signed by him consent to the removal forthwith from his body of the tissue specified in the consent and its implantation in the body of another living person.237

Although the Uniform Human Tissue Gift Act (UHTGA) does not state that the risk incurred is not to be disproportionate to the anticipated benefit, the common law recognizes this condition.238 However, unlike the Civil Code, the UHTGA excludes tissue capable of regeneration.239

Thus, insofar as sperm and even eggs (despite the fact that eggs are limited in number) are parts of the body capable of regeneration,240 statutory provisions on human tissue

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237. Supra, note 236, s. 1(c) defines ‘tissue’ as follows: ‘tissue’ includes an organ, but does not include any skin, bone, blood, blood constituent or other tissue that is replaceable by natural processes of repair.”


   Every one is protected from criminal responsibility for performing a surgical operation on any person for the benefit of that person if
   (a) the operation is performed with reasonable care and skill; and
   (b) it is reasonable to perform the operation, having regard to the state of health of the person at the time the operation is performed and to all circumstances of the case.

See also s. 14 of the Criminal Code.

239. Supra, note 236. The latest version of the statute, Uniform Human Tissue Donation Act, supra, note 236, available for adoption by the provinces, provides in s. 1 that “tissue” means a part of a living or dead human body, but does not include (a) spermatozoa or ova, or (b) an embryo or fetus. It remains to be seen whether the provinces will adopt these amendments.

240. See Knoppers, supra, note 196 at 109; Helene, supra, note 234 at 61. [Translation] “A medicine as we know it today, blood, milk, hair, skin, bone marrow and genetic material are considered parts of the body susceptible of regeneration.” Commenting on the Ontario statute (supra, note 236), the OLRC wrote in its report, supra, note 2 at 63:
donation do not apply, and the donation of gametes is governed by the common law: "Under the common law, an adult, if fully informed, can consent to having regenerative tissue removed from his body. Indeed, the Red Cross Blood Transfusion Service is wholly dependent on such donations." 241

While we may conclude (subject to the ambiguity surrounding the characterization of gametes) that our systems of law seem to cast aside any doubt as to the legality of gamete donations, the legitimacy of such donations is certain to draw comment because of the very nature of gametes.

For some authors, discussions of gamete and embryo donation are too often centred on the controls and conditions that may be imposed, thereby clouding the question of the ethical value of such donations:

[TRANSLATION]
Some essential questions concerning the donation of gametes have thus been purely and simply "medicalized" and therefore trivialized. Science has taught us to think that the problems lie not in the ethical value of the actual procedure, but rather, because the procedure is established, in the controls and conditions which may be imposed. Accordingly, the public and jurists have been conditioned to believe that the legitimacy and, consequently, the legality of gamete donation were no longer open to discussion in ethical and legal terms. All that remained was to look to the law for procedural management models. In focusing the debate on the question of "how," we truly lost sight of "why." Science has acted as if the only real problems were technical. 242

Some hold the view that the question of the legitimacy of gamete donation cannot be settled until an ethical and legal analysis has been conducted of the fundamental issues gamete donation raises for our society. Marie- Angèle Hermitte summarized the question as follows:

[TRANSLATION]
We have to decide whether or not we want to be a society that considers kinship to be subject to commercial transactions... To answer the question, we need to determine whether the "transaction" is to be analyzed as simply a donation of life or as a shift in the order of consanguinity. The analysis in the first case is materialistic and purely biological. AID is viewed as a somewhat magical treatment for sterility. In the second case, the donation

The critical question is, of course, whether the Act applies to the donation of sperm or ova, or, put another way, whether sperm or ova come within the definition of "tissue" in the Act. Section 1(e) provides that "tissue" includes an organ, but does not include any skin, bone, blood, blood constituent or other tissue that is replaceable by natural processes of repair...

There would appear to be little controversy that sperm comes within the closing flush of section 1(e) and, accordingly, is outside the purview of section 3(1). However, the same may not be said of ova. A woman's complement of ova is fixed and not replaceable; she loses one or more during menstruation from puberty to menopause.

241. Picard, supra, note 196 at 129. See also Dickens, supra, note 238 at 163-64.
of life is part of the logic of genealogy, and one realizes that a branch of the family tree is being broken. In the end, the questions remain the same: Are bodies nothing more than living matter which may be passed on according to the rules of trade? Are they not also the medium for the cultural representations that transcend them, at least in part? 243

Having briefly analysed the question of gamete donation, we must now turn our attention to embryo donation. Neither the civil nor the common law provides for the donation of embryos. Indeed, it may be difficult to consider embryos as tissues or parts of the body (whether capable of regeneration or not). The ambiguity of the embryo’s status gives rise to moral and social objections that have appeared with the creation and freezing of surplus embryos. What is at issue here is one’s image of the embryo: Is it a thing, a person, a potential person, or something else? The question of the legitimacy of embryo donation therefore remains completely unanswered. Do we wish to legalize or prohibit embryo donation?

To end this discussion of the legality and legitimacy of gamete and embryo donation, we might ask ourselves in more general terms whether we wish to treat gametes and/or embryos differently from other parts of the body or alienable cells, or in other words, create a special regime suited to the specific nature of gametes and embryos. 244

In concluding, we should address the question of the very nature of gamete and embryo donation. Are such donations blind or conditional? In other words, are donors entitled to attach conditions to the donation? May they withdraw their consent?

While the donation of tissues and body substances such as blood, milk and bone marrow creates few moral problems, the donation of gametes and embryos, which entails the potential to create a human life, is more problematic. In light of the significance of such donations, the donor may wish to attach conditions to how the gamete or embryo is used. For this reason, gamete and embryo donation should not be permitted without the free and informed consent of the donor, not only regarding the procedure and the risks of donating, but also the ultimate purpose of the donation. It is therefore essential that the donor be told how the donated gametes or embryo will be used.

243. Supra, note 207 at 337.

244. In its working paper on experimentation, the LRC indicated that: “Gametes and human embryos cannot be considered to be simple cells or simple tissues. The first are the virtual sources of new human life; the second already have life.” See Biomedical Experimentation Involving Human Subjects, supra, note 7 at 53.
It would appear that at common law conditions may be attached to donations of organs and tissues that do not regenerate. This is all the more reason why the donation of gametes and embryos should be subject to conditions, provided such conditions are not discriminatory.

The right of a donor to withdraw his or her consent to a donation is provided for in Quebec civil law in article 20 C.C.L.C. At common law, such right is determined by the nature of the contract. If the contract is deemed to be a contract for a service, the donor may withdraw his or her consent at any time. However, if the contract is for the sale of a good, the donor would not have the option of withdrawing consent. We might ask whether it is appropriate that the revocable or irrevocable nature of the consent should depend on the nature of the contract (sale of a good or contract for a service). Are not the nature of the donated product and the significance of a donation of life sufficient to warrant the right to revoke a donation of gametes or embryos?

2. Gamete and Embryo Donations: Free and Anonymous

Under current law in the common law provinces, the sale of tissues and parts of the human body is generally prohibited by provisions which deem such sale to be "contrary to public policy."

No person shall buy, sell or otherwise deal in, directly or indirectly, for a valuable consideration, any tissue for a transplant, or any body or part or parts thereof other than blood or a blood constituent, for therapeutic purposes, medical education or scientific research, and any such dealing is invalid as being contrary to public policy.

245. Uniform Human Tissue Gift Act, supra, note 236, s. 3(6): "If for any reason the tissue specified in the consent is not removed in the circumstances to which the consent relates, the consent is void."

246. See Staudenmaier, supra, note 210 at 45: "[TRANSLATION] A donation of sperm is not a 'blind' donation, but rather a deliberate, directed donation, a conditional donation that is part of a true agreement between the donor, the physician and, through the physician, the recipient or recipients. Donating the potential for human life is not an ethically neutral act and must therefore be evaluated and respected as such."

247. See supra at 47. An Act to add the reformed law of persons, succession and property to the Civil Code of Quebec, supra, note 235, art. 21, provides that consent may even be revoked verbally. Art. 24 of Bill 125, supra, note 186, provides likewise.

248. OLRCC, supra, note 2 at 62.

249. See Uniform Human Tissue Gift Act, 1971, supra, note 236, s. 10. It should be borne in mind that most of the provinces patterned their legislation after this model. Section 10 was amended as follows, omitting the notion of public policy (Uniform Human Tissue Donation Act, 1989, supra, note 236):

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The scope of this prohibition is not entirely clear, however. On the one hand, the definition of "tissue" excludes tissues that are capable of regeneration, which suggests that they are exempt from the prohibition. On the other hand, the language of the provision seems to indicate the intention also to include tissues that are capable of regeneration: "No person shall buy, sell . . . any tissue for a transplant, or any body or part or parts thereof other than blood or a blood constituent [emphasis added]."

In Quebec, article 20 C.C.L.C. permits the sale of parts of the body that are capable of regeneration. We may therefore conclude that if gametes are considered as tissues or parts of the body capable of regeneration, donations would not necessarily have to be gratuitous.

However, the nature of gametes leads us to consider the appropriateness of prohibiting all commercialization of gametes, permitting only the reimbursement of expenses and limiting the circulation and storage of gametes to hospitals and non-profit fertility clinics. Similarly, we must consider the need to regulate the import of eggs and sperm. We must also ask the same questions with regard to embryo donation.

15. (1) No person shall buy, sell or otherwise deal in, directly or indirectly, any tissue, body or body part for the purpose of a transplant or for a therapeutic purpose, medical education or scientific research.

(2) Any dealing in any tissue, body or body part that was lawful before this Act came into force shall continue to be lawful, provided this Act is complied with.

(3) A person who contravenes this section is guilty of an offence and liable on summary conviction to a fine of not more than $100,000 or to imprisonment for not more than 1 year, or to both.

As we saw supra, the definition of "tissue" in this latest version of the Uniform Human Tissue Donation Act, supra, note 236, does not include spermatozoa, ova, embryos or fetuses.

250. Ibid. For more details on this matter, see the forthcoming LRC working paper Procurement and Transfer of Human Tissues and Organs.

251. An Act to add the reformed law of persons, successions and property to the Civil Code of Québec, supra, note 235, art. 22, provides: "The alienation of a part of the human body not capable of regeneration shall be gratuitous." Therefore, the sale of a part of the body that is capable of regeneration should be permitted. See Ouellette, supra, note 235 at 21-22. However, subarticle 22(2), which is new law, prevents exploitation by stating that "[t]he alienation of a part of one's body shall not be repeated if it involves a risk to the health." However, art. 25 of Bill 125, supra, note 198, drops the distinction for parts of the body that are capable of regeneration. See supra, note 210.

252. See supra at 41ff and notes 207 and 208.

253. Section 11.5 of An Act to amend the Uniform Child Status Act, supra, note 199, provides as follows:

(1) No person shall, directly or indirectly, buy, sell or otherwise deal in human eggs, sperm or embryos.

(2) A person who contravenes this section is guilty of an offence and liable on summary conviction to a fine of not more than $100,000, to imprisonment for not more than one year or to both.

(3) This section does not prohibit a person from giving or receiving reimbursement for reasonable expenses necessarily incurred in donating her own eggs or his own sperm.
Another characteristic of gamete and embryo donation is that it is normally made on the condition, implied or express, that the donor remain anonymous. Donors must therefore be assured adequate protection against disclosure of their identity.

If the donor is considered a "patient," he or she is protected, under both the civil law and the common law, by the rules of confidentiality that normally apply to patient-physician communications. Physicians have always been required, either by medical ethics or by the law, not to disclose the medical information in their patients' records. It is clear law that a physician owes a duty of confidence to his or her patient. This duty is recognized at common law, in at least two provinces by legislation and may even be constitutionally guaranteed.

The two legal systems also share a rule whereby information may not be disclosed unless the patient gives his or her consent, or unless disclosure is required by law or as a matter of public policy.

Confidential information may also be disclosed in a court of law. However, the risk of such disclosure seems to be greater at common law because physicians do not enjoy any privileges as witnesses, whereas in the civil law a physician is bound to maintain confidentiality in a legal proceeding unless the patient releases him or her from that duty.

Given the special status of donors, we might ask whether it is appropriate to make them subject to the same legal scheme that applies to patients. Moreover, since gamete and embryo donations are made within the very specific framework of medically assisted

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254. Rule 5 of the Canadian Medical Association, Code of Ethics (Ottawa: The Association, 1990) reads: "An Ethical Physician... will keep in confidence information derived from a patient or from a colleague regarding a patient, and divulge it only with the permission of the patient except when otherwise required by law."


257. Casswell, supra, note 255 at 231.

258. Knoppers, supra, note 196 at 125; see also Hals v. Mitchell, supra, note 255 at 136.

259. See Medical Act, supra, note 255, s. 42, and s. 9 of the Quebec Charter.

260. There is uncertainty both in Quebec and in the common law provinces about considering donors as patients. See the OLRC report, supra, note 2 at 83; and Butha Maria Knoppers, "Vérité et information de la personne" (1987) 18 R.G.D. 819 at 830.
procreation, we cannot ignore the need for information of the other persons involved. In light of these factors, it may be reasonable to consider the establishment of another scheme that would protect the donor’s privacy and at the same time guarantee access to the medical, genetic and social information required by the other persons involved.

3. Donor Consent

As stated earlier, because the human body is inviolable, the donation of sperm, ova and embryos requires the free and informed consent of the donor. In Quebec, article 20 C.C.L.C. states that such consent must be in writing. The same requirement exists in the other provinces that adopted section 3 of the Uniform Human Tissue Gift Act. Further, in both civil and common law the risk incurred must not be disproportionate to the anticipated benefit.

Ovum donation, however, poses a special problem. In cases where consent was given for a procedure requiring egg retrieval (for any reason), it is not certain under the law as it now stands whether it is necessary to obtain the woman’s consent to dispose of her eggs if she does not specify the purpose for which they are intended:

Ova may be obtained ... for instance at a woman’s sterilization, investigation for subfertility, or hysterectomy. Further, where superovulation is stimulated to assist a woman to become pregnant by I.V.F., any surplus ova may be available to others ... It is not clear, however, that such sources of ova for I.V.F. of other women are legally required to give consent. If they take initiatives to control recovered ova, their wishes must be respected.

261. See infra at 157-60.
262. See supra at 46ff.
263. Supra, note 236. It should be noted, however, that the latest version of the Act available for adoption by the provinces does not require written consent. See Uniform Human Tissue Donation Act, supra, note 236, s. 5(4).
264. Bernard M. Dickens, ‘“Reproduction Law and Medical Consent”’ (1985) 35 U.T.L.J. 255 at 283, refers to Venner v. Maryland, 354 A.2d 483 at 499 (1976), in which the court ruled that “when a person does nothing and says nothing to indicate an intent to assert his right of ownership, possession, or control over such material, the only rational inference is that he intends to abandon the material.” See also OLRC, supra, note 2 at 89.

Assuming that a property right or a right of possession rests in the producer of the human genetic material — and, therefore, assuming that such rights in the material do not automatically pass to the hospital or physician storing or working with it — questions still arise concerning, for example, whether these rights may be lost merely by a failure to assert them or whether, notwithstanding the absence of specific directions, the producer of the ova or semen may correctly assume that the hospital or physician is under a legal duty to destroy the unused material, with perhaps some routine examination by a pathologist, but not experimentation.
Despite the existence of more general consent, the special nature of ova and the significance of the donation for the woman should justify the need for specific donor consent.

Consent to embryo donation raises other problems. Whether the embryo is derived from the gametes of the two spouses or was conceived using gametes from only one of the spouses and a donor, we may ask whose consent is required. Finally, as discussed earlier, the revocability of consent respecting a donation raises a number of issues for the common law provinces.

4. Donor Liability

A donor who provides false information or fails to disclose information about his or her medical history can endanger the child and the mother. If the donor conceals the fact that he or she carries a genetically transmissible disease and the child born using the gametes he or she donated is affected, the donor may be held liable. The donor would thus have a duty to disclose.

But who would be able to take action against the donor? In the civil law, a child that is conceived but not yet born enjoys some legal recognition, conditional on its birth. At common law, an action for "wrongful birth" is also open to the parents and the child.

265 Knoppers, supra, note 196 at 99-100.

266 See supra at 51.


268 Baudouin and Labrousse-Riou, supra, note 210 at 57; Knoppers, supra, note 267 at 16-17.

The protection of the unborn from negligent injury under the common law was first established with regard to its proprietary and successorship interests from conception onwards, provided the child was born alive. . . . Similarly, the common law in the United States and Canada has either adopted the notion of a pre-existing duty not to harm, or a conditional prospective duty, which crystallizes at birth as the basis for tortious liability. Another recent common law notion is the causal approach, which separates the concepts of injury and damage and looks simply at the causal link between the infant's condition at birth and the defendant's wrongful conduct, thus avoiding the issue of legal personality and the moment of injury altogether.
However, donor anonymity, evidentiary problems and the difficulty of establishing a causal link would make such legal action virtually impossible. Moreover, a liability suit could more easily be taken against the gamete and embryo bank for its failure in the selection and screening phases, or against the physician in charge.

Nevertheless, it should be noted that there is a public-order dimension to donor liability. A donor could be held criminally liable if, for example, he knew he was carrying a potentially fatal virus, still donated his sperm and deliberately withheld the information. It is therefore important that donors at least be identifiably and that their identity be revealed in the event of criminal prosecution for failure to disclose information.

C. Children

1. Legal Parentage

Parentage law organizes the legal relationships between a child and his mother and father. Parentage gives rise to certain rights and obligations, such as the obligation of

Dickens, supra, note 221 at 262:

Liability for pre-conception tests has recently been recognized under Common Law reasoning, so it may not matter whether the ovum was fertilized or unfertilized when damage occurred. Recognition of pre-conception tests has emerged only recently, however, and upon the basis of United States decisions, so that their status in Canada is at present unavoidably unclear.

See Jorgensen v. Mosaic Johnson Laboratories, Inc., 483 F. 2d 237 at 240 (1973), in which Judge Holloway ruled:

If the view prevailed that tortious conduct occurring prior to conception is not actionable in behalf of an infant ultimately injured by the wrong, then an infant suffering personal injury from a defective food product, manufactured before his conception, would be without remedy. Such reasoning runs counter to the various principles of recovery which Oklahoma recognizes for those ultimately suffering injuries proximately caused by a defective product or instrumentality manufactured and placed on the market by the defendant. . . .

We are persuaded that the Oklahoma courts would treat the problem of the injuries alleged here as one of causation and proximate cause, to be determined by competent medical proof. . . . And such treatment of the problem would accord with the predominant view that an action may be maintained for prenatal injuries negligently inflicted if the injured child is born alive.

See also Renslow v. Mennonite Hospital, 351 N.E. 2d 870 (1976).


270. R. v. Thorston (15 June 1989), Ottawa-Carleton 1814 (Ont. Dist. Ct. aff’d (1991) 1 O.R. 3d 480 (C.A.)). This very recent ruling by the Ottawa District Court found an individual guilty of public nuisance because he voluntarily donated his blood to the Red Cross even though he knew he was carrying the AIDS virus.


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parents to feed, care for and educate their children. Legal parentage also has an impact on the law of successions. The legal bond can be established by the biological link (parentage by blood) or by an act of will (adoption). How then is the parentage of children born as a result of medically assisted procreation to be determined?

Traditionally, the establishment of maternal filiation through the fact of childbirth reflected a biological and genetic certainty. Marriage, a social reality, made it possible to resolve the uncertainty of paternity by a presumption which, while favouring the social aspect of paternity, usually reflected a biological reality.

Some legislatures have over the years eased the traditional rules governing filiation by abolishing the distinction between illegitimate children and legitimate children and recognizing the predominance of biological truth.

In the common law provinces, however, there appears to be disparity between those that establish parentage solely on the basis of marriage and those that have passed laws to ease this common law rule by abolishing the difference between illegitimate and legitimate children.

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273. With respect to the civil law see D. Castelli, *supra,* note 271 at 122; with respect to the common law see Margo Wilson, “Impact of the Uncertainty of Paternity on Family Law” (1987) 45 U.T. Fac. L. Rev. 216 at 232. See *infra,* note 283.


