

Statement on human cloning*

National academies of science from all parts of the world are united in supporting a worldwide ban on the reproductive cloning of human beings, and in calling for cloning to obtain embryonic stem cells for both research and therapeutic purposes to be excluded from this ban.

Reproductive cloning

Cloning is currently the subject of intense global debate. Some countries have already banned the reproductive cloning of humans. We urge all other countries to introduce and support appropriate regulations to ensure that reproductive cloning is subject to a universal ban.

Human reproductive cloning by somatic cell nuclear transfer (Note 1) (see Box 1) raises many issues – ethical, social, economic and scientific. It is through scientific research that the prospect of human reproductive cloning has come to be an issue of public policy. Scientists therefore have a special responsibility in the associated public debate.

Scientific research on reproductive cloning in other mammals shows that there is a markedly higher than normal incidence of fetal disorders and loss throughout pregnancy, and of malformation and death among newborns. There is no reason to suppose that the outcome would be different in humans. There would thus be a serious threat to the health of the cloned individual, not just at birth but potentially at all stages of life – without obvious compensating benefit to the individual bearing this threat. Moreover, death of a fetus late in pregnancy could pose a serious threat to the health of the woman carrying it. Even on a purely scientific basis, therefore, it would be quite irresponsible for anyone to attempt human reproductive cloning given our current level of scientific knowledge.

It is not beyond the bounds of possibility that scientific knowledge could advance to the point where reproductive cloning by somatic cell nuclear transfer might be accomplished without undue risk. Such a situation would not of itself warrant the lifting of a ban on the prac-

tice, which would still face strong ethical, social and economic objections.

We therefore call on all countries worldwide to ban the reproductive cloning of human beings.

Cloning for research and therapeutic purposes

Similar to reproductive cloning, cloning for research or therapeutic purposes involves generating a human blastocyst (Note 2) via somatic cell nuclear transfer. However, the crucial difference is that the cloned blastocyst is never implanted into the uterus. Instead, cells isolated from the blastocyst are used to make stem cell lines for further study and clinical applications.

Research studies using such nuclear transfer techniques could be important for improving our basic knowledge of, for example, how the cell nucleus can be re-programmed to switch on the set of genes that characterize a particular specialized cell, or for understanding the genetic basis for human diseases, or for enhancing our understanding of re-programming faulty human genes. A more long-term goal would be to learn how to re-programme somatic cells into stem cells (see Box 2) and thus provide a way of obtaining stem cells, genetically compatible with the patient, without any need

for the use of eggs and embryos. It is, of course, only justified to carry out this research using human eggs where animal studies fail to provide a suitable alternative.

Nuclear transfer techniques also offer the prospect of therapeutic applications for patients requiring cell, tissue or organ transplants, by producing embryonic stem cells that are genetically compatible with the recipient and thus circumventing the problem of rejection. However, aside from the scientific challenges, there are problems with the cost of customized treatments and obtaining a supply of unfertilized human eggs. At present, as cloning is an inefficient process, it is likely many eggs would be required to make a single embryonic stem cell line. It remains to be established if cloning for therapeutic purposes will be clinically viable. Research into additional strategies to overcome immune rejection is therefore strongly to be encouraged and such research may require the use of human embryonic stem cells derived from early human embryos.

Cloning for research and therapeutic purposes therefore has considerable potential from a scientific perspective, and should be excluded from the ban on human cloning. Both policies should be reviewed periodically in the light of scientific and social developments.

List of IAP Member Academies which endorsed the statement:

Box 1. What is cloning

Cloning of an organism commonly involves a technique called somatic cell nuclear transfer, where the nucleus of an egg cell (containing its genetic material) is removed and replaced with the nucleus of a somatic cell taken from the body of an adult. If the reconstructed egg cell is then stimulated successfully to divide, it may develop to the pre-implantation blastocyst stage. In reproductive cloning, the cloned blastocyst is then implanted in the uterus of a female and allowed to continue its development until birth. However, in cloning for research or therapeutic purposes, instead of being implanted in the uterus, the cloned blastocyst is converted into a tissue culture to make a stem cell line for research or clinical applications.