Subsection 56(1) of the *Law and Equity Act,* R.S.B.C. 1979, c. 224, as am. S.B.C. 1985, c. 68, s. 80, reads: Subject to the *Adoption Act* and *Family Relations Act,* for all purposes of the law of British Columbia,

(a) a person is the child of his natural parents,

(b) any distinction between the status of a child born inside marriage and a child born outside marriage is abolished, and

(c) the relationship of parent and child and kindred relationships flowing from that relationship shall be determined in accordance with this subsection.
It should be noted, however, that these changes have sparked controversy, both in Quebec and in the common law provinces, regarding the priority to be given to the various means of proving parentage.275

It is questionable whether the legislatures truly intended to choose between the biological and social aspects of parentage. Indeed, whether they established the certainty of maternity and the presumption of paternity or did away with the difference between legitimate and illegitimate children, was their objective to preserve the child’s interest, or were they more concerned about ensuring a clearer application of the law (better administration of justice)?

While it is as difficult to determine what the current rules on parentage should reflect (social likelihood or biological truth) as it is to identify legislative intent, applying and adapting the rules to medically assisted procreation is becoming especially problematic. Problems that may arise include: the attribution of the responsibilities of fatherhood to a husband who did not consent to the conception of the child (or to a donor who had absolutely no intention of being a father276; disavowal of a child originally wanted277 by

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This suggests that in the provinces where this traditional distinction has been abolished, the law would still conceal the biological truth under the presumption of legitimacy related to marriage and would thus favour a social truth. However, the British Columbia Supreme Court ruled as follows in *B. (B.J.) v. K. (J.)* (1989), 56 D.L.R. (4th) 150 at 158.

While it is correct to say that s. 56 of the *Law and Equity Act* has, by abolishing “any distinction between the status of a child born inside marriage and a child born outside marriage” abolished the status of illegitimacy... it certainly has not abolished the reality of what we know as “legitimacy” and “illegitimacy”... ...

In my opinion, it would require a clear and unambiguous expression of intention by the legislature to displace such a longstanding presumption, which, in my view, provides a just and useful rule... ...

As no such clear and unambiguous expression appears in s. 56 of the *Law and Equity Act*, I hold that the presumption of legitimacy remains in effect in British Columbia.


276. See, e.g., s. 56(1(a) of the *Law and Equity Act*, supra, note 274.

277. Subject to art. 586 C.Q. in Quebec; s. 13(3) of the *Children’s Act*, supra, note 197, in the Yukon; and s. 12(3) of *The Children’s Law Act*, supra, note 197, in Newfoundland.

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parents who change their mind at some point during the procedure; and the possibility of a challenge of paternity by third parties or the donor and a claim of paternity by the donor. 278

Application of the current rules governing parentage to technologies that use donated ova has had the effect, for the first time, of dividing maternal biological filiation into genetic filiation and gestational filiation. This situation has faced lawmakers with a choice that they could not have foreseen: Should gestation and delivery prevail over genetic link?

2. Biological Parentage

Children may wish to trace their origin, either to obtain the medical histories of their forebears or to satisfy a psychological need to establish their identity. We can safely say that a child’s interest in knowing about his or her medical and genetic history meets with little objection in our society.279 In practice, however, access to such information is not guaranteed:

A major inadequacy of present legislative regimes is that, even when they accommodate the preferences of active participants in artificial insemination, as in the case of A.I.D. [artificial insemination with donor], they do not necessarily protect the children consequently

278. See Dickens, supra, note 218 at 68; Canadian Bar Association, Report of the Special Task Force Committee on Reproductive Technology of the British Columbia Branch, 1989 at 13 [unpublished]. See arts 586 and 588 C.C.Q. and art. 580 of Bill 125, supra, note 196. The Unon has provided for the absence of a legal relationship between a donor and a child born of the product of the donation if the donor is not the husband of the mother. Children’s Act, supra, note 197, s. 13(6); In An Act to amend the Uniform Child Status Act, supra, note 199, s. 14(2) provides that: "A man whose sperm is used in an assisted conception and who is not presumed to be the father of a child pursuant to section 9 is deemed not to be the father of the child." Art. 579 of Bill 125, supra, note 196, is similar:

Participation in the parental project of another person by way of a contribution of genetic material to medically assisted procreation does not allow the creation of any bond of filiation between the contributor and the child born of that procreation.

279. Baudouin and Labrusselle-Riou, supra, note 210 at 55; and Bernard M. Dickens, “Legislating for the Brave New Children” in Barbara Landau, ed., Children’s Rights in the Practice of Family Law (Toronto: Carswell, 1986) 345 at 347. To ensure that records are kept on the genetic origin of children born as a result of medically assisted procreation and that access to those records is made possible, s. 11.6 of An Act to amend the Uniform Child Status Act, supra, note 199, provides that:
born. . . . A larger problem concerns children born of donated sperm and/or ova, whose medical care and later reproductive counselling may be dependent upon knowledge of their genetic parentage. The absence of means of tracing at least genetic profiles of biological parents may place them at a disadvantage and perhaps at risk. The adoption model may expose children to disadvantage, but it usually permits discovery of at least a birth mother’s characteristics; the practice of birth through donated sperm or ova reveals a default of legislative attention that exposes an increasing number of children to the risk of grave disadvantage to health.280

Further, access to this information, assuming it were available, raises the problem of disclosure of the use of a donation from a third party to the conception. We therefore have to decide whether such disclosure should be left to the parents’ discretion or whether the child should be recognized as having a right to the information upon reaching the age of majority.

The notion of the child wanting information because of a psychological need to establish his or her identity is not unfamiliar. It is being recognized more and more in the context of adoption. Systems have been put into place to enable adopted children to search for

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280. Dickens, supra, note 279 at 355; see also Knoppers, supra, note 267 at 15 n. 64.
their biological parents. In the area of medically assisted procreation, however, because of the confusion over record keeping and the "protection" afforded the donor's anonymity, children are generally denied the right to know about their origins.

We must first ask whether it is appropriate to compare the situation of children born as a result of a donation with that of adopted children and then question the appropriateness of granting them the right to know about their origins or certain rights to information. In any event, it is important to find a middle ground so that the privacy of the donor and the parents is protected. Should no formal right to information be granted, we would anticipate cases in which anonymity might be lifted.

3. Legal Status

Most provinces and territories in Canada have done away with the distinction between legitimate and illegitimate children, but some, such as Alberta and Nova Scotia, have not.

In these provinces, children conceived by use of donated gametes or embryos will be denied the benefits enjoyed by legitimate children, not only in cases where the parents are not married, but also in cases where it is proved that the child was conceived as a

281. In Quebec, e.g., a summary of the child's history may be delivered upon request to an adoptive parent or to the child if he or she is aged 16 years or older, provided anonymity is respected (Youth Protection Act, R.S.Q., c. P-34.1, ss 131.1 and 131.2). Further, the legislature has provided for the reunion of biological parents and adoptees who have reached the age of majority (art. 632 C.C.Q.). The search takes into account the intentions of the parties. See, e.g., art. 632 C.C.Q. (referred to in art. 577 of Bill 125, supra, note 196) and, in Ontario, Ministry of Community and Social Services, Adoption Disclosure Services (Toronto: Queen's Printer for Ontario, 1987). It should be noted, however, that in Quebec, art. 583 of Bill 125, supra, note 196, provides for the confidentiality of identifying information about those involved in the medically assisted procreation of a child. An exception is made where confidentiality could cause grave injury to the child's health.

282. According to a study in the United States, barely one-third of the physicians interviewed kept permanent records on children conceived by artificial insemination, and fewer than one-third kept permanent records on donors. See Martin Currie-Cohen, Lesleigh Luttrel and Sander Shapiro, "Current Practice of Artificial Insemination by Donor in the United States" (1979) 300:11 New Engl. J. Med. 585 at 588, quoted in Ann T. Lampert, "The Genetics of Secrecy in Adoption, Artificial Insemination, and In Vitro Fertilization" (1983) 14:1 Am. J. L. Med. 109 at 116-17, particularly at 118: "Often the semen used in artificial insemination is collected by a urologist and the insemination [is] done by an obstetrician who may not actually deliver the child."

283. The Civil Code of Quebec has removed the existing inequities between the different known types of filiation: legitimate, natural and adoptive. See art. 594 C.C.Q. (referred to in art. 536 of Bill 125, supra, note 196). See also Law and Equity Act (B.C.), supra, note 274, s. 56(1); Child Status Act, R.S.P.E.I. 1988, c. C-6, s. 1(1) and (4); Family Services Act, S.N.B. 1980, c. F-2.2, s. 96(1) and (4); The Family Maintenance Act, R.S.M. 1987, c. F.20, s. 17; Children's Law Reform Act (Ont.), supra, note 228, ss 3(1) and (4) and 2(1); Judicature Act, R.S.N.W.T. 1988, c. J-1; Children's Act ( Yukon), supra, note 197, s. 5(1) and (4); The Children's Law Act (Nfld), supra, note 197, s. 3; The Children's Law Act, S.S. 1990, c. C-8.1, s. 40; and Uniform Child Status Act, supra, note 199, s. 2.
result of a donation (despite the fact that artificial insemination with donor is no longer considered adultery). Even though the presumption of paternity covers AID by making the husband the father of the child, there are questions to be asked regarding the appropriateness of clarifying the legal status of children born as a result of gamete or embryo donation:

[W]here the distinction exists between legitimacy and illegitimacy, the children of proven A.I.D. [artificial insemination with donor] are held to be illegitimate, with all of the legal disadvantages they bear in their social families due to that status. The fact of A.I.D. is frequently concealed, because children born to married women are legally presumed to be their husbands', and no one has an interest to rebut that presumption.

D. Medical Personnel

1. Liability of Physicians

A physician incurs liability when he or she acts in a negligent manner in administering treatment. However, physicians are held only to an obligation of means — or general duty of prudence and diligence —, not to an obligation of result — or absolute duty.

In medically assisted procreation, negligence on the part of the physician may occur before or after conception. Negligence may pertain to the administration of the procedure used, the performance of the duty to inform or the respect of the duty of confidentiality. For the purposes of our study, we will look specifically at the duty to inform, which stems from the duty of every physician to obtain the free and informed consent of his or her patient.

The scope of this obligation to inform appears to vary depending on whether the treatment is therapeutic, elective or experimental. Although the Supreme Court did not


285. Dickens, supra, note 279 at 347.

286. Baudouin and Labrèse-Riou, supra, note 210 at 57.

287. Ibid.; Picard, supra, note 196 at 670.

288. Baudouin and Labrèse-Riou, supra, note 210 at 58; Sharpe, supra, note 256 at 181 ff. This matter has already been discussed; see supra at 53.

289. The requirement of consent flows from application of the rule, protected by the civil law, the common law and the criminal law, that a person's physical integrity may not be violated without the person's consent. With respect to the criminal law, see L.R.C., Revisifying Criminal Law: Revised and Enlarged Edition of Report 30, Report 31 (Ottawa: The Commission, 1987) at 61.

290. Picard, supra, note 196 at 92.
comment specifically in *Reibl v. Hughes* or *Hopp v. Lepp* on the scope of the duty to inform in cases of elective surgery, other decisions, both in the civil law and the common law, interpret the obligation more broadly.

In the field of research, the obligation to disclose information and risks would be even stricter. “Not only must the research subject consent but his or her consent must be explicit and based on what might well be called a ‘perfect’ disclosure.”

The problem in medically assisted procreation is determining whether the different technologies that are used are therapeutic, elective or experimental treatments and, consequently, what standard of disclosure should be required.

Although it is not within the scope of our study to decide this matter, we feel it would be desirable for our courts, in considering this issue, to take into account the specific characteristics of each technology (risks, success rates, and so on).

2. Liability of Gamete and Embryo Banks

Donor selection is vital because it not only increases the chances of success, but also protects the health of the person receiving the donation and prevents the transmission of serious infectious or genetic diseases. Since it was discovered that the AIDS virus can be transmitted through sperm, physicians have begun using frozen sperm for insemination.

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291. *Supra*, note 238.

292. With respect to the common law, see Picard, *supra*, note 196 at 93:

> In interpreting the Supreme Court of Canada’s decision in *Reibl v. Hughes* (*Supra*, note 238), the provincial courts have said that for an elective procedure, minimal or possible risks and alternative procedures and their comparative risks must be explained voluntarily... There is authority for requiring a doctor effecting a sterilization to explain other methods or techniques. It would seem that this can be generalized to cover any elective procedure.


293. Picard, *supra*, note 196 at 118. See also *Biomedical Experimentation Involving Human Subjects*, *supra*, note 7 at 301. In Canada, the decision in *Haluchuk v. University of Saskatchewan* (1985), 53 D.L.R. (4th) 416 (Sask. C.A.), sets the standard and scope of the duty to disclose in the area of experimentation. See at 464: “The subject of medical experimentation is entitled to a full and frank disclosure of all the facts, probabilities and opinions which a reasonable man might be expected to consider before giving his consent.” There can be no exceptions to the ordinary requirements of disclosure in the case of research as there may well be in ordinary medical practice.”

See also the recent decision by the Quebec Superior Court in *Weiss v. Solomon*, [1989] R.J.Q. 731.

294. See chap. 1.

295. Lori B. Andrews, *Medical Genetics: A Legal Frontier* (Chicago: American Bar Foundation, 1987) at 168: “[S]ome women who have been inseminated with donor sperm have contracted venereal disease from the sperm. In addition, children have been born with genetic defects that were passed on by the sperm donor.”
Gamete and embryo banks may be liable if they are negligent in screening for genetic defects and diseases. In the civil law, "the laboratory is held to an obligation of result and in the case of error will have to exculpate itself by demonstrating some external force or event." In the common law, "since the standard of care is proportionate to the risks involved, the standard of care could very well be the same." 296

Unlike donations of other human products, such as blood, the donation of gametes and embryos is not currently subject to any national regulatory scheme. In 1977, an advisory committee examined the question of the storage and use of human sperm; in 1981, the committee submitted a report to the Minister of National Health and Welfare. One of the committee's recommendations was that "Federals regulations governing standards for the acquisition, preservation and importation of human sperm be established." 299 In 1988, the Canadian Fertility and Andrology Society adopted guidelines dealing, among other things, with the selection of donors and genetic screening. However, as stated in the preface, the document was "not intended to be exhaustive, nor to replace any other guidelines or be considered as a rigid set of procedures and standards." 300

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296. Knoppers, supra, note 267 at 10 n. 47.

297. Ibid.

298. The Food and Drugs Act, R.S.C. 1985, c. F-27 and the Food and Drug Regulations (C.R.C. 1978, c. 870) have set standards respecting, inter alia, advertising, labeling, sale, import, processing, storage and the number of donations permitted. The Canadian Red Cross Society also has standards respecting some of these subjects; see Procurement and Transfer of Human Tissues and Organs, supra, note 250.


300. Canadian Fertility and Andrology Society, supra, note 11 at 3; current practice is therefore fraught with uncertainty respecting the application of uniform criteria. The OLRC wrote in 1985:

Practices respecting donor screening appear to vary considerably. The most common tests, which are either given by the doctors or required by them to be done in a laboratory, are blood group and type, semen analysis and culture, and VDRL [Venereal Disease Research Laboratory]. Less frequent tests are CBC [Complete Blood Count] and hepatitis, with still fewer responses reporting karyotype [sic], genetic screening, and the taking of a family or general history of the donor. Other tests were listed by some practitioners.
The rapid development of medicine makes it difficult to regulate medical selection and the prescription of genetic screening tests. On the other hand, the tremendous uncertainty regarding the uniform application of selection criteria and storage and import standards poses a risk for the unborn child and the future parents. It is therefore important to determine who should be responsible for ensuring the uniformity of the standards used and how this should be carried out. Could the medical profession alone take on this task?

We should also point out that gamete and embryo banks have a duty to respect the donor’s consent and may be held liable if they show negligence in the storage of gametes and embryos.

We have seen that many rules of law apply to medically assisted procreation, but in general few are adapted to this new reality. In the chapter listing our recommendations, we will endeavour to address the problems identified in this first section. We will now examine the specific problems posed by surrogate motherhood.

II. Surrogate Motherhood

A. Issues

1. Legality and Legitimacy

As the law now stands, surrogacy runs counter to the principles of contract law. For both the civil and the common law, any contract or agreement, even where there is no payment, between a surrogate and prospective parents is quite probably null and void as

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OLRC, supra, note 2 at 22.

Further, there are doubts as to the ability of practitioners to carry out appropriate genetic screening of donors. See Lampert, supra, note 282 at 117. The author refers to Currin-Cohen, Lar nell and Shapiro, supra, note 282 at 586.

Geneticists and others recognize that there is a serious lack of knowledge of genetics in those who perform artificial insemination. Donors are commonly screened for hereditary disorders, but the screening is illusory. Family histories are taken, but they are usually superficial. Biochemical testing is only performed in about 28.8% of the cases. . . . The doctors participating in the Currin-Cohen study, 92% said that they would reject a donor with a chromosomal translocation or trisomy, but only 12.5% actually examined the potential donor’s karyotype to see if the chromosomes were abnormal. Also, 71.4% said that they would reject a donor who had hemophilia in his family, despite the fact that it is an X-linked trait and would be impossible for a man to transmit unless he carried the gene and exhibited the trait himself.

Lampert writes, ""To date, this is the only study of its kind."" See also Barratt, Chablan and Cooke, supra, note 161; and "Screening Gametes and Embryo Donations," supra at 32.

301. See supra, note 231.
being against public policy.\footnote{302}{Moreover, such contracts are also at odds with a fundamental principle of family law: the custody of a child must be determined according to the best interests of the child rather than the wishes expressed by the parents in a contract.\footnote{303}{Clear, the absolute nullity of such an agreement does not prevent the parties from giving effect to a contract where the surrogate does not object to surrendering the child.\footnote{304}{302} With respect to the common law, see Dickson, \textit{supra}, note 218 at 71: "It is commonly accepted that, in the absence of approving legislation, surrogate motherhood agreements will be held void by the courts as against public policy." OLRC, \textit{supra}, note 2 at 220. "Although not otherwise prohibited, it would appear that such arrangements are illegal and unenforceable at common law as being against public policy." With respect to the civil law, see Baudouin and Labranque-Rieu, \textit{supra}, note 210 at 115; art. 582 of Bill 125, \textit{supra}, note 196, is explicit: "Procreation or gestation agreements on behalf of another person are null." Recall also, the U.S. decision in \textit{Baby M.}, 537 A. 2d 227 at 1234 (N.J. 1988): "We invalidate the surrogate contract because it conflicts with the law and public policy of this State."}

\footnote{303}{303} The precedence of the best interests of the child over contractual freedom is made clear in such provisions as ss 52(1)(c), 53(1)(c) and 56(1) of Ontario's \textit{Family Law Act, 1986}, \textit{supra}, note 196. For example, s. 52(1)(c) reads:

\begin{quote}
52(1) A man and a woman who are married to each other or intend to marry may enter into an agreement in which they agree on their respective rights and obligations under the marriage or on separation, on the continuance or dissolution of the marriage or on death, including,

\begin{itemize}
\item the right to direct the education and moral training of their children, but not the right to custody of or access to their children; . . .
\end{itemize}
\end{quote}

Paragraph 53(1)(c) provides likewise for persons "who are cohabiting or intend to cohabit." See also s. 56(1):

\begin{quote}
"In the determination of a matter respecting the support, education, moral training or custody of or access to a child, the court may disregard any provision of a domestic contract pertaining to the matter where, in the opinion of the court, to do so is in the best interests of the child."
\end{quote}

\footnote{304}{With respect to the common law, see the OLRC report, \textit{supra}, note 2 at 99: "With respect to the common law . . . the courts have long held that, subject to very few exceptions, parental rights and responsibilities are inalienable and incapable of transfer as a matter of contract." The illegality of such an agreement would be determined primarily by the interests of the child. Thus the Supreme Court has recognized that some custody agreements, the main objective of which were the best interests of the child, were not illegal; see \textit{Chisholm v. Chisholm} (1908), 40 S. C. R. 115. \textit{A contrario}, see \textit{Re Hutchinson} (1913), 28 O.L.R. 114 (C.A.). See the discussion of surrogate contracts and transfer of custody of the child in the OLRC report, \textit{supra}, note 2 at 94-102. With respect to the civil law, see Rivet, \textit{supra}, note 275 at 850: [\textbf{TRANSLATION}]


\footnote{305}{OLRC, \textit{supra}, note 2 at 99-100:}

\begin{quote}
While a surrogate motherhood agreement may not be enforced as a matter of contract law, the existing legal regime does not make it completely impossible to give effect to the wishes of
We should point out, however, that this is made possible only through the application of rules governing filiation in both the civil law and the common law.\textsuperscript{306}

Since the current law does not sanction surrogate motherhood and indirectly even permits it, we must ask if it would be appropriate to alter the situation. If so, legislation could be introduced that would make surrogacy contracts legal and set out the terms and conditions governing them, or would specifically prohibit such contracts as being contrary to public policy.

[Translation]
It may be that culture and traditional family law are drastically altered by the notion of surrogate motherhood, but surrogacy should not necessarily be seen as something negative to be restricted or prohibited. The mere fact of surrogacy means that a choice must be made between confirmation and prohibition in positive law.\textsuperscript{307}

2. Commercial Aspects of Surrogacy

The monetary aspect of a surrogacy contract raises not only the potential for exploitation of the parties, but also the prospect of trade in children, which is currently prohibited in Canada by adoption and youth protection legislation:

Though each Canadian province has its own legislation governing adoption, the basic statutory framework is similar throughout the country. The legislation restricts who may adopt a child and who may be adopted. Throughout North America concern exists about a practice sometimes known as baby farming, the unscrupulous placement of babies for adoption by operators motivated by a desire for profit and invariably acting with little regard for the welfare of the child. As a result, legislation restricts who may arrange adoptions and how they are to be arranged; in particular, there are restrictions about receiving payment for placing a child or doing other work in connection with an adoption.\textsuperscript{308}

\begin{footnotesize}
\footnote{the parties to such an agreement. There are procedures by which a child, born or about to be born to a surrogate mother, might be "naturalized" in the care and custody of the prospective social parents, at least where the social father is also the biological father. The focus of attention here is not on the validity or enforceability of a surrogate motherhood agreement, but on the steps that may be taken today where a child is born or about to be born and the surrogate mother is willing to transfer the child. See also Baudouin and Labouesse-Riou, supra, note 210 at 127-28; Rivet, supra, note 275 at 850-51, regarding the use of adoption rules in Quebec.}
\footnote{See infra at 69ff.}
\footnote{Baudouin and Labouesse-Riou, supra, note 210 at 120.}
\footnote{Nicholas Bala, Heino Lilles and Georges Thomson, Canadian Children's Law (Toronto: Butterworths, 1982) at 294-85.}
\end{footnotesize}
For example, subsection 33(2) of The Adoption Act of Saskatchewan reads as follows:

(2) [No person shall:
(a) give or receive; or
(b) agree to give or receive;
any payment or reward, whether directly or indirectly, for any purpose related to the adoption of a child.\textsuperscript{309}

Since it is illegal to receive payment in return for arranging the adoption of a child, these provisions may also prohibit a surrogate who consents to the adoption of her child from being reimbursed for expenses.\textsuperscript{310}

\textsuperscript{309} The Adoption Act, S.S. 1989-90, c. A-5.1. In Quebec, see s. 135.1 of the Youth Protection Act, supra, note 281:

Whether the placement or the adoption takes place in Quebec or elsewhere and whether or not the child is domiciled in Quebec, any person who
(a) gives or receives or agrees to give or receive, directly or indirectly, a payment or a benefit either for finding a placement or contributing to a placement with a view to adoption, or for obtaining the adoption of a child, . . .
is guilty of an offence and liable, on summary proceedings, in addition to costs, to a fine of $2,000 to $5,000, in the case of an individual, and to a fine of $5,000 to $10,000, in the case of a corporation.

This provision does not, however, invalidate the placement or adoption. See also Child Welfare Act, S.A. 1984, c. C-8.1, s. 71, as am. S.A. 1988, c. 15, s. 35; Family Services Act, (N.B.), supra, note 283, s. 95; The Child and Family Services Act, S.M. 1985-86, c. 8, ss 63 and 85; The Adoption of Children Act, 1972, S.N. 1972, No. 36, s. 5, as am. S.N. 1974, No. 9, s. 3 and S.N. 1979, c. 35, Sch. A, Item 1; Child Welfare Act, R.S.B.C. 1979, c. C-6, s. 108; Children and Family Services Act, S.N.S. 1990, c. 5, s. 89(3); Adoption Act, R.S.P.E.I. 1988, c. A-4, s. 23, Children's Act (Yukon), supra, note 197, s. 102. Ontario and British Columbia permit payment under certain conditions; see Child and Family Services Act, 1984, S.O. 1984, c. 55, ss 159-160, as am. S.O. 1987, c. 4, s. 8, and S.O. 1989, c. 72, s. 20, and Adoption Act, R.S.B.C. 1979, c. 4, as am. S.B.C. 1980, c. 36, s. 2. A decision by the British Columbia Supreme Court allowed the natural mother to be reimbursed by the adoptive couple for reasonable expenses related to the adoption of her child. Justice Hudspeth specified, however, "On another day and in other circumstances another judge might have different criteria but the adopting parents have established to my satisfaction that my criteria for approving a payment to the natural mother have been met."

Re Adoption Act (1982), 27 R.F.L. (2d) 72 at 75.

\textsuperscript{310} Dickens, supra, note 218 at 71: "Known participants complying with their terms in Canada have not been subjected to legal proceedings, for instance for violation of prohibitions against offering and receiving money for consent to adoption."

It should be noted, however, that adoption laws in all provinces except Manitoba\(^{311}\) are aimed solely at transactions intended to result in the adoption of a child. Any other transaction that does not constitute an adoption would therefore not be subject to these laws.

In the *Criminal Code* there are currently no provisions that specifically prohibit traffic in and the sale and purchase of children. On the other hand, the offences referred to in sections 279(1) (kidnapping), 279.1 (hostage taking), 280 (abduction of person under sixteen), 281 (abduction of person under fourteen), 282 (abduction in contravention of custody order) and 283 (abduction where no custody order) are not appropriate charges in all situations involving surrogate motherhood.\(^{312}\)

**B. Children**

1. **Legal Parentage**

   We have already outlined the problems the various technologies of medically assisted procreation pose for parentage law.\(^{313}\) We must now consider the specific problem of the parentage of children born to a surrogate.

   The parentage of a child born to a surrogate can depend on a number of factors, among them whether or not the woman voluntarily surrenders the child; whether the woman is married or single; and whether the surrogate provides the ovum or is only the gestational mother.\(^{314}\)

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311. Unlike the other provinces, Manitoba does not limit the offence to adoption. Section 84 of *The Child and Family Services Act*, supra, note 309 reads as follows:
   Any person who gives or receives or agrees to give or to receive any payment or reward either directly or indirectly in consideration for
   (a) the purported sale of a child for any purpose; or
   (b) procuring or assisting in procuring the purported sale of a child for any purpose;
   is guilty of an offence punishable on summary conviction and liable to a fine of not less than $1,000.00 and not more than $10,000.00 or to imprisonment for a term not exceeding 6 months or both.


313. *Supra* at 36ff.

If an unmarried surrogate surrenders the child at birth, she will nevertheless be deemed to be the legal mother but she may also consent to adoption by the father’s wife.\textsuperscript{315} If she does not so consent, a motion may in theory be filed seeking deprivation of the surrogate’s parental authority for having abandoned the child.

If an unmarried surrogate refuses to surrender the child, the social father, if he is also the child’s biological father, may claim paternity.\textsuperscript{316} He must also file an application for custody with the court, which will rule on the application according to the best interests of the child.\textsuperscript{317}

If there is no dispute (the child is surrendered) but the surrogate is married, her husband is presumed to be the child’s father.\textsuperscript{318} The appropriate procedure is for the surrogate’s

\textsuperscript{315} See River, supra, note 275 at 850-52. See also the recent decision in Re Ontario Birth Registration Number 88-05-065848 (12 February 1990), Windsor A072/89 (Ont. Prov. Ct.), where the judge approved the adoption of a child whose mother had been artificially inseminated with her father-in-law’s sperm under a surrogacy agreement.

\textsuperscript{316} Dickson, supra, note 218 at 68:

Some donors . . . intend specifically to rear the children born to women who have acted as surrogate mothers. . . . All these expectations, however, are subject to displacement, sometimes quite arbitrarily, by legislative provisions drafted with no regard for the different forms of artificial insemination and reproduction. The Nova Scotia Family Maintenance Act, for instance, defines a "possible father" as one who has "had sexual intercourse with . . . the mother of a child," thereby excluding a donor for sexual reproduction.

See also art. 589 C.C.Q.

\textsuperscript{317} Canadian Family Law Guide, supra, note 274 at 2461: "The conflict between the common law principle of the prima facie right of a mother to the custody of her illegitimate child and the equitable principle that the welfare of the child is the paramount consideration is reflected in the case law relating to custody disputes between the parents of illegitimate children." See also at 2410, which quotes the decision in D. (W.) v. P. (G.), [1984] 5 W.W.R. 289 (Alta C.A.):

The traditional rule is that the natural father of a child born out of wedlock is a deemed stranger to the child. As such, he cannot obtain custody of the child from the mother without first demonstrating that she has either abandoned or neglected the child, or without offering other serious or commanding reasons. That is a court-made rule, however, and as such can be changed by the court.

\textsuperscript{318} Dickson, supra, note 218 at 69-70: "[A] sperm donor who seeks to establish his paternity may face legal obstacles, particularly if his object is to assert custody rights to a child born of a surrogacy motherhood agreement made with a married woman." Knoppers and Sioss, supra, note 269 at 716: "Presumptions of paternity. . . . with respect to children born of artificial insemination would work against any biological father where the gestational mother was married or cohabiting with a man."
husband to disavow his paternity or for the surrogate herself to challenge her husband’s paternity. The biological father may then claim paternity.

If a married surrogate refuses to surrender the child at birth, the social father will have to challenge the presumed paternity of the husband before it is confirmed by possession of status, in order to establish his own paternity through evidence that he is the biological father.

Even if the surrogate is not genetically linked to the child and there is a dispute, she will probably be deemed the legal mother. However, we may ask whether it would be possible for the genetic mother to challenge the gestational mother’s maternity and file a claim of maternity on the grounds of her genetic link to the child. Since the rule that establishes maternity by the fact of childbirth is not a substantive rule but a rule of

319. In Quebec, see art. 581 C.C.Q., subject to application of the defences at bar in art. 586 C.C.Q. to medically assisted procreation technologies other than artificial insemination. Note that art. 589 of Bill 125, supra, note 196, eliminates this ambiguity. For the Yukon and Newfoundland, see supra, note 277. See Knoppers, supra, note 284 at 220. Since, in the common law provinces, the presumption of paternity is rebuttable, a challenge to paternity may be made provided the evidence is sufficiently clear and convincing. Accordingly, the Canadian Family Law Guide, supra, note 274 at 2401, para. 4305, states:

It is a strong presumption in law that children born in wedlock are in fact the legitimate offspring of the husband and wife. Where the husband had opportunity of access, a mere denial of paternity is not enough to rebut the presumption (Re Johnston and Johnston (1975), 10 O.R. (2d) 249 (Prov. Ct); Gutierrez v. Gutierrez (1976), 28 R.F.L. 30 (Man. Q.B.)): mere is an admission of paternity by another (Re Brown and Argue, [1925] 3 D.L.R. 873 (Ont. C.A.); Re Anderson, [1947] 3 D.L.R. 302 (N.B.C.A.)). Admission of paternity by another, however, when coupled with a temporary assumption of the child’s support and rehabilitation with the mother, was found to be enough to rebut the presumption in Gray v. Foster (1974), 19 R.F.L. 12 (Ont. Prov. Ct).

See also B. (B.J.) v. K. (J.), supra, note 274 at 151:

The presumption may be rebutted by evidence which satisfies the court, on the balance of probabilities, that the child’s mother and her husband did not engage in sexual intercourse by which the child could have been conceived. The H.L.A. tissue typing test results should be taken into account, together with all the other relevant evidence, in determining whether the respondent has met the onus placed upon him. The weight given to the test results depends on the credibility of the parties.

320. For Quebec, see art. 589 C.C.Q. For the common law provinces, see Dickens, supra, note 279 at 355:

In a surrogate motherhood transaction involving a married woman, her husband’s name might have to be recorded as father of the child, having the intended father to seek a separate judicial declaration of paternity before he could gain a right of custody. He might also have to adopt his child before a birth certificate could be issued naming him as his child’s father.

Dickens, supra, note 218 at 70: “In contrast to an ovum donor, a man entering an agreement and donating his sperm for the insemination will in law be entitled to recognition as father of the child.”

321. See Martine Nolin and Hélène Guay, “Le phénomène des femmes porteuses: le droit à l’écoute de la science et de la société” in Martine Nolin, Réflexions juridiques sur le phénomène des femmes porteuses d’enfants (Cowansville, Que.: Yvon Blais, 1983) at 54; see also Rabellin-Devichi, supra, note 314 at 488.
— it does not appear in any legislation — and by analogy with the establishment of paternity, some analysts claim that it would be possible to rebut the presumption of maternity on the basis of genetic filiation.

In any event, it is clear from this brief review of the various issues that the current rules are inadequate.

322. Monique Hardae, "Réflexions sur la maternité" in Mélanges offerts à Pierre Raynaud (Paris: Dalloz-Sirey, 1985) 27 at 50. [TRANSLATION] "While the result is surely that evidence of childbirth is enough to establish maternity, the fact of delivery is by no means the very essence of the link, and one could not seriously argue that the authors of the Code civil intended to settle, through the evidentiary scheme they established, a substantive problem of which they did not have the faintest idea."

323. Ibid. at 30-31:

[TRANSLATION]
It springs from the need imposed on the interpreter to adopt in respect of the father and the mother the same notion of the biological element whose role is today predominant in the components of blood parentage... The basis for paternity then lies in heredity, and it seems that it is heredity as well, that is, the furnishing of a root cell, which forms the primary biological foundation of maternal parentage.

324. Knoppers and Sloss, supra, note 269 at 716:

If the gestational mother refused to surrender custody the law would presume her right to the child as its gestational mother subject to later proof of paternity. If the child was actually the result of in vitro fertilization utilizing the sperm and ovum of the social parents, the societal parents would have recourse against her only in so far as they could prove their genetic link to the child.

However, this situation is subject to unchallengeable presumptions of paternity in cases where artificial insemination is used, if these provisions apply to gestational surrogate mothers; see supra, note 277. Faced with the division of the biological and gestational aspects of motherhood, a Michigan court recognized the right of the genetic parents to have their names on the birth certificate and to be deemed the legal parents. Smith v. Jones (14 March 1986), Michigan 8552011660 Wayne Co. Cir. Ct., in Sherrill Cohen and Nadine Taub, eds., Reproductive Laws for the 1990s (Clifton, N.J.: Humana, 1989) at 388. In a recent U.S. decision we find the following comment:

[A] surrogate mother has lost her bid to be named the third parent of a test-tube baby she bore for an infertile couple.

Judge Richard Parslow of the U.S. Superior Court [sic] ruled... that Anna Johnson does not have any parental rights to the baby boy born a month ago, and he granted permanent custody to Mark and Crispina Calvert, the couple who paid Ms Johnson $100,000 (U.S.) to carry their fertilized embryo... Despite her contribution, the judge said, 'a surrogate carrying a genetic child for a couple does not acquire parental rights.'


325. The Uniform Law Conference of Canada, in An Act to amend the Uniform Child Status Act, supra, note 199, proposes the following amendments:

11.3. A woman who gives birth to a child before or after the coming into force of this section is deemed to be the mother of the child whether or not the child is conceived using the woman's egg.

11.4(1) A woman whose egg is used in an assisted conception and who does not give birth to the child conceived using her egg is deemed not to be the mother of the child.
2. Custody

From parentage stems parental authority, and parental authority is the basis for custody. In medically assisted procreation, if parentage is established as vesting in the surrogate and her husband, they will have custody of the child unless they are deprived of their parental authority and/or unless the interests of the child prevent them from retaining custody.

On the other hand, if legal parentage is established as vesting in the surrogate and the social father, it may be difficult to determine custody of the child. The criterion then is the best interests of the child, as determined in light of the circumstances in each case.

326. D.-Castelli, supra, note 271 at 182.
327. To deprive a person of parental authority, it must first be demonstrated that the holder of such authority has been guilty, by action or inaction, of a serious and unjustified failure to perform the parental duty (C. (G.) v. V.-F. (T.), [1987] 2 S.C.R. 244 at 246). Further, such deprivation must be in the interest of the child (see art. 654 C.C.Q., and art. 400 of Bill 125, supra, note 196). See also Dickens, supra, note 218 at 76-77.
328. With respect to the common law, see Dickens, supra, note 218 at 52:

In most cases . . . the State's role is now seen to be to pursue the individual child's best interests, established by legal process, in the conflict between the two principles [the "natural rights" of the parents and the best interests of the child], it seems to be accepted, in Canada and elsewhere in the common law world, that the "best interests of the child" principle has prevailed.

See also M. Joyce Schlosser, "Third Party Child-Centred Disputes: Parental Rights v. Best Interest of the Child" (1984) 22 Alta L. Rev. 394 at 398 and 401. With respect to the civil law, see D.-Castelli, supra, note 271 at 185; the author refers to the Supreme Court decision in C. (G.) v. V.-F. (T.), supra, note 327, in which the interest of the child was affirmed as a primary consideration. In the ruling, custody was awarded to third parties, taking into account the interest of the children without deprivation of parental authority or loss of custody rights. Beets J. at 266-67: "In such a situation, the holder is deprived of the exercise of custody but not of the right itself.

329. D.-Castelli, supra, note 271 at 186. See art. 30 C.C.L.C. and arts 33 and 34 of Bill 125, supra, note 196. See also Divorce Act, 1985, S.C. 1986, c. 4. For the determination of the interests of the child, see Chad vs. Chad (1978) 2 D.L.R. (3d) 641 at 647 (Ont. C.A.): "I do not think it safe to proceed on the assumption that a child will receive greater love and a more understanding upbringing if it is returned to a mother who did not want it at the time of its birth, than it would if left in the hands of those who sought it out for their love and care." Professor Dickens comments:

Similarly, it would be perverse, and possibly harmful to the child's best interests, to place the child with strangers, when the latter had not been shown to have violated legally mandated minimum standards of child protection. . . . In Ontario . . . section 55(1) of the Family Law Reform Act, know section 56 of the Family Law Act, 1985, supra, note 196] provides that:

"In the determination of any matter respecting . . . custody of or access to a child, the court may disregard any provision of a domestic contract pertaining thereto where, in the opinion of the court, to do so is in the best interests of the child." This provision embodies the position at common law, and is applicable in principle to disputed custody of a child born in a surrogacy agreement.

In Clark v. Clark (1992), O.W.N. 671 at 671-72 (H.C.), Barlow J. stated this position thus: "The agreement as to custody is not binding on the Court if the Court in its discretion is of the opinion that it is not in the best interests of the child's physical, moral, emotional and spiritual welfare." See also the OLRC report, supra, note 3 at 96-97, and the U.S. decision in Baby M., supra, note 102.

330. See Dickens, supra, note 218 at 53: "The concept of ‘best interests’ has become interpreted to mean the least detrimental alternative."
The parent who is not granted custody nevertheless retains parental authority, but the ability to exercise that authority may be reduced to a simple right of supervision. 331 The parent without custody may also enjoy access rights, but such rights are neither automatic nor guaranteed.

There are two options open to lawmakers: either to resolve the custody problem by giving legal effect to the private agreement between the surrogate and the prospective parents, or to let the dispute be resolved according to the interests of the child.

In summary, the problematic issues here are the contractual and commercial aspects of surrogate motherhood, and the parentage and custody of the resulting child. We will return to these issues in our chapter of recommendations.

III. The Canadian Charter of Rights and Freedoms

To complete our study of the different legal questions raised by medically assisted procreation, we will examine the impact that the Canadian Charter of Rights and Freedoms is likely to have on any attempt by the government to regulate the use of and access to reproductive technologies. 332

We will first examine whether a right to procreate is entailed in the right to liberty and security of the person enshrined in section 7 of the Charter, and whether such a right would entail a right of access to in vitro fertilization, artificial insemination, or a right to enforce a surrogacy contract. We will then turn to section 15 of the Charter, and discuss the impact that equality rights are likely to have on possible attempts to limit access to reproductive technologies. We conclude with a brief discussion of section 1 of the Charter.