- African Academy of Sciences
- The Caribbean Academy of Sciences
- Latin American Academy of Sciences
- Third World Academy of Sciences
- The Academy of Sciences of Albania
- National Academy of Exact, Physical and Natural Sciences, Argentina
- Australian Academy of Science
- Bangladesh Academy of Sciences
- National Academy of Sciences of Belarus
- National Academy of Sciences of Bolivia
- Brazilian Academy of Sciences
- Bulgarian Academy of Sciences
- Cameroon Academy of Sciences
- Chinese Academy of Sciences
- Academia Sinica, China Taiwan
- Croatian Academy of Arts and Science
- Cuban Academy of Sciences

*Issued by the InterAcademy Panel on International issues, Third World Academy of Sciences, 34014, Trieste, Italy.

Box 2. What are stem cells?

Stem cells are cells that can replicate themselves and also generate specialized cells as they multiply. Stem cells could be used to generate replacement cells and tissues to treat many diseases and conditions including Parkinson's disease, leukaemia, stroke, diabetes, spinal cord injury and skin conditions, including burns. Damaged organs or tissues would be colonized with sufficient normal cells with an appropriate scaffold for their reconstruction.

Stem cells occur at all stages of development from embryo to adult but their versatility and abundance gradually decrease with age. While embryonic stem cells may be able to produce any of the 200 different types of specialized cells that make up the human body, adult stem cells appear to be capable of producing only one or a limited number of types of cell. Recently some have argued that adult stem cells have proved sufficiently versatile and therefore there is no need to derive stem cells from very early human embryos. We believe the scientific findings that have been reported so far do not support this conclusion. Therefore research on both adult and embryonic stem cells is vital for a proper evaluation of the prospects of stem cell therapy for the treatment of serious disease and injury.

- Academy of Sciences of the Czech Republic
- Royal Danish Academy of Science and Letters
- Academia de Ciencias de la República Dominicana
- Academy of Scientific Research and Technology, Egypt
- Estonian Academy of Sciences
- The Delegation of the Finish Academies of Science and Letters
- Académie des Sciences, France
- Georgian Academy of Sciences
- Academy of Athens, Greece
- Hungarian Academy of Sciences
- Indian National Science Academy
- Indonesian Academy of Sciences
- Israel Academy of Sciences and Humanities
- Science Council of Japan
- Royal Scientific Society of Jordan
- Kenya National Academy of Sciences
- National Academy of Sciences of the Kyrgyz Republic
- Latvian Academy of Sciences
- Lithuanian Academy of Sciences
- Macedonian Academy of Sciences and Arts
- Academia Mexicana de Ciencias
- Academy of Sciences of Moldova
- Mangolian Academy of Sciences
- Academy of the Kingdom of Morocco
- The Royal Netherlands Academy of Arts and Sciences
- Academy Council of the Royal Society of New Zealand
- Nigerian Academy of Sciences
- Norwegian Academy of Sciences and Letters
- Pakistan Academy of Sciences
- Palestine Academy for Science and Technology
- Academia Nacional de Ciencias del Peru
- National Academy of Science and Technology, Philippines
- Romanian Academy
- Russian Academy of Sciences
- Académie des Sciences et Techniques

- Singapore National Academy of Sciences
- Academy of Science of South Africa
- National Academy of Sciences of Sri Lanka
- Royal Swedish Academy of Sciences
- Academy of Sciences of the Republic of Tajikistan
- Thai Academy of Science and Technology
- Turkish Academy of Sciences
- The Uganda National Academy of Sciences
- The Royal Society, UK
- US National Academy of Sciences
- Academia de Ciencias Fisicas, Matemáticas y Naturales de Venezuela

The following IAP Member Academies have also endorsed the statement since its release on 22 September 2003:

- Academy of Sciences of the Islamic Republic of Iran
- Academy of Sciences and Arts of Bosnia and Herzegovina
- Union of German Academies of Sciences and Humanities
- Akademi Sains Malaysia

Notes

1. Somatic cells are all types of cell other than egg or sperm cells or their precursors.
2. Approximately 5–6 days after a human egg is fertilized, it is known as a blastocyst and consists of about 100 cells, the majority of which are already specialized to form the placenta. Most countries that allow *in vitro* fertilization (IVF) treatment allow the use of embryos up to day 14 after fertilization.