But first it is necessary to stress that, in the absence of legislative regulation or other government intervention, the issues raised by medically assisted procreation are not constitutional issues. Pursuant to section 32, the Charter applies "to the Parliament and government of Canada" and "to the legislature and government of each province": it does not apply directly to the activities of private individuals. 333 Accordingly, the activities of doctors, hospitals or other non-governmental individuals or entities are not subject to the Charter unless their practices or policies are dictated by the government. 334

331. See D.-Caselli, supra, note 271 at 190-91.
332. The question of the constitutional division of legislative powers relating to these issues will not be addressed here.
Thus, a decision by government to abstain from regulating reproductive technologies will not give rise to constitutional challenges.\textsuperscript{335} On the other hand, a decision to regulate the use of and access to reproductive technologies will give rise to a number of potential \textit{Charter} challenges. In particular, a legislative restriction on access to reproductive technologies may violate either section 7 or section 15 of the \textit{Charter}.

A. Section 7: The Right to Life, Liberty and Security of the Person

Section 7 reads as follows:

\textit{Everyone} has the right to life, liberty and security of the person \textit{and} the right not to be deprived thereof except in accordance with the principles of fundamental justice [emphasis added].

The "\textit{and}" in the middle of the text (as well as the semi-colon used in the French version) suggests that the section could be read disjunctively to provide two rights, both a right to "life, liberty and security of the person" and a right "not to be deprived thereof except in accordance with the principles of fundamental justice." However, the courts have rejected this interpretation, finding that section 7 provides one right, a right not to be deprived of life, liberty or security of the person except in accordance with the principles of fundamental justice.\textsuperscript{336} Thus, establishing a violation of section 7 involves a two-step process: first, an individual must establish that his or her right to life, liberty or security of the person has been violated;\textsuperscript{337} and second, that the violation was not in accordance with the principles of fundamental justice.

The term "\textit{everyone}" raises the question of whether the unborn are included and can thus claim the benefit of the right not to be deprived of life except in accordance with the principles of fundamental justice. If the unborn do have section 7 rights, or if they are "individuals" for the purposes of section 15, then the \textit{Charter} could have an impact, for example, on the handling of embryos frozen for purposes of IVP.

Prior to the enactment of the \textit{Charter}, Canadian law recognized the legal existence of a fetus only upon its subsequent live birth.\textsuperscript{338} The courts that have addressed the issue

\textsuperscript{335} Of course, actions by non-governmental individuals or entities restricting access to reproductive technologies could be the subject of complaints under provincial human rights codes if individuals are discriminated against on the grounds of sex, family status, marital status, or sexual orientation.


\textsuperscript{337} The case law has established that the three interests protected by s. 7 — life, liberty and security of the person — are independent interests each of which must be given independent meaning. See \textit{Singh}, supra, note 336 at 204-05; \textit{Reference re Section 34(2) of the Motor Vehicle Act (B.C.)}, [1985] 2 S.C.R. 486 at 500; \textit{Morgenstuder}, supra, note 336 at 52.

under the Charter have followed the traditional common law position and held that the unborn do not have Charter rights.\textsuperscript{339} This position is consistent with the law in the United States,\textsuperscript{340} in England,\textsuperscript{341} and under the European Convention for the Protection of Human Rights and Fundamental Freedoms.\textsuperscript{342} By contrast, the Irish Constitution explicitly protects the right to life of the unborn,\textsuperscript{343} and courts in former West Germany have found that a fetus falls within the constitutional guarantee of the right to life.\textsuperscript{344}

The Supreme Court recently declined to resolve this issue in \textit{Borowski v. Canada (Attorney General),}\textsuperscript{345} holding that the issue was moot and that the appellant no longer had standing to pursue the action.\textsuperscript{346} Notwithstanding this uncertainty, it seems unlikely that the traditional Anglo-Canadian position on the rights of the unborn will be reversed under the Charter.\textsuperscript{347} Accordingly, we will proceed on the basis that the legal treatment of a fetus or embryo is not subject to constitutional constraints flowing from the constitutional status of the unborn.

\textsuperscript{339} \textit{Borowski v. Attorney-General for Canada} (1987), 39 D.L.R. (4th) 731 (Sask. C.A.); \textit{Campbell v. Attorney-General of Ontario} (1987), 58 O.R. (2d) 209 (H.C.), aff'd (1987) 60 O.R. (2d) 617 (C.A.), leave to appeal to the Supreme Court refused, [1987] 1 S.C.R. vi. But see the recent decision of the Quebec Court of Appeal in \textit{Duigle, supra}, note 219, in which the Court held, three to two, that a fetus has a right to life under the Quebec Charter, \textit{supra}, note 255. However, this judgement was reversed by an unanimous decision of the Supreme Court of Canada.

\textsuperscript{340} See \textit{Roe v. Wade}, 410 U.S. 113 at 161 (1973): "[T]he law has been reluctant to endorse any theory that life, as we recognize it, begins before live birth or to accord legal rights to the unborn except in narrowly defined situations and except when the rights are contingent upon live birth." However, in a recent ruling, \textit{Webster v. Reproductive Health Services}, 492 U.S. 490 at 518 and 526 (1989), a majority of the Court indicated that the decision in \textit{Roe} may be overruled in the near future. Four judges subscribed to the view that \textit{Roe} was "unsound in principle and unworkable in practice," and Justice O'Connor suggested that the Court should "reexamine Roe . . . carefully" in a future case.

\textsuperscript{341} \textit{Paton v. Trustees of BPAS}, [1978] 2 All E.R. 987 at 989 (Q.B.): "The foetus cannot, in English law, in my view, have any right of its own at least until it is born and has a separate existence from its mother."

\textsuperscript{342} Also known as \textit{European Convention on Human Rights} (1955) 213 U.N.T.S. 221. See \textit{Paton v. United Kingdom} (1980), 3 E.H.R.R. 408: a fetus does not have a right to life under art. 2 of the Convention, at least not in the initial stages of pregnancy.

\textsuperscript{343} Section 40.3.3. added to the Irish Constitution after a referendum in 1983, provides that: "The state acknowledges the right to life of the unborn and, with due regard to the equal right to life of the mother, guarantees in its laws to respect, and as far as practicable, by its laws to defend and vindicate that right."

\textsuperscript{344} See the discussion in \textit{Borowski} (1987), \textit{supra}, note 339 at 747-48.

\textsuperscript{345} [1989] 1 S.C.R. 342.

\textsuperscript{346} The 1989 \textit{Borowski} case, \textit{supra}, note 345, began as a challenge to s. 251, the old therapeutic abortion provision of the Criminal Code. In January 1989, the Supreme Court released its decision in \textit{Morgentaler, supra}, note 336, in which s. 251 was struck down as a violation of a woman's rights under s. 7 of the Charter. Thus, when the \textit{Borowski} appeal was argued later in the year, the legal basis for the challenge no longer existed.

\textsuperscript{347} For a full discussion of the many legal and ethical difficulties that would follow from such a holding, see Catherine Tolton, "Medical Implications of Constitutional Status for the Unborn: 'Ambulatory Choices' or 'Priorities and Aspirations'" (1988) 47 U.T. Fac. L. Rev. 1.
1. The Right to Procreate

Is a right to procreate entailed in the right to life, liberty or security of the person? Surely, a person’s life is not threatened by a denial of access to reproductive technologies. At the most, it is the possibility of creating a new life that is being denied to the person. Thus, an individual’s “right to life” is not relevant here. However, it may be that the right to have access to the means to attempt procreation is an element of either the right to liberty or the right to security of the person.

The meaning of the “right to liberty” has yet to be clearly set out by the Supreme Court of Canada. At least we know that the phrase encompasses deprivations of physical liberty such as imprisonment. Beyond instances of physical restraint, it is, as the Court has noted, a phrase “capable of a broad range of meaning.” Yet thus far only Justice Wilson has explored in her judgments the potential breadth of the “liberty” protected by section 7; the other justices have not yet found occasion to do so. For example, in R. v. Jones, Justice La Forest was willing to assume that the right to liberty included a “right of parents to educate their children as they see fit,” but he did not find it necessary to decide the issue.

Justice Wilson has articulated a definition of liberty that would entail a right to procreate. In her view,

the right to liberty contained in s. 7 guarantees to every individual a degree of personal autonomy over important decisions intimately affecting their private lives.

In Jones, she held that the right to liberty protects the parents’ right to raise their children in accordance with their conscientious beliefs. In Morgentaler, she held that a woman’s decision whether or not to terminate a pregnancy fell within the class of decisions protected from state interference by the right to liberty. Such a decision has “profound psychological, economic and social consequences for the pregnant woman.” Could we not say the same about the decision to bear and raise a child conceived with the assistance of a reproductive technology?

348. See Reference re Section 94(2) of the Motor Vehicle Act (B.C.), supra, note 337.
349. See Singh, supra, note 336 at 206.
350. Supra, note 336 at 302.
351. See Morgentaler, supra, note 336 at 171. See also Jones, supra, note 336 at 318, where Justice Wilson offered the following rationale for the right to liberty:

I believe that the framers of the Constitution in guaranteeing “liberty” as a fundamental value in a free and democratic society had in mind the freedom of the individual to develop and realize his potential to the full, to plan his own life to suit his own character, to make his own choices for good or ill, to be non-conformist, idiosyncratic and even eccentric — to be, in to-day’s parlance, “his own person” and accountable as such.

352. Supra, note 336.
353. Supra, note 336.
354. Ibid. at 171.
In defining liberty, Justice Wilson relied heavily on a series of U.S. constitutional cases establishing an area of personal autonomy over reproductive decisions as an element of the constitutional guarantee of "liberty" in the due process clause of the Fourteenth Amendment. For example, in Singh\textsuperscript{355} and Jones\textsuperscript{356} she relied on a passage in Meyer v. Nebraska\textsuperscript{357} in which the United States Supreme Court stated that:

[Liberty] denotes not merely freedom from bodily restraint but also the right of the individual ... to marry, establish a home and bring up children ... \textsuperscript{358}

And in Morgentaler,\textsuperscript{359} Justice Wilson approved of the U.S. cases, discussed below, that established a right of access to contraception and abortion as an element of liberty protected by the Fourteenth Amendment. In sum, Justice Wilson's position is that liberty would be infringed by any state interference with an individual's access to the means to procreate. However, whether such a broad conception of liberty will garner the support of the majority of the Court remains to be seen.

The right to security of the person, like the right to liberty, is capable of a broad range of meaning.\textsuperscript{360} The core meaning of the concept, in the words of then Chief Justice Dickson in Morgentaler, is that "the human body ought to be protected from interference by others."\textsuperscript{361} In Singh, Justice Wilson stated that security of the person protects an individual from the threat of physical punishment or suffering as well as freedom from the actual punishment or suffering itself.\textsuperscript{362} And the courts have indicated that security of the person extends to the control of one's psychological well-being as well as one's physical integrity.\textsuperscript{363} As Justice Lamer argued in Mills v. The Queen:\textsuperscript{364}

[S]ecurity of the person is not restricted to physical integrity; rather, it encompasses protection against "overlong subjection to the vexations and vicissitudes of a pending criminal accusation." ... These include stigmatization of the accused, loss of privacy, stress and anxiety resulting from a multitude of factors, including possible disruption of family, social life and work, legal costs, uncertainty as to the outcome and sanction.\textsuperscript{365}

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\textsuperscript{355} Supra, note 336 at 205.
\textsuperscript{356} Supra, note 336 at 317-18.
\textsuperscript{357} 262 U.S. 390 (1923).
\textsuperscript{358} Ibid. at 399.
\textsuperscript{359} Supra, note 336 at 167-71.
\textsuperscript{360} Singh, supra, note 336 at 206.
\textsuperscript{361} Supra, note 336 at 53.
\textsuperscript{362} Supra, note 336 at 207.
\textsuperscript{364} [1986] 1 S.C.R. 865.
\textsuperscript{365} Ibid. at 919-20.
In *Morgentaler*, the five majority justices found that serious state-imposed psychological stress violated security of the person. Justice Beetz held that security of the person "include[s] a right of access to medical treatment for a condition representing a danger to life or health without fear of criminal sanction." According to then Chief Justice Dickson,

state interference with bodily integrity and serious state-imposed psychological stress, at least in the criminal law context, constitute a breach of security of the person. It is not necessary in this case to determine whether the right extends further, to protect either interests central to personal autonomy, such as a right to privacy, or interests unrelated to criminal justice.  

It is possible that the anxiety and stress caused to a person otherwise unable to procreate by the denial of access to reproductive technology could fall within the definition of security of the person offered by the justices in the *Morgentaler* case. However, as then Chief Justice Dickson noted, the psychological component of security of the person has yet to be applied by the Supreme Court outside the context of a criminal prosecution.

The Court has also yet to decide whether security of the person extends beyond the protection of physical or psychological integrity to a broader right of privacy or autonomy that might encompass the right to procreate and other rights related to family life. However, a number of lower courts have suggested that security of the person does entail a right of autonomy over personal and intimate decisions.

Given the uncertainty regarding the meaning of "liberty" and "security of the person" in these early stages of *Charter* interpretation, it will be useful to consider the U.S. constitutional jurisprudence as it relates to the right to procreate. As we noted above, Justice Wilson found it useful in interpreting section 7 of the *Charter*, and in the absence of guiding Canadian precedent, U.S. case law will continue to be an influential source for *Charter* interpretation.

The United States Supreme Court first recognized the importance of procreative autonomy in *Skinner v. Oklahoma*, holding that forced sterilization of habitual criminals violated the equal protection clause. Justice Douglas termed the right to have offspring "a sensitive and important area of human rights... a right which is basic to the perpetuation of a race." Later in the opinion he added that:

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366. *Supra*, note 335 at 81.
368. See *Re T and Catholic Children’s Aid Society* (1984), 46 O.R. (2d) 347 (Prov. Ct) (right to security of the person includes the right to individual privacy or family autonomy); *S. (S.F.) v. Director of Child and Family Services*, [1987] 5 W.W.R. 309 (Man. Q.B.) (same); *R.L. Cronin Inc. v. Croucher* (1983), 6 D.L.R. (4th) 478 at 502 (Sask. Q.B.): "[T]he phrase ‘security of the person’ includes a right to personal dignity and a right to an area of privacy or individual sovereignty into which the State must not make arbitrary or unjustified intrusions."
We are dealing here with legislation which involves one of the basic civil rights of man. Marriage and procreation are fundamental to the very existence and survival of the race. . . . [The person sterilized by the state] is forever deprived of a basic liberty.  

Most of the U.S. cases protecting procreative autonomy have relied on the right to privacy guaranteed by the Fourteenth Amendment. Although the U.S. Constitution does not explicitly recognize a right of privacy, the Court has found that one aspect of "liberty" protected by the due process clause of the Fourteenth Amendment is "a right of personal privacy, or a guarantee of certain areas or zones of privacy." This right to personal privacy includes "the interest in independence in making certain kinds of important decisions," such as the choice to make use of contraceptives to avoid procreation. As Justice Brennan put it:

If the right of privacy means anything, it is the right of the individual, married or single, to be free from unwarranted governmental intrusion into matters so fundamentally affecting a person as the decision whether to bear or beget a child.

The Court has also recognized the rights to marry and to raise and educate children as fundamental rights under the Fourteenth Amendment. Similarly, in 1973 the Supreme Court held that the right to privacy outlined in these cases was "broad enough to encompass a woman's decision whether or not to terminate her pregnancy." The Court has held, however, that the state may refuse to pay for abortions even if they are medically necessary to preserve the mother's life or health, and that the state may prohibit the performance of abortions by public employees or in public hospitals.

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371. Ibid. at 541.
374. Griswold v. Connecticut, 381 U.S. 479 (1965) (state cannot make the use of contraceptives by married persons a crime); Eisenstadt v. Baird, 405 U.S. 438 (1972) (invalidating a regulation which made contraceptives less available to the unmarried than to married couples); Carey v. Population Services International, 431 U.S. 678 (1977) (invalidating a state ban on the commercial distribution of non-medical contraceptives). See Carey, ibid. at 687: "Read in light of its progeny, the teaching of Griswold is that the Constitution protects individual decisions in matters of childbearing from unjustified intrusion by the State."
375. Eisenstadt, supra, note 374 at 453.
376. Loving v. Virginia, 388 U.S. 1 (1967) (race may not be a basis for restricting an individual's right to marry); Zablocki v. Redhail, 434 U.S. 374 (1978) (striking down a statute requiring a certain class of state residents to obtain court permission to marry); Meyer, supra, note 357 at 399 (recognizing a constitutional right to "marry, establish a home and bring up children"); Pierce v. Society of Sisters, 268 U.S. 510 (1925) (child rearing and education).
380. Webster, supra, note 340.
The right to privacy in U.S. constitutional law thus protects the individual from state interference with his or her procreative potential. In addition, an individual has the right to be free from state interference with his or her access to the means of contraception. However, the state has no obligation to provide the individual with resources adequate to ensure access to the means of contraception.

Courts in the United States have not yet resolved the issue of whether the right to privacy includes the right to have access to existing reproductive technology free from state interference. A number of authors have argued that the right to privacy entails the right to unrestricted access to the means to attempt to conceive, John Robertson being the most forceful advocate of this position. In his view, any state regulation of access to reproductive technologies must be justified by a compelling state purpose: "the state must carry the burden of showing actual harm from [the] use of these techniques." But, as Knoppers observes, the ultimate judicial recognition of a constitutional right to procreate by whatever means available as an expression of personal liberty or privacy is uncertain, particularly in light of the current Supreme Court’s reluctance to expand this branch of substantive due process.

If uncertainty on these issues reigns under the U.S. Constitution, this is all the more true in Canada, where Charter jurisprudence is still in its early stages. In only one case has a Canadian court faced an argument that a right to procreate is included in section 7. In *E. (Mrs.) v. Eve*, the Supreme Court considered whether a court had the power, under its *parents patriae* jurisdiction, to authorize the non-therapeutic sterilization of a mentally handicapped woman. The Court concluded as follows:

> The grave intrusion on a person’s rights and the certain physical damage that ensues from non-therapeutic sterilization without consent, when compared to the highly questionable advantages that can result from it, have persuaded me that it can never safely be determined that such a procedure is for the benefit of that person. Accordingly, the procedure should never be authorized for non-therapeutic purposes under the *parents patriae* jurisdiction.

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386. *Rid.* at 431.
Relying on U.S. precedent, counsel for the party seeking the sterilization argued before the Court that section 7 entailed a right to free procreative choice, including a right to choose to have or not to have children and to implement that choice by means of contraception.\textsuperscript{387} Justice La Forest, speaking for the Court, did not find it necessary to decide this point:

\text{[A]}ssuming for the moment that liberty as used in s. 7 protects rights of this kind (a matter I refrain from entering into), counsel’s contention seems to me to go beyond the kind of protection s. 7 was intended to afford. All s. 7 does is give a remedy to protect individuals against laws or other state action that deprive them of liberty. It has no application here.\textsuperscript{388}

Although the Court did not find it necessary to rely on the Charter in deciding the case, it emphasized the fundamental nature of the right to procreate. Justice La Forest pointed out the “growing legal recognition of the fundamental character of the right to procreate,”\textsuperscript{389} “the great privilege of giving birth”\textsuperscript{390} and “[t]he importance of maintaining the physical integrity of a human being . . . particularly as it affects the privilege of giving life.”\textsuperscript{391} He characterized the proposed sterilization as a “grave intrusion on a person’s rights”\textsuperscript{392} and an “irreversible and serious intrusion on the basic rights of the individual.”\textsuperscript{393}

In conclusion, Canadian courts have not yet addressed the question of whether the right to liberty and the right to security of the person guaranteed by section 7 entail a right to procreate. However, in light of the expansive definition of liberty advanced by Justice Wilson, the influence of U.S. jurisprudence, and the strong language of Justice La Forest in the \textit{Eve} case underlying the importance to an individual of the ability to procreate, it seems likely that either liberty or security of the person, or both, will be found in a future case to include the right to procreate.

2. The Deprivation of Liberty or Security of the Person and the Principles of Fundamental Justice

As noted above, establishing a violation of section 7 involves a two-step process. First, one must establish that there has been an interference with life, liberty or security of the person and, second, one must show that the interference is not in accordance with the

\textsuperscript{387} \textit{Ibid.} at 436.
\textsuperscript{388} \textit{Ibid.}
\textsuperscript{389} \textit{Ibid.} at 419-20.
\textsuperscript{390} \textit{Ibid.} at 428.
\textsuperscript{391} \textit{Ibid.} at 434.
\textsuperscript{392} \textit{Ibid.} at 431.
\textsuperscript{393} \textit{Ibid.} at 432. The English courts reached the same conclusion in a case involving a fact situation similar to that in \textit{Eve}. In \textit{Re D (a minor)}, [1976] 1 All E. R. 326 at 332, the Court stated:

The type of operation proposed is one which involves the deprivation of a basic human right, namely the right of a woman to reproduce, and therefore it would, if performed on a woman for non-therapeutic reasons and without her consent, be a violation of such right.
principles of fundamental justice. Assuming that the courts do hold that the right to liberty or security of the person entails a right to procreate, legislation limiting access to reproductive technologies will not necessarily violate section 7. That will only be the case if the legislative limits are imposed in a manner that violates the principles of fundamental justice.

The principles of fundamental justice have both a procedural and a substantive component. In their procedural aspect, the principles of fundamental justice are similar to common law notions of procedural fairness. As Justice Wilson put it in Singh, procedural fairness means that:

[T]he tribunal which adjudicates upon [a person's] rights must act fairly, in good faith, without bias and in a judicial temper, and must give to him the opportunity to adequately state his case.

This means that persons seeking to exercise their right to procreate through the use of a reproductive technology must be treated in a procedurally fair manner by any law limiting access to that technology. The operation of the administrative structure must not be unfair or arbitrary, and the criteria determining accessibility must not be vague. An applicant who is initially denied access must be granted an opportunity to defend his or her rights. The applicant must be made aware of the reasons for the denial of access prior to the hearing, and at the hearing the applicant must be given an adequate opportunity to state his or her case. It should be noted, however, that an oral hearing will not be necessary in cases in which credibility is not at issue. As Justice Wilson stated in Singh:

I am prepared to accept the submission that procedural fairness may mean different things in different contexts. ... Thus it is possible that an oral hearing before the decision-maker is not required in every case in which s. 7 of the Charter is called into play.

I should note, however, that even if hearings based on written submissions are consistent with the principles of fundamental justice for some purposes, they will not be satisfactory for all purposes. In particular, I am of the view that where a serious issue of credibility is involved, fundamental justice requires that credibility be determined on the basis of an oral hearing.

According to the Supreme Court, the substantive aspects of the principles of fundamental justice are to be found in the basic tenets of our legal system. They do not lie in the realm of general public policy but in the inherent domain of the judiciary as guardian of the justice system.

394. Reference re Section 94(2) of the Motor Vehicle Act (B.C.), supra, note 337 at 497-99.
396. See Morgentaler, supra, note 336.
397. Ibid.
399. Ibid. at 213-14.
400. Reference re Section 94(2) of the Motor Vehicle Act (B.C.), supra, note 337 at 503.
An example of such a principle is the notion that the morally innocent shall not be punished; any law that has the potential of depriving the morally innocent of their liberty will accordingly violate section 7.\textsuperscript{401}

The substantive aspects of the principles of fundamental justice enable the courts to go beyond the examination of the fairness of the administration of the law to an evaluation of the substance of the legislation to determine whether it complies with ‘the basic tenets of our legal system.’ This appears to require the courts to pass judgment on the wisdom of a legislative policy that interferes with life, liberty or security of the person, a task that courts prefer to leave to the legislature. Accordingly, this is a branch of section 7 review that the courts are likely to apply with some caution.\textsuperscript{402} Thus far the Court has not developed any substantive principles of fundamental justice outside the context of criminal procedure.

As long as any restrictions on access to reproductive technologies are not arbitrary,\textsuperscript{403} it is unlikely that such restrictions would conflict with substantive principles of fundamental justice. The Court has stated that the future growth of the principles of fundamental justice will be based on ‘historical roots’\textsuperscript{404} that ‘have been developed over time as presumptions of the common law’ or ‘have found expression in the international conventions on human rights’ or ‘have been recognized as essential elements of a system for the administration of justice which is founded upon a belief in ‘the dignity and worth of the human person’ and on ‘the rule of law’.’\textsuperscript{405} An unrestricted right of access to the means necessary to procreate does not have such privileged roots in our legal traditions.

B. The Application of Section 7 to Certain Aspects of Medically Assisted Procreation

In light of the above general discussion of the principles governing the interpretation of section 7, we will now turn to the more specific issues raised by surrogate motherhood and access to medically assisted procreation.

1. The Enforceability of Surrogacy Contracts

In this section we will consider whether legislation rendering surrogacy contracts enforceable or unenforceable, or regulating the circumstances in which such contracts would be enforceable in court, would interfere with rights protected by section 7 of the \textit{Charter}.

\textsuperscript{402} See the comments of Dickson C.J. in \textit{Morgentaler, supra, note 336 at 53.}
\textsuperscript{404} Reference re Section 94(2) of the Motor Vehicle Act (B.C.), \textit{supra, note 337 at 513.}
\textsuperscript{405} \textit{Ibid.} at 503.
The only case that discusses the relationship between a constitutional right to procreate and the enforceability of surrogacy contracts is the U.S. case of Baby M,\textsuperscript{406} which arose out of the competing claims of the biological mother and father in a dispute over the custody of a child conceived in performance of a surrogacy contract. The biological father, Mr. Stern, argued that the right to procreate included the right to enforce a surrogacy contract. In holding that he had a constitutional right to custody of the child, the lower court reasoned as follows:

[If one has a right to procreate coitally, then one has the right to reproduce non-coitally. If it is the reproduction that is protected, then the means of reproduction are also to be protected. The values and interests underlying the creation of family are the same by whatever means obtained. This court holds that the protected means extends to the use of surrogates. . . . It might even be argued that refusal to enforce these contracts and prohibition of money payments would constitute an unconstitutional interference with procreative liberty since it would prevent childless couples from obtaining the means with which to have families. . . . A woman and her husband have the right to procreate and rear a family. The means to do so can be withheld from them only on a showing of a compelling state interest.\textsuperscript{407}]

The Court therefore held that the rights of the parties under the surrogate contract were constitutionally protected. On appeal, the New Jersey Supreme Court rejected this reasoning, holding that:

The right to procreate very simply is the right to have natural children, whether through sexual intercourse or artificial insemination. It is no more than that. Mr. Stern has not been deprived of that right. . . . The custody, care, companionship, and nurturing that follow birth are not parts of the right to procreation. . . . To assert that Mr. Stern’s right of procreation gives him the right to the custody of Baby M would be to assert that Mrs. Whitehead’s right of procreation does not give her the right to the custody of Baby M; it would be to assert that the constitutional right of procreation includes within it a constitutionally protected contractual right to destroy someone else’s right of procreation. . . . There is nothing in our culture or society that even begins to suggest a fundamental right on the part of the father to the custody of the child as part of his right to procreate when opposed by the claim of the mother to the same child.\textsuperscript{408}

The Baby M decision is in accord with the earlier case of Doe v. Kelley,\textsuperscript{409} a challenge to a Michigan adoption statute that prohibited the exchange of money or other consideration in connection with an adoption or related proceedings. A couple sought to engage in a surrogacy arrangement and to rely on an adoption proceeding to secure their legal right to the child. The court found that the statute did not violate the couple’s right to procreate:

\textsuperscript{407} Ibid. at 1164-65. See the criticism of this reasoning in Laurence A. Tribe, American Constitutional Law, 2d ed. (Mineola, N.Y.: Foundation, 1988) at 1360-62.
\textsuperscript{408} Baby M, supra, note 302 at 1253-54. See the criticism in Robertson, supra, note 382 at 23-24.
\textsuperscript{409} Supra, note 310.
The statute in question does not directly prohibit John Doe and Mary Roe from having the child as planned. It acts instead to preclude plaintiffs from paying consideration in conjunction with their use of the state's adoption procedures. In effect, the plaintiffs' contractual agreement discloses a desire to use the adoption code to change the legal status of the child — i.e., its right to support, intestate succession, etc. We do not perceive this goal as within the realm of fundamental interests protected by the right to privacy from reasonable government regulation. 410

Robertson has disputed these courts' claim that the right to procreate does not "include[] within it a constitutionally protected contractual right to destroy someone else's right of procreation." 411 He argues that the parties to a surrogacy arrangement should have the freedom of contract to dispose of their constitutional rights as they see fit:

A strong argument based on the autonomy of couples and surrogates can be made that the preconception agreement of the parties, which made the very existence of the child possible, should prima facie be determinative, just as it would be with sperm or egg donors. ... It simply is unclear why [the] agreement, if knowingly and freely made, should not control in [the] circumstances. 412

In the Canadian context, the argument that the biological father has a right under section 7 of the Charter to enforce a surrogacy contract depends on a number of problematic propositions: first, that the right to procreate entails custody rights once the child is born; second, that denying the father custody of the child would violate the principles of fundamental justice; and, finally, that the biological mother is free to contractually waive her own constitutional right to procreate (to the extent that it too would entail a right to custody of the child) and that she is not free to revoke that waiver when the child is born.

On the other hand, a biological mother's claim to a constitutional right to retain custody of the child conceived in performance of a surrogacy agreement would also depend upon the assertion of a number of problematic propositions: first, that the right to procreate entails a right to custody once the child is born; second, that denying her custody of the child would violate the principles of fundamental justice; and, finally, that a pre-conception contractual agreement to turn over the child is an ineffective waiver of her constitutional rights.

As far as the right to custody is concerned, an argument can be made that the right to procreate is an empty one if the state permits a child to be taken from its parents at birth. 413 However, there are obvious difficulties in superimposing a constitutional framework on the competing claims made by parties to custody disputes. Our law gives primacy to the best interests of the child rather than to the alleged Charter rights of the parents.

410. Ibid. at 441.
411. Robertson, supra, note 382 at 23; see also Baby M, supra, note 302 at 1254.
412. Robertson, supra, note 382 at 23-24.
413. Ibid. at 23: "[R]ejecting [is] the result that makes conception itself so worthy of protection ... Mr. Stern's interest or having a surrogate is precisely to conceive a child whom he will then mire."
It is not surprising that courts have been reluctant to allow the Charter to add further procedural complexity to child-protection and custody disputes.\textsuperscript{414}

Even assuming the biological parents could each assert a constitutional right to custody, we would nonetheless be left with reciprocal, competing constitutional claims, as in the Baby M case.\textsuperscript{415} It has been argued that the way out of this deadlock is to allow the contractual determination of rights to prevail: the biological father's constitutional claim would prevail because the biological mother waived her constitutional rights when she entered into the surrogacy agreement.\textsuperscript{416}

The Supreme Court of Canada has in fact held that individuals are free to waive their constitutional rights.\textsuperscript{417} The Court has stated that an individual "cannot be compelled to take advantage of rights for his or her benefit even if such rights may have a public interest aspect."\textsuperscript{418} To be effective, however, any waiver of rights must be voluntary, and it "must be premised on a true appreciation of the consequences of giving up the right."\textsuperscript{419} On this test, the question is whether a biological mother, at the time of entering into a surrogacy contract, could ever have a "true appreciation" of the emotional difficulty she might experience in surrendering the child to the biological father. Another difficulty is that a biological mother seeking to retain custody of the child is clearly no longer freely waiving her alleged constitutional rights. It is not clear that a waiver of constitutional rights would be effective if it is subsequently revoked in these circumstances.

At present, surrogacy contracts are probably unenforceable in civil and at common law as being contrary to public policy,\textsuperscript{420} although the issue has yet to be tested in a Canadian court. Recent legislation in England has rendered surrogacy arrangements unenforceable and prohibited all commercial agreements,\textsuperscript{421} a development in line with the recommendations of most of the studies carried out in this area.\textsuperscript{422} As Knoppers and Sloss have pointed out:

These recommendations are based on the negative psychological and emotional implications for the gestational mother on having to give up a child she carried to term, the repugnance of contracting to surrender parental rights, and the potential of commercial exploitation of such arrangements (which one British report has equated with prostitution).\textsuperscript{423}

\textsuperscript{414} See Re T and Catholic Children's Aid Society, supra, note 368; Re McTavish and Director, Child Welfare Act (1986), 32 D.L.R. 4th 394 (Alta Q.B.).

\textsuperscript{415} Supra, note 302.

\textsuperscript{416} See Robertson, supra, note 382.

\textsuperscript{417} See Clarkson v. The Queen, [1986] 1 S.C.R. 383 at 396 (right to counsel cannot be forced upon an unwilling accused); R. v. Turpin, [1989] 1 S.C.R. 1296 (waiver of right to a jury trial).

\textsuperscript{418} Turpin, supra, note 417 at 1316.

\textsuperscript{419} Clarkson, supra, note 417 at 396.

\textsuperscript{420} See "Legality and Legitimacy," supra at 65.


\textsuperscript{422} For more details, see appendix A, infra at 173.

\textsuperscript{423} Supra, note 269 at 709.
If legislation were passed to make clear that surrogacy contracts were legally unenforceable, it is unlikely that the courts would find a violation of section 7 of the Charter. Even assuming that any such legislation would be found to constitute an interference with liberty because of the burden on the right to procreate of men with infertile partners, and assuming also that the courts would hold that the biological mother is free to contractually waive her constitutional rights, restrictions on conception arrangements involving a consideration would not violate the principles of fundamental justice.

Rendering surrogacy contracts unenforceable is consistent with two fundamental principles of existing family law. The first is that custody of children be determined in accordance with the best interests of the child rather than the contractually expressed wishes of the parents. The primacy placed on the best interests of the child over freedom of contract is evident in family law legislation that prohibits parties from dealing with custody or access to children in marriage contracts or cohabitation agreements, and that empowers a court to disregard any provision of a domestic contract pertaining to the support, education, moral training, custody of or access to a child if it is in the best interests of the child to do so. Second, it is illegal to receive payment in return for arranging the adoption of a child. Such legislation, far from violating the fundamental tenets of the Canadian legal system, is consistent with those tenets.

In summary, there are many legal obstacles in the way of the argument that custody disputes following the birth of a child to a surrogate can be resolved by recourse to section 7 of the Charter. It seems likely that any legislation passed in this area, whether it renders surrogacy contracts enforceable or unenforceable, will not offend the section 7 rights of either biological parent.

2. **In Vitro Fertilization**

The theory that the right to procreate entails a right of access to IVF technology was argued before a U.S. federal court, but the judgment left the issue unresolved. In Smith v. Hurtig, an infertile couple alleged that an Illinois statute rendered in vitro fertilization illegal and thus “infringe[d] on their privacy interests because the provision prevent[ed] them from effectuating their only hope for conceiving a child.” While the defendant attorney general conceded that the “plaintiffs’ situation presents the strongest case for a fundamental right to [in vitro fertilization],” the court found it unnecessary to address the constitutional issue, having held that the statute did not prohibit in vitro fertilization in the circumstances.

424. See, e.g., ss 52(1)(a) and 53(1)(b) of the Ontario Family Law Act, 1986, supra, note 190.
425. Ibid., s. 56(1).
426. See supra at 67H.
428. Ibid. at 160.
429. Ibid. at 161.
It is not possible to predict whether a Canadian court would interpret the right to procreate to include access to IVF services. It may be that the right to procreate will be limited to natural procreation only, or extended no further than to include artificial insemination. Nevertheless, the prudent course would appear to be to assume that access to IVF might be held to fall within the scope of section 7 of the Charter.

Even were a court to so conclude, it would not mean that access to IVF could not be limited or regulated. Canadian case law suggests that the guarantee of Charter rights and freedoms does not impose affirmative obligations on governments to initiate laws or programs, because prior government action is necessary to invoke the Charter.430 Thus, the government has no obligation to initiate the provision of IVF services. But were the government to pass legislation restricting access to existing IVF services, then the necessary government action would be present and the right to procreate of infertile women and their partners could be found by a court to have been violated. If so, any such legislative restriction would have to be in accordance with the principles of fundamental justice.

It is difficult to predict what procedures a court would require to comply with the principles of fundamental justice in such a case. At the very least, fundamental justice would require that access to IVF services be determined in accordance with fair and rational criteria that are communicated to applicants. Furthermore, fundamental justice would require that applicants for IVF services be treated in a procedurally fair manner, and given an opportunity to present their case fully in their application for access. It is unlikely that a court would insist that there be an oral hearing at this stage, as long as the criteria are clear and no issues of credibility are raised.431 It is even more difficult to predict whether in this context fundamental justice would require that there be a system of appeal from decisions denying access to IVF services.432

430. See, e.g., RWDSU v. Dolphin Delivery Ltd., supra, note 333. However, where government has initiated a program on a limited basis, a remedy can be obtained under the Charter ordering the extension of benefits or services in a manner consistent with s. 15 equality rights. See, e.g., Schachter v. Canada, infra, note 584. Some sections of the Charter, such as the language rights in ss 16-23, do expressly create entitlements to a particular government service and may thus be invoked in the absence of government action; however, those sections are not relevant to the present discussion of the right to procreate.

431. See Singh, supra, note 356.

432. Ibid. In Singh, where the threat to liberty arose from the possibility of persecution (or even death) on the return of refugee applicants to their home country, the Supreme Court stated that fundamental justice required that a right of appeal be granted. In our view, the degree of interference with the liberty interest of an applicant for IVF services is not as extensive as was the case in Singh. Since the procedural protection required by s. 7 will vary with the degree of interference with the liberty interest, it may well be that a court would find an appeal right unnecessary in this context.
3. Artificial Insemination (AI)

The only reported cases dealing with AI involve the issue of the child's parentage. In the only constitutional challenge to a possible restriction of AI to exclude single women, the Michigan chapter of the American Civil Liberties Union filed a complaint on behalf of a single woman who was refused AI, but the case was settled when the clinic accepted the woman's application.

AI is unlike IVF in that minimal or non-existent technological requirements, public expense or medical complications are involved. The procedure can be performed very simply in the privacy of one's home without the intervention of experts. And unlike surrogacy, the involvement of the genetic donor is not an ongoing one: the genetic donor's involvement in the process is often over once he has made his contribution of semen. For these reasons, attempts to regulate the performance of AI raise serious privacy concerns that are not present with the other techniques of non-coital procreation. Just as regulation "is neither desirable nor practicable in the case of natural reproduction," it may be argued that the state has no place in the bedroom of a woman attempting to artificially inseminate herself. Regulation of such private behaviour would constitute an interference with liberty and might, it may be argued, infringe the principles of fundamental justice. A basic tenet of our legal system is the principle of non-interference in private behaviour that causes no harm to others.

Regulation of sperm banks or of the medical performance of AI does not raise such privacy concerns. The purpose of such regulation would be to ensure that genetic and other health problems are avoided. Ensuring the safety of the AI procedure in this way would not raise constitutional objections.

4. The Right to Be Informed of One's Biological Origins

The issue to be addressed in this section is whether a child conceived by means of medically assisted procreation has a constitutional right to information regarding his or her genetic origins.

The psychological need to know one's biological roots has gained increasing recognition in the context of adoption, and a growing number of adoptees have been pressuring courts and legislatures to relax the closed-record policies traditionally followed by adoption


435. OLRC, supra, note 2 at 154.
Recent studies have recognized that the disclosure of information to an adult adoptee is desirable as long as such disclosure does not conflict with the birth parents’ right to withhold identifying information if they so desire. It was recommended that the right of a birth parent or the donor of genetic material to remain anonymous should prevail over the child’s right to be informed.

In the United States, constitutional challenges to closed-adoption-record laws on the grounds that adoptees have a fundamental right to know their biological origins have not succeeded. The courts have generally held that closed-record laws achieve a desirable balance between the birth parents’ privacy rights, the state’s interest in protecting the integrity of the adoption process, and the adoptees’ need for information.

Section 7 of the Charter requires that a distinction be drawn between information that discloses the identity of the biological parents and that which does not. The release of non-identifying genetic and medical information is often necessary to the physical well-being of the child conceived through medically assisted procreation. The majority in Morgentaler recognized that security of the person included a right of access to medical treatment for a condition that represents a danger to life or health. Refusal to disclose information necessary to the preservation of life and health would similarly interfere with the security of the person conceived by means of medically assisted procreation. Legislation that provided for the automatic release of non-identifying information would protect the interests of the person so conceived without compromising the privacy rights of donors.

On the other hand, the disclosure of identifying information raises a clear conflict between the interests of the child in knowing the identity of his or her biological parents and the interests of donors in remaining anonymous. Many donors participate in medically assisted procreation programs on condition that they remain anonymous. Provisions for


437. See supra, note 260.


439. Supra, note 336.
the disclosure of identifying information would therefore jeopardize the operation of programs and interfere with the privacy interests of donors. In the face of these equally compelling competing claims, section 7 would not be violated by legislation that prohibited the disclosure of identifying information without the consent of the donor.\textsuperscript{440}

C. Section 15 Equality Rights

Section 15 provides, in part, as follows:

15. (1) Every individual is equal before and under the law and has the right to the equal protection and equal benefit of the law without discrimination and, in particular, without discrimination based on race, national or ethnic origin, colour, religion, sex, age or mental or physical disability.

A framework for the analysis of equality claims was set out recently by the Supreme Court in Andrews v. Law Society of British Columbia.\textsuperscript{441} Justice McIntyre, writing for a majority of the Court on the interpretation of subsection 15(1), expressed the goal of section 15 as follows:

[The admittedly unattainable ideal should be that a law expressed to bind all should not because of irrelevant personal differences have a more burdensome or less beneficial impact on one than another.\textsuperscript{442}]

As is the case for section 7, to establish a violation of section 15 one must follow a two-step process: first, one must establish a violation of one of the four basic equality rights guaranteed by section 15,\textsuperscript{443} and second, one must establish that the impact of the law is discriminatory.\textsuperscript{444}

The Court has defined the "minimal content of the right to equality before the law" as follows:

The guarantee of equality before the law is designed to advance the value that all persons be subject to the equal demands and burdens of the law and not suffer any greater disability in the substance and application of the law than others.\textsuperscript{445}

\textsuperscript{440} Most recent reports on both adoption and medically assisted procreation have recommended that non-identifying information be made readily available and that identifying information be released only with the consent of the biological parent. See Kneppers and Sloss, supra, note 269 at 693-96; Kneppers, supra, note 269 at 832-33; Ontario, Ministry of Community and Social Services, supra, note 436.

\textsuperscript{441} [1989] 1 S.C.R. 143.

\textsuperscript{442} Ibid. at 165.

\textsuperscript{443} Turpin, supra, note 417. A litigant must show that he or she is not receiving equal treatment before or under the law or that the law has a differential impact on him or her in the protection or benefit accorded by the law.

\textsuperscript{444} Andrews, supra, note 441 at 182.

\textsuperscript{445} Turpin, supra, note 417 at 1329.
With respect to the concept of discrimination, in the Andrews decision Justice McIntyre referred to and adopted the following definition of discrimination from the Action Travail des femmes case:

Discrimination . . . means practices or attitudes that have, whether by design or impact, the effect of limiting an individual’s or a group’s right to the opportunities generally available because of attributed rather than actual characteristics . . .

It is not a question of whether this discrimination is motivated by an intentional desire to obstruct someone’s potential, or whether it is the accidental by-product of innocently motivated practices or systems. If the barrier is affecting certain groups in a disproportionately negative way, it is a signal that the practices that lead to this adverse impact may be discriminatory.\footnote{446}

Justice McIntyre then proposed the following definition of discrimination:

[D]iscrimination may be described as a distinction, whether intentional or not but based on grounds relating to personal characteristics of the individual or group, which has the effect of imposing burdens, obligations, or disadvantages on such individual or group not imposed upon others, or which withholds or limits access to opportunities, benefits, and advantages available to other members of society. Distinctions based on personal characteristics attributed to an individual solely on the basis of association with a group will rarely escape the charge of discrimination, while those based on an individual’s merits and capacities will rarely be so classed.\footnote{447}

Not every legislative distinction will constitute discrimination. The Supreme Court has made it clear that only laws that have an unequal impact on groups of persons having in common any of the characteristics that constitute an enumerated or analogous ground can constitute discrimination for the purpose of section 15. A law that has an unequal impact on persons who cannot be associated with a group contemplated in section 15 cannot be considered discriminatory. The Court emphasized that the test for the eligibility of a group for section 15 protection lay in a situation of social, political or legal disadvantage related to a ground analogous to those enumerated in section 15. In Andrews, the Court stressed that non-citizens lack political power, have suffered a history of discrimination, and therefore constitute a good example of a discrete and insular minority deserving of section 15 protection.

On the other hand, in Reference Re Workers’ Compensation Act, 1983 (Nfld),\footnote{448} the Court summarily dismissed a challenge to a workers’ compensation statute that denied access to the courts to victims of workplace accidents and their dependents. The court was unanimously of the view that the statute did not create any discrimination within the meaning of section 15, since


\footnote{447} Andrews, supra, note 441 at 174-75.


\footnote{449} Ibid. at 924.
Similarly, in Turpin\textsuperscript{50} the Court rejected a challenge by accused persons outside the province of Alberta to a provision of the Criminal Code that provided accused persons in Alberta, but not elsewhere, the right to choose to have a trial by judge alone. Speaking for the Court, Justice Wilson rejected the claim on the grounds that accused persons outside Alberta did not constitute an analogous group for the purposes of section 15:

[...] it would be stretching the imagination to characterize persons accused of one of the crimes listed in s. 427 of the Criminal Code in all the provinces except Alberta as members of a "discrete and insular minority." [...] Differentiating for mode of trial purposes between those accused of s. 427 offences in Alberta and those accused of the same offences elsewhere in Canada would not, in my view, advance the purposes of s. 15 in remedying or preventing discrimination against groups suffering social, political and legal disadvantage in our society. A search for indicia of discrimination such as stereotyping, historical disadvantage or vulnerability to political and social prejudice would be fruitless in this case because what we are comparing is the position of those accused of the offences listed in s. 427 in the rest of Canada to the position of those accused of the offences listed in s. 427 in Alberta. [...] Persons resident outside Alberta and charged with s. 427 offences outside Alberta do not constitute a disadvantaged group in Canadian society within the contemplation of s. 15.\textsuperscript{51}

It follows that any attempt to regulate access to reproductive technologies by the use of rules that would have a disparate impact on an enumerated or analogous group would violate section 15. Conversely, rules that have a disparate impact on groups not analogous to the enumerated groups would not be discriminatory for the purposes of section 15. For example, an argument by an infertile couple that a rule limiting access to reproductive technologies discriminates against the infertile as a class is not likely to succeed. The infertile do not constitute a discrete and insular minority that have suffered a history of disadvantage or a lack of political power.\textsuperscript{52}

On the other hand, rules that discriminate or have a disparate impact on groups with respect to an enumerated ground such as sex, or an analogous ground such as marital status, parental status,\textsuperscript{53} family status or sexual orientation, will be vulnerable to a section 15 challenge.

One of the clearest examples of a non-enumerated, but nevertheless protected, ground of discrimination is marital status. It is a ground of discrimination expressly prohibited

\textsuperscript{50} Supra, note 417.

\textsuperscript{51} Ibid, at 1333.

\textsuperscript{52} It may be that rules will impose an unequal burden on a subset of the group of infertile people defined by their sex, class, marital status or sexual orientation. In that case, the possibility of making out a case of discrimination on a prescribed ground arises. See Martha A. Field, Surrogate Motherhood, expanded ed. (Cambridge, Mass.: Harvard University Press, 1990) at 47-49 for a discussion of these issues.

\textsuperscript{53} For an example of discrimination on the basis of parental status, see Synes v. Canada, [1989] 3 F.C. 59 at 81-82 (T.D.). It should be noted that the trial decision was reversed on other grounds (19 June 1991) A-290-89 (C.A.).
by human rights legislation in every Canadian jurisdiction. In addition, family status is a prohibited ground of discrimination in federal legislation, and in Manitoba, Ontario and the Northwest Territories. Discrimination against an individual based on his or her status as a single, married, divorced or widowed person is prohibited. Case law interpreting these legislative human rights guarantees has increasingly affirmed the principle that rules cannot be based on distinctions drawn upon stereotypical characteristics associated with the family status of an individual without a consideration of his or her actual situation or merits as they relate to the benefit or service at issue.

It is generally recognized that marital and family status are grounds of discrimination that ought to be recognized under section 15 of the Charter. For example, the Parliamentary Committee on Equality Rights concluded as follows:

We believe that section 15 of the Charter should be read against the historical background of the treatment in law of married women and the recognition nationally and internationally that marital and, in many cases, family status deserve protection by the state. Accordingly, while section 15 does not specifically prohibit discrimination on the basis of marital or family status, we believe that the ground can be properly read into the open-ended language of the section. In other words, marital or family status is implicitly covered by section 15.


455. The Human Rights Code, supra, note 454, s. 9(3)(c).


458. Brossard (ville) v. Quebec (Commission des droits de la personne), [1988] 2 S.C.R. 279 at 295 (municipality's blanket anti-nepotism policy discriminated on the basis of civil status); Carlin v. Canadian Broadcasting Corporation, [1988] 1 F.C. 494 (C.A.); Saskatchewan Human Rights Commission v. Saskatchewan Department of Social Services) (1985), 52 D.L.R. (4th) 253 (Sask. C.A.) (lower welfare benefits to single persons constitutes discrimination on the basis of marital status under the provincial human rights code); Schoop v. Canadian Armed Forces, [1989] 3 F.C. 172 at 184 (C.A.), Hugessen J.: employer's policy of granting family accommodation only to married couples perpetuates a stereotype, namely, that a relationship between a man and a woman has a lesser social value if it does not have the status of marriage. . . . It is a commonplace that the existence of the marriage bond is no guarantee of the permanency and stability of a relationship, just as its absence is no sure indicator of a more passing fancy.


In *Re MacVicar and Superintendent of Family and Child Services*, the Court concluded that discrimination on the basis of marital status was prohibited by the *Charter*:

Marital status is a direct result of a personal decision whether or not to marry with the blessing of the State and thereby obtain certain rights and incur certain obligations. . . . Marital status, in itself, bears no relationship to ability to nurture a child and consider its best interest.461

Similarly, there is a growing recognition of the need to provide legal recognition of the rights of sexual minorities.462 The Parliamentary Committee on Equality Rights concluded that "'sexual orientation' should be read into the general open-ended language of section 15 of the *Charter* as a constitutionally prohibited ground of discrimination."463 Sexual orientation is a ground of discrimination not enumerated in section 15, but is analogous to the enumerated grounds in that gay men and lesbian women constitute discrete and insular minorities that have suffered a history of discrimination.464 Sexual orientation is now a prohibited ground of discrimination in Quebec, Ontario, Manitoba and the Yukon.465

In sum, there is little doubt that any legislative limitations placed on access to reproductive technologies will have to be tailored to avoid discrimination on the basis of family status, marital status and sexual orientation. On the other hand, legislation requiring applicants to be assessed on the basis of their merits and capacities as potential parents would not violate section 15. Such an approach would also be in line with current Canadian adoption law and the recommendations contained in several provincial reports concerning reproductive technologies. For example, Ontario adoption law has recently been changed.

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465. *Quebec Charter*, *supra*, note 233, s. 10; *Human Rights Code, 1981* (Ontario), *supra*, note 454, as am. S.O. 1986, c. 64, s. 18; *The Human Rights Code* (Man.), *supra*, note 454, s. 9(2)(b); *Human Rights Act* (Yukon), *supra*, note 454, s. 6.
to allow for equal consideration of single applicants, as is the case in all other Canadian jurisdictions. Similarly, three recent provincial reports on reproductive technology recognized that limiting access to married couples would violate human rights laws. The Ontario Law Reform Commission concluded that restricting access to reproductive technologies to couples

would appear to contravene human rights legislation applicable in this Province. Moreover, any a priori exclusions based simply on membership in a particular group (such as married persons) would automatically eliminate from consideration single persons or unmarried couples who, by any standard, would make suitable parents.

The Ontario Law Reform Commission accordingly recommended that eligibility to participate in a medically assisted procreation program "should be limited to stable single women and to stable men and stable women in stable marital or nonmarital unions." Similarly, the British Columbia Royal Commission proposed that the guiding standard should be an applicant's "ability to nurture":

[A]n attempt to judge the recipient in terms of her conformity to prevailing mores about marriage and lifestyle should be made in the context of their current state of flux and, more importantly, should concentrate on the conduct of the individual which can be shown to relate directly to her ability to nurture. As suggested above, the central concern in evaluating the prospective AID recipient should focus directly (and singularly) on her ability to be a successful parent. It is the potential child's interest which must be paramount in this situation.

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466. Child and Family Services Act, 1984, supra, note 309, s. 140. Previously the law would only allow for the consideration of unmarried applicants in "special circumstances." The change was explained in the background paper to the legislation as follows:

Aside from a recognition of the changing concepts of marriage and the family, requirements imposed by Ontario's Human Rights Code have contributed to the elimination of the "special circumstances" qualification for unmarried adoptive applicants. Hence, single applicants will be eligible for consideration as adoptive parents. It is, however, anticipated that in most cases adoption practice would continue to reflect a strong preference for a two-parent family based on considerations of the child's best interests.


467. Child Welfare Act (Alta), supra, note 309, s. 56, as am. S.A. 1988, c. 15, s. 35; Adoption Act (B.C.), supra, note 309, s. 3; The Child and Family Services Act (Man.), supra, note 309, s. 66(1), 71(1); Family Services Act (N.B.), supra, note 283, s. 66; The Adoption of Children Act, 1972 (N.B.), supra, note 309, s. 4(1); Children and Family Services Act (N.S.), supra, note 309, s. 72(1) and (2); Adoption Act (P.E.I.), supra, note 309, s. 3(1); C.C.Q., arts 598-599; The Adoption Act (Sask.), supra, note 309, s. 17(2)(b); Children's Act (Yukon), supra, note 197, s. 79(1).

468. OLRC, supra, note 2 at 158.

469. Ibid. at 275.


[It] would appear that if a physician or medical institution which generally offers artificial insemination services to the public refuses to inseminate an unmarried woman solely because of her marital status, The Saskatchewan Human Rights Code would be violated.
If a limit on access is considered necessary, such an approach, similar to the criteria applied in Canadian adoption law, would be needed to satisfy the requirements of section 15 of the Charter. In addition, in light of the definition of discrimination set forth by the Supreme Court, care will have to be taken to ensure that the criteria chosen to regulate access to reproductive technologies do not unjustly burden a group of prospective parents defined by reference to one of the grounds enumerated in section 15 (or a ground analogous to those enumerated in section 15).

For example, let us consider infertility as a potential criterion for access to a reproductive technology. In the case of IVF, legislation making demonstrated infertility a precondition for access would be constitutionally unimpeachable. IVF is a procedure that will mainly be sought by women who are infertile; fertile women can bear children without the assistance of IVF. However, making infertility a precondition for access to surrogate motherhood and AI would give rise to section 15 objections. Such a requirement would impose a disproportionate burden on fertile men and women who wish to exercise their right to procreate non-coitally. These people are defined by their lack of a heterosexual partner, which is to say, by their marital status or their sexual orientation. Legislation imposing infertility as a requirement for access to AI and surrogacy arrangements would effectively deny these men and women the right to procreate. This disproportionate burden suffered by individuals as a result of their marital status or sexual orientation would constitute discrimination for the purposes of section 15.

In addition, care will have to be taken to ensure that criteria neutral on their face in terms of their impact on groups protected by section 15 or analogous groups, such as “ability to parent,” are not in practice applied in such a way as to impose an unequal burden on unmarried, single, gay or lesbian applicants for access to reproductive technologies.

D. Section 1

A law that infringed a right or freedom guaranteed in the Charter would not be unconstitutional provided that it conformed to the criteria set out in section 1 of the Charter. Section 1 provides as follows:

The Canadian Charter of Rights and Freedoms guarantees the rights and freedoms set out in it subject only to such reasonable limits prescribed by law as can be demonstrably justified in a free and democratic society.

471 See Andrews, supra, note 441; Turpin, supra, note 417.
For a limitation on rights to be upheld, it must be prescribed by law. This requirement was explained by Justice Le Dain in the following terms:

The requirement that the limit be prescribed by law is chiefly concerned with the distinction between a limit imposed by law and one that is arbitrary. The limit will be prescribed by law within the meaning of s.1 if it is expressly provided for by statute or regulation, or results by necessary implication from the terms of a statute or regulation or from its operating requirements. The limit may also result from the application of a common law rule.472

This means that any restrictions on access that would constitute an infringement of the Charter must be set out in legislation. Limitations that resulted from the application of policy guidelines or other directives without legal force would not be considered as "prescribed by law" for the purposes of section 1.

With respect to the other requirements of section 1, the basic framework was established by the Supreme Court in R. v. Oakes.473 In Oakes, the Court held that the legislative objective must be sufficiently important to justify the infringement of rights, and that the means chosen must be reasonable and justified in terms of the objectives sought.

There are certain limitations on the kinds of arguments that courts will entertain under section 1 with respect to such objectives. First, one cannot rely upon a legislative objective that is ultra vires Parliament or the legislature or is otherwise itself a violation of the Charter.474 Second, the objective pleaded in justification must be that which actually underlay the legislation, not one that was invented after the fact for the purposes of argument before the courts.475

These limitations aside, courts have tended to accept legislative objectives as sufficiently important to justify limitations on rights: most cases turn on the question of whether the limitations on rights were reasonable under the circumstances. In this respect, it is difficult to predict how a court would apply section 1 to any law restricting access to medically assisted procreation. It is now a commonplace to observe that the Supreme Court is divided on the question of how to apply section 1 of the Charter, and it is impossible to conclude that section 1 is being applied in a consistent manner.476 In some cases courts have ruled

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474. R. v. Big M Drug Mart Ltd., [1985] 1 S.C.R. 295 at 362 (per Dickson J., as he then was).
475. Ibid. at 362.
that government must give compelling reasons for its laws, while in others courts have shown a considerable degree of deference to the legislature.\textsuperscript{477} Indeed, recent cases leave unresolved the central question concerning the standard of review to apply in equality rights cases: whether different standards are to apply depending on the nature of the case and, if so, what those standards are.\textsuperscript{478} This considerably complicates the task of predicting how a court might apply section 1 to the issues considered in this study.


CHAPTER THREE
The Role of the State

There is currently no legislation in Canada that deals specifically with medically assisted procreation, but the technologies are subject to other types of control. We must therefore determine the adequacy of these controls before we can decide if legislation in this area is appropriate. Generally speaking, we cannot recommend greater state intervention in medically assisted procreation without considering the role of the state and of the federal government in this field.

I. The Scope of Existing Controls

Among the control mechanisms now in place are individual control based on the parties' individual responsibility and freedom of choice; professional and socio-professional control focusing on the quality of medical practice and the ethical aspects of applying scientific discoveries; community control exercised by permanent ethics committees (provincial and/or national) and the courts; and legislative and regulatory control.

479. With the exception of provisions in some provinces respecting parenthood. See supra, chap. 2.

[Translation]
The medical profession has behind it a long tradition of self-regulation. In this respect it is a very old legal order outside the state itself. In the absence of legislation or precedents, the College of Physicians has established its own rules and standards and a code of ethics that serve as law for members of the association. Through ethics committees, disciplinary committees or other means, physicians in hospitals set standards to regulate their conduct and their relations with their patients (and other medical personnel). The medical profession is a typical case of agents or instruments being recognized as having the authority to develop, interpret and enforce rules that have all the characteristics of law but do not originate with the state. . . . Of course, none of this stands in the way of state regulation. What must be considered, then, is the exact source of state regulation: what non-governmental legal order has the credibility, power and influence needed to ensure acceptance of its standards? What compromise will state regulation (by a legislature or a court of law) strike between the standards of several concurrent non-governmental legal orders?
A. Individual Control: The Private Ordering Approach and Individual Responsibility

Absolute priority is given here to individual freedom and the right to privacy. The individuals concerned make their own arrangements and no value-judgment is made regarding their choice. The assessment of the risks and benefits that are part of medically assisted procreation and the decision whether or not to use a technology are left to the free will of the people involved. The role played by lawmakers may be active or passive in that a legislature may specifically entrench freedom of choice in a statute or regulation, or may give tacit approval to such freedom. Without intervening at the outset, it can nevertheless regulate some of the outcomes of private decisions. For example, while it does not become involved in a couple’s decision to marry and raise a family, it establishes the property and other rights and obligations of the spouses and the children, as well as the parentage of the children.

This approach has the advantage of focusing on individual responsibility and initiative. It allows every member of a pluralistic society to conduct himself or herself according to his or her own beliefs, without undue interference by the state and without the state imposing a specific set of morals on everyone. It ensures that there is no conflict with the freedom and privacy rights guaranteed by the Constitution and at issue in procreation. The decision to conceive is therefore a purely private one, a moral choice to be left to individuals.

The state does not intervene in natural reproduction. As the Ontario Law Reform Commission has pointed out, the “best interest of the child” is never put forward to justify an intrusion into the private lives of couples or individuals. This criterion was used in the past for eugenic and financial reasons in legislation providing for the forced sterilization of mentally retarded persons, criminals and members of ethnic minorities, but this sort of motivation is no longer considered acceptable; the human rights charters and court rulings guarantee this.

Furthermore, treatment for sterility (hormone therapy, surgery, and so on) is one of the medical services available to the public, and the marital status and psychological profile

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482. OLRC, supra, note 2 at 106ff.
484. OLRC, supra, note 2 at 106ff.; Robertson (1986), supra, note 381 at 1040; The American Fertility Society, The Ethics Committee, supra, note 187 at 58.
485. It may intervene to sanction illicit sexual relations. In this case, however, the purpose is to protect persons who are exposed to exploitation, not to intervene in a joint decision to procreate: OLRC, supra, note 2 at 106ff.
of those undergoing such treatment are not scrutinized.\textsuperscript{487} If this is the case, why would we wish to treat "artificial" reproduction differently than natural reproduction? Is artificial procreation, too, not a part of the "private ordering approach" insofar as the actual decision to conceive is concerned?\textsuperscript{488}

For those opposed to state intervention in medically assisted procreation, society is in the process of coming to terms with all these discoveries and will ultimately consider their use a matter of course, as has been the case with so many other discoveries in the past. However, most defenders of this approach recognize the need to attach certain legal consequences to individual choices and to impose controls over the way at least some of the technologies are applied. These controls will be examined later, but suffice it to say for the time being that, in general, they are designed to ensure the quality of the services provided and, by extension, to protect the "recipients." In any event, defenders of this position do not question the legitimacy of using the various technologies associated with medically assisted procreation.

Individual freedom can also be advocated at this stage simply because any intervention is considered premature, as discussion of the issues is not yet sufficiently advanced to permit standards to be issued. What is therefore needed is community-wide reflection.\textsuperscript{489}

A full private ordering approach does not, however, have widespread support. It fails to answer any of the fundamental questions raised by the application to humans of scientific discoveries in the area of procreation, and it does not provide conscientious researchers with the guidelines they are seeking. It also creates difficulties in terms of the administration of justice. For example, the legal status of children born of medically assisted procreation has yet to be resolved, and control of gametes and embryos is foundering in a legal void.

The role of individual control in medically assisted procreation can be increased with proper public awareness and education. The task is to present the issues associated with new technologies to the public as objectively as possible. First to be addressed are the issues that affect individuals: the advantages and disadvantages for those using the technologies, the success and failure rate of each technology in the context in which it is used; and the physical and psychological risks for all of the people concerned. Next are the issues that affect future generations and society as a whole: the longer-term issues for the embryo, the fetus, the unborn child and the future of the human race; the fate of surplus embryos; the possible benefits to researchers — in short, all the major issues.

\textsuperscript{487} OLRC, supra, note 2 at 110.

\textsuperscript{488} ibid. at 118. Thomas A. Eaton. "Comparative Responses to Surrogate Motherhood" (1986) 65 Neb. L. Rev. 686 at 707.

\textsuperscript{489} Rubellin-Devichi, supra, note 314 at 457-59.
Currently being debated in various forums. However, it is difficult to guarantee that public education will be objective because it will always be based on information from many different sources. It could therefore be influenced by the fragmentation of the sources of information and might even be distorted by the unbridled confidence of some researchers and the sensationalism that sometimes characterizes the media.

This approach may also be accompanied by a promotion of values designed to instil a sense of responsibility in all the individuals concerned, and remind them that respect for human dignity must always take precedence over immediate individual interests. The positions taken by moral and religious authorities reflect this trend.490

Similar views can be seen in the positions adopted by various ethics bodies, which we will discuss later. Again, these positions must be conveyed to the general public, as the National Advisory Ethics Committee for the Life and Health Sciences in France is endeavouring to do.491 The recommendations of the American Fertility Society also stress the dissemination of information and moral counsel and the use of persuasion to enable people to make informed choices and gradually to alter public attitudes.492

These and other similar measures are vital to a better understanding by the general public of the real issues at stake in medically assisted procreation. But the basic information must from the start be non-denominational and non-political. And it should not be associated with specific groups, such as scientists, health professionals and women, although these groups do have an important role to play. No matter how active they might be in public education, these groups do not always display the objectivity needed to provide neutral information.493

One might ask what type of audience is reached by ethics bodies and moral and religious authorities? For Jacqueline Rubellin-Devichi, the opinions they issue have a considerable influence on public opinion.494 This is probably the case in Europe, but does it also hold true in Canada?


493. For example, some view reproductive technologies as nothing more than exploitation of women’s bodies for scientific research and “production” and as a result fail to see their contribution to procreation. This type of attitude was evident during the international forum, Sortir la maternité du laboratoire (Québec: Conseil du statut de la femme, 1988). On the other hand, the attitudes reflected in studies such as the Conseil du statut de la femme, Nouvelles technologies de la reproduction: pratiques cliniques et expérimentales au Québec (Québec: The Council, 1986) and the study by the Alberta Advisory Council on Women’s Issues, Discussion Paper on New Reproductive Technologies: Medical, Legal and Ethical Implications (Edmonton: The Council, 1988), bear witness to the important contribution of these groups.

494. Supra, note 314 at 457.
The position taken by the Catholic Church, which opposes medically assisted procreation outright, raises some reservations.\textsuperscript{495} For the United Church, artificial insemination with donor may be a legitimate choice in terms of ethics and morality.\textsuperscript{496} Surrogate motherhood itself could be admissible in very special circumstances.\textsuperscript{497}

Is education alone enough, given the issues raised by medically assisted procreation? Apparently not: education will raise awareness but has no compelling influence. It therefore offers no protection against the dangers of abuse or against commercialization of the technologies. As Jean-Louis Baudouin and Catherine Labrousse-Riou point out, it is unrealistic to rely on self-discipline.\textsuperscript{498} Public education is an important, even essential, step because we are looking for the solution or solutions to what is first and foremost a problem for all of society and not just an individual problem. Public education can discourage the abuse of the technologies made possible by science, but it is unlikely to have any impact on unscrupulous people.

B. Professional Control

The quality of a medical treatment may be controlled as to both its indication and its performance. Control may go beyond this practical stage and question the moral value of applying scientific discoveries; this is the role of ethics. In this regard, standards may be issued by professional associations or recommended by multidisciplinary committees.

I. The Medical Profession

Procreation assistance technologies are a type of therapeutic treatment and are naturally a part of the medical profession. This is certainly true of the more complex technologies that require special resources and a special environment. Artificial insemination, on the other hand, can easily be performed without special training. However, it is not without risk.

Physicians are required to comply with the standards of good medical practice as defined by their professional associations within limits set by the courts. These standards are designed to protect the public from unqualified individuals\textsuperscript{499} — in other words, malpractice or unlawful practice — and to set a benchmark for professional practice that

\textsuperscript{495} Congregation for the Doctrine of the Faith, supra, note 490 at 5ff.; the Congregation bases its opposition on the separation of sexual relations and procreation, which in its view fails to respect the dignity of the person. It fears the "technologization" of procreation. The Ethics Committee of the American Fertility Society has published a special discussion paper on this subject in response to the Instruction from Rome: "Ethical Considerations of the New Reproductive Technologies" (1988) 49:2 (Supp. 1) Fertil. Steril.

\textsuperscript{496} United Church of Canada, Division of Mission in Canada, "A Brief to the Royal Commission on New Reproductive Technologies," 1991, at 9-10 [unpublished].

\textsuperscript{497} Ibid. at 12.

\textsuperscript{498} Supra, note 210 at 208.

\textsuperscript{499} Ibid.
may be considered by the courts in the event of a dispute. Assuming that professional associations adopt positions on the basic legitimacy of assisted procreation, as some have already done, and set ethical guidelines for the use of new technologies by their members, effective sanctions must be devised for those who violate these guidelines. Standards, where they do exist, have no legal force unless they are included in regulations, as is the case, for example, with the Quebec codes of professional ethics.

In France, the Conseil d'État recommends specific sanctions for professional misconduct and urges professional associations to endeavour to prevent the misuse of new technologies. It even suggests that criminal sanctions be imposed for violating basic rules and procedures and for ignoring the advice of an ethics committee. Even then there is the problem of enforcement, especially when such fundamental choices are involved. Further, there is nothing of a tradition of secrecy and moral independence surrounding professional associations, and this could raise doubts about their ability to ensure real control that is sufficiently credible to the public. And if control of the application of discoveries were left to professional associations, standards would perhaps vary from one association to another, from one province to another and from one country to another, and it would be impossible to set the standard rules that most of those involved would like to see adopted.

Finally, as the members of the Quebec task force noted, regulation of human reproduction technologies is a matter of social policy that must not be identified with the development of standards for the professional quality of the services. The task must therefore not be left to the medical profession or to the other professions that are directly involved.


501. On all or only some technologies. For example, The American Fertility Society, *supra*, note 187 at 62 ff., favours considerable reproductive freedom but is opposed to surrogate motherhood for convenience and has reservations about the actual process. The Swiss Academy of Medical Sciences in *Directives médicales-éthiques pour la procréation médicalement assistée* (Zürich: P. Kober, 1990) recognizes the validity of in vitro fertilization for medical reasons, provided it is performed in the case of married couples or unmarried couples living a conjugal life (directive 3.1 at 2) and rejects surrogate motherhood (directive 12.6 at 5). The Conseil de l'Ordre des médecins in France accepts procreation assistance within the couple, has reservations about the intervention of unknown donors, and rejects surrogate motherhood and embryo donation in *Comité consultatif national d'éthique pour les sciences de la vie et de la santé*, supra, note 491 at 33. In Canada, the ethics committees of the Canadian Fertility and Andrology Society and the Society of Obstetricians and Gynaecologists of Canada have published a document listing their recommendations on all medically assisted procreation technologies; see *Ethical Considerations of the New Reproductive Technologies* (Toronto: Ribostride Communications, 1990).


Without standards from their professional associations or regulations developed at another level, physicians are faced with a dilemma: in dealing with medically assisted procreation, should they simply meet the demand, or should they instead question the values involved, consider the common good and exercise their own judgment? Leaving health professionals alone to set the ground rules in so complex a field puts a very heavy burden on them.

2. Local Ethics Committees

There are presently two major types of hospital or institutional ethics committees — research ethics committees and clinical ethics committees — and each type has a role to play in medically assisted procreation. Rather than give a complete background, we will simply state briefly that research ethics committees have been mandatory in Canada since 1978 for any research funded by the Medical Research Council of Canada: before a project is submitted to the Council, it must be approved by a local committee in accordance with the Council’s general criteria. Multidisciplinary committees are desirable but not mandatory; in some institutions, the committee is made up entirely of scientists. The main objective of research ethics is also to protect subjects in terms of both the inviolability of the person and the right to privacy. The Medical Research Council has issued special guidelines for research on embryos. The guidelines are extremely important because in most centres the use of in vitro fertilization produces surplus embryos. This raises a number of questions about embryos that will not be given to another infertile couple, among them storage, experimentation, destruction and ex utero development.

Since research on the treatment of infertility is costly and is for the most part funded by official agencies, ethical controls are generally applied. However, these controls are not always entirely successful: the local approval process is not always as thorough as it should be, and there is much criticism of the lack of project follow-up once initial approval has been obtained. What is more, when the research is carried out by a private company the controls do not apply. In addition, the controls no longer come into play once the technology is past the experimental stage. In effect, most medically assisted procreation technologies can now be used in fertility clinics and would therefore fall under the jurisdiction of clinical ethics committees.


507. Medical Research Council of Canada, Guidelines on Research Involving Human Subjects (Ottawa: Supply and Services Canada, 1987) at 33-34. Most of the papers, reports and recommendations mentioned in this working paper consider this aspect as well. See also Biomedical Experimentation Involving Human Subjects, supra, note 7.

508. The standards of the Medical Research Council of Canada can nevertheless be applied by hospital ethics committees when they consider research projects funded by the private sector.
The number of clinical ethics committees around the world has grown astronomically over the past ten years, and much has been written about them.\textsuperscript{509} These multidisciplinary bodies are generally able to address the problems referred to them from a much broader perspective. Their role is essentially an advisory one. In 1986, the Canadian Hospital Association recommended the creation of ethics committees in all major hospitals.\textsuperscript{510}

Clinical ethics committees are usually consulted when the attending physician feels he or she has a difficult decision to make or when there is a disagreement with the family or the medical team. But unless the institution has internal regulations requiring consultation in specific cases, the attending physician is free to treat patients for infertility as he or she sees fit; in other words, the physician can decide according to his or her own professional judgment to use medically assisted procreation technologies as soon as they are past the experimental stage. It should therefore come as no surprise that the criteria for using the technologies vary considerably.

In France, the centres for the study and storage of human sperm, or CÈCOS, have endeavoured to draft a common ethics policy. [TRANSLATION] "It became clear that the problems encountered were more than merely technical and demanded consideration of the very nature of artificial procreation through gamete donation, the significance of and rationale for the procedure, the inherent risks and the limits to be imposed."\textsuperscript{511} However, this policy applies only to frozen sperm, not fresh sperm, which physicians remain free to use as they wish.\textsuperscript{512} These regulations can also be circumvented by consulting private gynecologists.\textsuperscript{513} Opinion is also divided on the issue of compensation. The Quebec departmental task force\textsuperscript{514} points out that each clinic has its own rules.

Local committees still have undeniable advantages: apart from playing a role in education, as discussed above, they have very flexible operating structures and, with the proper membership, can be used to take the pulse of society.\textsuperscript{515} The ethical rules that emerge can in the medium term provide inspiration for lawmakers. However, as has been noted,\textsuperscript{516} they are but a temporary substitute for legislative intervention because they are not of any real effect unless they are accepted by patients. In disputes, they cannot be used as a basis for a court ruling unless they have somehow been incorporated in a statute or in regulations.\textsuperscript{517}

\textsuperscript{509} Baudouin, Ouellette and Molinarì, supra, note 506 at 7.
\textsuperscript{510} Durand, supra, note 505 at 553; and Baudouin, Ouellette and Molinarì, supra, note 506 at 7.
\textsuperscript{512} Rabellin-Deviechi, supra, note 314 at 460; Alain Sériaux, "Droit naturel et procréation artificielle : quelle jurisprudence?" D. 1985, 53 at 54, no. 4.
\textsuperscript{513} Jacqueline Rabellin-Deviechi, "Le droit, les pères et la paternité" (1988) 34-3 Rev. Droit santé et soc. 425 at 435.
\textsuperscript{514} Ministère de la Santé et des Services sociaux, supra, note 504 at 107ff.
\textsuperscript{515} This is even more true of national committees. LeRoy Walters, "Ethics and New Reproductive Technologies: An International Review of Committee Statements" (1987) 17:3 (Supp.) Hastings Cent. Rep. 3.
\textsuperscript{516} Dominique Thuessen, "Éthique et droit en matière biomédicale" D. 1985, 21 at 24-25, nos 12-13.
\textsuperscript{517} Ibid. at 23, no. 7.
Codes of ethics typically impose nothing more than minimum standards, and in dealing with patients, physicians must take into account their interests and their wishes. Ethical rules alone are not sufficient to deal with the problems raised by medically assisted procreation technologies.\footnote{518}

C. Community Control

1. Permanent Provincial or National Ethics Committees

Reference was made above to the regulations currently in place governing research. The need to regulate experimentation became apparent at the end of World War II, when it was discovered that horrifying experiments had been conducted without regard for the human subjects of them, on the grounds that the pursuit of knowledge justified whatever means might be used. Research ethics committees are in a position to impose rules because of their role in determining funding. Before issuing regulations, they normally consult broadly with the parties directly involved and with the general public. Their activities do not seem to give rise to opposition.\footnote{520}

When it comes to dealing not with research, that is, the acquisition of new knowledge, but rather with the application of that knowledge, ethical issues take on a very strong social dimension. For this reason, a number of authors and reports propose the creation of a national ethics committee where one does not already exist.\footnote{520} The American Fertility Society has also emphasized the need for national consultation. Gorovitz writes that “we need an independent agency of reflection and recommendation that is respectful of the full range of sentiment on such matters, and at the same time has the freedom and courage to take the stands that seem most justified — in full awareness that any stand on these matters will draw opprobrium from some quarters.”\footnote{521}

\footnote{518} Knoppers and Sloss, supra, note 269 at 671 n. 7 and the reports mentioned therein.

\footnote{519} For an overview of the main committees, see Durand, supra, note 505 at 352; Robertson, supra, note 483 at 8. See also Biomedical Experimentation Involving Human Subjects, supra, note 7; and Baudouin, Ouittellet and Molinari, supra, note 506.


\footnote{521} Gorovitz, supra, note 520.
Without compelling authority, how can ethical choices be imposed on those with different views? More to the point, what can be done to prevent commercialization, the proponents of which do not see beyond the law of supply and demand? The protracted search for an unattainable consensus is allowing new technologies to develop uncontrolled. It reflects [TRANSLATION] "the disarray of a society which, lacking a clear direction, intends to rely on a committee of scholars. . . . Is it possible that the future will be one of ethical rather than political laws?" 522 And are we to see in this an abdication of public authority?

2. The Courts

In the absence of clear legislation, the courts are among the first public institutions we turn to for settling disputes. Reference to the courts brings fundamental issues to the attention of the public and sparks the debate that is needed for any form of action, including legislative action. 523 The publicity afforded such issues by the media enables the courts to play a role in increasing public awareness. "Public policy may be affected by litigation and media attention. This in turn may affect the legislative approach that is chosen." 524

Existing laws do not deal specifically with medically assisted procreation. 525 Some, however, include provisions that may serve to govern a number of new situations. [TRANSLATION] "The concept of the interest of the child, an elastic concept open to a wide range of interpretations, has until now been used as the basis for decisions in cases involving new reproductive technologies and especially surrogate motherhood." 526

However, the application of existing rules to deal with a situation not anticipated when such rules were developed, such as recourse to the principles of natural justice, is not without problems.

522. Thurowlin, supra, note 516 at 26, no. 15.
523. Take, e.g., the Quiltan case (70 N.J. 10 (1976)) referred to in Baudouin, Ouellette and Molinari, supra, note 506 at 7. See also the comments by the Supreme Court of New Jersey in Baby M, supra, note 502 at 1264.
525. See supra, note 479.
526. Michelle Rivet, "Le rôle du juge et des parlements en matière de procréation assistée" (1990) 1:1 International Journal of Bioethics 49 at 52:

[TRANSLATION]
In the whole area of parenthood, the shift that has come about from a "social" truth to be protected to a "biological" truth to be recognized has given greater place to some aspects of "fictitious" or conventional truth. Now the trend is toward a "psychological" truth that springs from the very notion of the interest of the child, as a person with legal existence. Our courts will no doubt still have to rule on claims of parenthood and custody disputes between surrogate mothers who want to keep their children and biological fathers who claim the offspring, and they may then use this concept of the interest of the child.
See also supra, chap. 2.
For some of the issues raised by medically assisted procreation, the legislation now in place is of little use to the courts. This may lead the courts to avoid ruling on the fundamental questions underlying the disputes brought before them. This is what occurred, for example, in the Parpalaiai case in France, where the judgment dealt only with the return of frozen sperm to the widow, while the real social and ethical issue was post mortem insemination. 527 Further, while the discretion granted to judges may prove beneficial to some parties in a dispute, it also creates tremendous disparity in court rulings. The result for those who are parties to court actions is uncertainty as to the outcome of the case.

The judicial route therefore has its limits and does not offer a comprehensive solution to the problems associated with medically assisted procreation.

D. Legislative and Regulatory Control

1. Legislative Control

To the extent that courts, ethics committees, professional associations and citizens’ sense of personal responsibility are inadequate to ensure the desired control over all medically assisted procreation technologies or specific aspects of those technologies, legislative intervention, with all the necessary reservations and qualifications it implies, may be needed. It may be also needed to fill the gaps in positive law regarding some of the consequences of medically assisted procreation, including the parenting of children and control over gametes and embryos, in order to uniformize the situation in a given province or state, within a country or even internationally. 528 This approach may also be useful in examining the values of our society at a given point in history. There is not a single report or national committee that does not recommend some form of legislative intervention. 529

Public education may make the general population more aware of the issues and make freedom of choice more informed. Professional control and ethical review will limit the risk of abuse in the application of the technologies involved. The courts, by definition, deal only with disputes brought before them, which represent only a small proportion of the number of cases of medically assisted procreation, even surrogacy. 530

527 See supra, note 202. See Jones, supra, note 147.

528 D.G. Dickman, “Social Values in a Brave New World: Toward a Public Policy Regarding Embryo Status and In Vitro Fertilization” (1985) 29 St. Louis U. L.J. 817; Robertson (1986), supra, note 381 at 952 n. 48; Cornfield, supra, note 320 at 269 and 271; Stéaux, supra, note 542 at 54, no. 4; Knoppers and Sloos, supra, note 269 at 65 ff.; Bauduin and Labrousse-Rieu, supra, note 280 at 252.

529 See, inter alia, Warmack Report, supra, note 421; Benda Report, supra, note 520; Walker Report, supra, note 520; OLRC, supra, note 2; Ministère de la Santé et des Services sociaux, supra, note 504; Bureau du Québec, supra, note 3; and Conseil d’État, supra, note 503.

530 See in particular the studies on surrogacy in Canada: Margrit Eichler and Phebe Poole, “The Incidence of Preconception Contracts for the Production of Children among Canadians,” study prepared for the LRC, September 1998 [unpublished]; Ministère de la Santé et des Services sociaux, supra, note 504; Conseil du statut de la femme, supra, note 493; Alberta Advisory Council on Women’s Issues, supra, note 493.

Some writers suggest that positive state law is not the way to control all these issues\footnote{532. Rubellin-Devichi, supra, note 314 at 457-59 and n. 7; Gorovitz, supra, note 520 at 271.}. A law that is enacted and subsequently proves to be unenforceable risks being ignored, thereby missing its target altogether\footnote{533. Rubellin-Devichi, supra, note 314 at 457-59; Gorovitz, supra, note 520 at 271; Council for Science and Society, *Human Procreation: Ethical Aspects of the New Techniques* (Oxford: Oxford University Press, 1984) at 84, no. 8.6; Kiedler, supra, note 9.}. There are also fears that, despite appearances, legislation would not necessarily give everyone adequate protection\footnote{534. Hutton Brown et al., “Legal Rights and Issues Surrounding Conception, Pregnancy, and Birth” (1986) 39 Vand. L. Rev. 597 at 665.}. Others point to the risk of passing legislation hastily under pressure from the courts, public opinion and the media\footnote{535. Payne, supra, note 524 at 178.} when there is still no consensus\footnote{536. OLRG, supra, note 2 at 124; Payne, supra, note 524 at 180; Rubellin-Devichi, supra, note 314 at 496; Ministère de la Santé et des Services sociaux, supra, note 504 at 140; Council for Science and Society, supra, note 533 at 84, no. 8.9. For a general discussion of the subject see Kiedler, supra, note 9.}. Of course, it is important not to make a decision too swiftly. We must take the time to weight the pros and cons of legislative intervention, consider public opinion, define the ethical bases on which to act\footnote{537. Payne, supra, note 524.} and take pains to avoid dogmatism.

2. Regulation of Procedures

Legislative intervention may be sweeping or compartmentalized or may simply consist in regulatory control of practices. The proliferation of public and private centres in which medically assisted procreation technologies are applied obviously makes the task of applying controls more difficult, whether the controls are aimed at application standards, the quality of professional services or the cost of health care. Many reports therefore suggest that licences be issued to institutions and/or professionals providing procreation assistance services\footnote{538. See infra, appendix A at 173.}. In England, according to the Warnock Committee, the licensing authority should have the power to set standards for professional practice and research on medically assisted procreation and advise the government on specific matters\footnote{539. Warnock Report, supra, note 421 at 75ff. The Benda Report, however, supra, note 520, holds the view that such standards should be set out in legislation.}. This same general
approach has been adopted by some states in Australia. In this context, professional associations are without question very much involved in setting standards in the public interest. For the Quebec task force, the responsible agency would be the Department of Health and Social Services, which would set up an evaluation team to visit institutions periodically. France’s Conseil d’État recommends a certification system in accordance with the law and subject to administrative sanctions (revocation of licence) or criminal sanctions (for directors) in the event of a violation, as well as civil liability.

This type of regulation, which is inevitable if social service agencies are to take charge of infertility treatment, also makes it possible to guarantee uniform standards for all centres and therefore greater fairness to the public and more transparent practices. It would not prevent local ethics committees from ruling on specific cases. This approach would also permit better management of all the data related to artificial procreation. The Council of Europe recommends the establishment of a registry of all duly certified and authorized centres. The Ontario Law Reform Commission notes that the remoteness of centres may have an impact on accessibility; the Commission adds that this phenomenon is not unique to infertility treatment but holds for all types of fairly specialized health care.

II. The Role of the State and the Federal Government in the Reform Process

A. The Role of the State

Any legislative or state intervention in medically assisted procreation should be aimed at promoting values that society holds fundamental, such as the right to privacy and procreative autonomy, respect for the physical and mental integrity of patients.

540. See infra, appendix A at 198ff. Compare OLRC, supra, note 2 at 129 and 275ff.
541. Ministère de la Santé et des Services sociaux, supra, note 504 at 152.
542. Conseil d’État, supra, note 503 at 63-64, ss 9-12. The use of medically assisted procreation as provided for in the law must be under the supervision of the social security ministry. Ibid. at 67, s. 53.
543. See Ministère de la Santé et des Services sociaux, supra, note 504 at 152; Comité consultatif national d’éthique pour les sciences de la vie et de la santé, supra, note 491 at 30.
545. For France, see Catherine Labrousse-Ricaud, “Servitudes, servitudes” in Bernard Edelman and Marie-Ange Hermette, eds., L’Homme, la nature et le droit (Paris: Christian Bourgois, 1988) 308 at 326, where she writes that “[T]ranslation] ‘if we wish to humanize life, then the role of public policy is, on the one hand, to recognize the human being as different from objects or from another living form and, on the other, to protect the integrity, the dignity and the nature of individuals as subjects.’”
546. See R. v. Jones, supra, note 336 at 318-19 and Sterilisation: Implications for Mentally Retarded and Mentally Ill Persons, supra, note 486 at 63 (procreative liberty); Crimes against the Fetus, supra, note 7 at 39.
547. R. v. Morgentaler, supra, note 336 at 60-63.
equality, special protection of children and those who are otherwise unable to protect themselves or who are vulnerable to harm or exploitation by reason of incapacity. Indeed, Canadian society tends to regard many of these values as fundamental rights that give legal content to the moral concept of human dignity.

Such values and rights help define the various roles the state may play in medically assisted procreation. On the one hand, the rights to privacy, confidentiality and autonomy do not admit of burdensome or unwelcome governmental intrusion into reproductive choices. This view inspired former Prime Minister Trudeau’s famous comment — “[t]he state has no place in the nation’s bedrooms” — when, over two decades ago, the promotion of contraceptives, among other things, was decriminalized. It also inspires impassioned arguments against state control of assisted procreation technologies and against state power to decide who may procreate.

On the other hand, a societal commitment to children’s interests and rights, and to protecting those who cannot protect themselves, would seem to call for an active and supportive government role in advancing reproductive health and family life. This view helps legitimate laws that grant children born using medically assisted procreation access to confidential records for medical or genetic information essential to their health.

Between the polar extremes of the state as oppressor and the state as protector-liberator lies a spectrum of supportive roles the state may assume in fulfilling its democratic mandate respecting medically assisted procreation: namely, as dispute arbiter, health service provider, public financier, lawmaker, researcher, protector of public health, safety and

548. See Canadian Human Rights Act, supra, note 454, s. 3, prohibiting discrimination, inter alia, on grounds of family or marital status, gender, disability; and s. 15 of the Canadian Charter of Rights and Freedoms, supra, note 10. For a consideration of federal government obligations under international law in preventing discrimination and promoting equality by virtue of its international treaty obligations, see International Covenant on Civil and Political Rights, 16 December 1966, Can. T.S. 1976 No. 47, 999 U.N.T.S. 171, s. 26.


550. See E. (Mrs.) v. Eve, supra, note 385.

551. See Morgentaler, supra, note 336 at 164-66.

552. John Robert Colombo, ed., New Canadian Quotations (Edmonton: Hurtig, 1987) at 311. See also S.C. 1968-69, c. 41, s. 13 (amending former s. 159(2)(c) — now s. 162(2)(c) — of the Criminal Code). The amending law, also incorporated into the federal Food and Drugs Act, supra, note 295, contains express provisions for the regulation of contraceptive devices.


human life, strong and benevolent defender and promotor of human rights, or mere administrator of birth records. The precise roles are determined partly by historical and still evolving state roles in reproductive health, and partly by fundamental and evolving values that define and structure legal relations between the individual, the family and the state.

Many of the values underlying these various state roles are given expression by local, regional and national governments. Thus, provincial and territorial governments may license and regulate fertility and sterility specialists and clinics, as they do midwives, obstetricians and hospitals. They might help provide and pay for artificial insemination and IVF services for their residents, as they do for surgical treatment of infertility. They might prohibit discrimination in the delivery of fertility services, as they do for adoption or hospital services.

B. The Role of the Federal Government

The federal government derives its national health role and public responsibilities from the same value base. The value of protecting life, expressed constitutionally in the duty to protect Canadian public health and safety under the criminal law power, underlies the federal responsibility for certifying the safety and efficacy of therapeutic agents that affect fertility and sterility. Thus, the federal government regulates prosthetic tubes used in reconstructive surgery of the fallopian tubes, as it does fertility drugs and condoms — the latter of which help prevent the spread of sexually transmitted diseases that cause sterility. Such jurisdiction over safety issues — complemented by federal jurisdiction over interprovincial and international commerce — extends to regulating the import and export of reproductive tissues, drugs, and medical devices.

555. See Jones, supra, note 147 at 540 n. 75.


558. Compare Wemore, supra, note 549 at 288, 293 (federal Food and Drugs Act protection of national public safety and health based on the criminal law power) and Quarantine Act, R.S.C. 1985, c. Q-1, based on s. 9(11) and 2(7) of the Constitution Act, 1867 (U.K.), 30 & 31 Vict., c. 3. For further discussion of the criminal law power, see François Chevereau and Herbert Mars, Droit Constitutionnel (Montreal: P.U.M., 1982) at 742-45; and Peter Hugg, Constitutional Law of Canada, 2nd ed. (Toronto: Carswell, 1985) at 399-402.

559. See Food and Drugs Act, supra, note 298; federal licensing of the fertility drug clomiphene citrate was granted some 25 years ago (see supra, note 33).

560. For a discussion of federal import-export and quarantine duties under the Food and Drugs Act, supra, note 298, see Procurement and Transfer of Human Tissues and Organs, supra, note 250. In this regard, Agriculture Canada has for years administered a national health protection regime to govern the thousands of animal embryos and millions of semen doses that cross the border annually. See the Animal Disease and Protection Regulations, C.R.C. 1978, c. 290, ss 32, 30, 39, 64 and 115, adopted under the Animal Disease and Protection Act, R.S.C. 1985, c. A-11, recently replaced by the Health of Animals Act, S.C. 1990, c. 21, ss 2, 14, 16 and 19. These initiatives would seem to flow from the federal criminal law, quarantine, trade and commerce, and agricultural powers.
Similarly, the federal government promotes the values of liberty, autonomy and fairness by prohibiting unfair and misleading advertising of medical products and services.\textsuperscript{561} This role has been demonstrated by recent U.S. government prosecutions for illegal, false advertising of IVF success rates.\textsuperscript{562} Consistent with principles of informed decision making, such laws may help ensure that the Canadian consumer is not harmed, defrauded, or deceived in the purchase of infertility or sterility products and services. Moreover, the funding of infertility research by Health and Welfare Canada, the provision of infertility treatment to members of the Canadian Forces,\textsuperscript{563} and the legal entitlement of employees to paid pregnancy and adoption leave\textsuperscript{564} — rights predicated on the view that society should no longer place women in the position of having to choose prejudicially between family life and employment opportunities — all illustrate complementary avenues through which the federal government may actively promote reproductive health. Indeed, its obligations with respect to (a) protecting public health, safety, and human life-forms through its criminal law power, (b) regulating international and interprovincial trade through its trade and commerce and quarantine powers, (c) protecting and promoting the national interest by responding to pressing "national concerns," (d) protecting human rights through its Charter and human rights obligations, and (e) funding medical research and health services through its spending power — endow the federal government in the modern Canadian state with significant and diverse constitutional and legal bases for protecting reproductive health.\textsuperscript{565}


\textsuperscript{563} Members of the Canadian Forces may receive, at the National Defence Medical Centre or affiliated hospitals, such fertility services as artificial insemination by husband/partner, fertility drugs, reversal of tubal ligation, reversal of vasectomy. Personal communication, Department of National Defence Canada, Office of the Surgeon General, 1990.


\textsuperscript{565} On these constitutional bases, the federal government has already structured a number of relevant national safety, health protection, health services, and medical financing schemes through the Food and Drugs Act (supra, note 298), Quarantine Act (supra, note 558), Canada Health Act (R.S.C. 1985, c. C-6), Department of National Health and Welfare Act (R.S.C. 1985, c. N-10), Competition Investigation Act (R.S.C. 1985, c. C-34) (now the Competition Act) and the Medical Research Council Act (R.S.C. 1985, c. M-4). See supra, notes 558-561 and infra, note 574. See also Andrée Lajoie, Patrick A. Mollner, and Jean-Marie Aurey, Traité de Droit de la santé et des services sociaux (Montreal: P.U.M., 1981) at 891-92.
Finally, the state may also play a critical — and perhaps the quintessentially democratic — role in fostering societal inquiry, debate, study, education and reflection on medically assisted procreation through public forums in which the whole range of viewpoints may be expressed. The recently created Royal Commission on New Reproductive Technologies has been given the mandate to assist government authorities in fulfilling this role through its public hearings in some 20 locales in Canada. Such public debate may influence private reproductive choices in that it makes possible a better understanding of the current implications and provides guidance for future applications of medically assisted procreation.

Indeed, these public processes and forums seem likely to serve as the foundation and catalyst for national consensus, standards and reform. The public pronouncements of prior Canadian royal commissions and the legal initiatives to which they gave rise, as well as more recent public pronouncements from similar bodies in Australia, France, and Denmark are evidence of this. Compelling national concerns for which the federal government bears particular responsibilities continue to grow in Canada over such issues as the necessity of guaranteeing minimal but rigorous screening of gamete donors to protect potential recipients' offspring and the public from transmissible diseases; the absence of minimal national requirements for the collection, processing, storage and interprovincial distribution of gametes; the moral and legal status of frozen embryos; the legal relation between anonymous gamete donors and resulting offspring;

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566. See Jones, infra, note 147 at 545.
569. Some aspects of medically assisted procreation issues that have attained national dimensions may require uniform, national legislative solutions, as demonstrated by an inability of some provinces to address issues in such a way as to avoid "grave consequences" to residents of other provinces. In considering whether a problem is of "national concern," and warrants redress under the federal government's "peace, order and good government" powers, courts and lawmakers must ask whether the "national concern" poses the country with a single, distinct or somehow indivisible issue or problem that requires one national law. One test is whether provincial co-operation may realistically solve the problem, given the consequences that may flow from the failure of one province to co-operate. If the analysis reveals the need for one national law, federal intervention must still be reconcilable with the powers and responsibilities of the provinces.
570. Such standards have been called for in Canada for 15 years. See British Columbia Royal Commission on Family and Children's Law, supra, note 470; and Report on Human Sperm, supra, note 148.
571. Assuming that frozen surplus embryos should never be treated "as mere objects," what specific interests, rights or duties define their legal status in different contexts like death or divorce of the donors, or similar situations? See Biomedical Experimentation Involving Human Subjects, supra, note 7 at 49; Divorce Act, 1985, supra, note 329, ss 2.16, In re Estates of Elsa and Mario Ross, supra, note 384 (no inheritance rights for frozen embryos). See generally "Control over Gametes and Embryos," supra at 39.
the potential exploitation of children and women in surrogate motherhood arrangements; and the misleading but avoidable variations in the reporting of IVF success rates. Whether it be because such issues may imperil health, safety and life, or contravene fundamental values and rights, or challenge competing visions of the modern family, they combine to make legal and public policy reforms inevitable in Canadian society.

If legal reforms are to remedy inadequacies in the law, they may begin to do so by minimizing current ambiguities. This may be achieved by better defining the rights, duties, and interests of those affected by medically assisted procreation and the protection they should be afforded. Of course, reasonable minds may differ over whether a particular allocation of rights and duties is fair. But a just allocation of rights, duties and interests should be the common goal in a democratic society. Reformers should also be guided by an active, creative consciousness of the role the law may play in responding to the human need for both freedom and dependency in the various relations between child, adult, family, community and state: "The goal for the future is to devise reforms that help people help themselves - reforms that acknowledge the public as well as private influences on and preconditions for human relationships." 573

To instil these principles into the legislative reforms that must be carried out in the area of medically assisted procreation in Canada will require a concerted, sustained imaginative effort by federal, provincial and territorial governments. The articulation of public policy and national medical and bioethical standards will, of course, also depend heavily on the private sector and various communities. How should the process go forward?

The Royal Commission on New Reproductive Technologies is a welcome beginning. Still, in many respects the melding of efforts and views may well require the kind of common vision that underlay the enactment of universal health insurance in Canada years ago. If the enactment of such laws fundamentally and positively altered the relationship between the individual, the hospital and the state in Canadian society, it was because there was a shared vision of individual well-being, social justice and a national "co-operative partnership." 574 Whether changes of that magnitude are in order remains to be seen. 575 Still, similarly co-operative federal-provincial initiatives have led to the Canadian Blood Committee's management of national blood policy, the recently created Canadian Coordinating Office of Health Technology Assessment, 576 and the role of national expert

572. See supra at 13ff.
573. See Minow, supra, note 564 at 23.
574. See the preamble to the Canada Health Act, supra, note 565.
575. Much would seem to depend on societal consensus, even in the important and sensitive domain of national public policy, respecting the necessity, form, and content of laws and policies that express basic principles of justice in the area of procreation.
576. In this sense, the provision of medically assisted procreation services across Canada might be modeled, in part, on the provision of blood transfusion services. Canadians entering hospitals who are in need of blood may currently rely on three levels of federal-provincial co-operation. First, the hospital or clinic that a patient enters is likely to be licensed and regulated under provincial health facilities legislation. Second, to ensure the safety of the nation's blood supply, such clinics, hospitals, and blood banks across Canada must also comply with minimal federal standards on the collection, screening, processing, storage, shipping, and labeling of blood products. Third, the overall management and development of national blood policies and principles are overseen by the national Canadian Blood Committee. See Procurement and Transfer of Human Tissues and Organs, supra, note 250.
committees in developing Canadian environmental regulations.\textsuperscript{577} Such co-operation has been recommended by the Ontario Law Reform Commission.\textsuperscript{578} Perhaps a similar vision and similar partnership today would best enable law and policy makers to fashion as able and as just a response to the challenges of medically assisted procreation.

\textsuperscript{577} See \textit{Canadian Environmental Protection Act}, S.C. 1988, c. 22, s. 6: "For the purpose of establishing a framework for national action and taking cooperative action in matters affecting the environment and for the purpose of avoiding conflict between, and duplication in, federal and provincial regulatory activity, the Minister shall, in cooperation with the governments of the provinces, establish a federal-provincial advisory committee to advise the Minister on (a) regulations proposed." See also ibid., s. 34; and Environment Canada, \textit{Canadian Environmental Protection Act: Report for the Period Ending March 1990} (Ottawa: Supply and Services Canada, 1990) at 11.

\textsuperscript{578} See OLRC, supra, note 2 at 275, 277, recs 2, 17.