



Open

distinguish...

- ethical aspects of a technology *as such*
 - e.g. the ethics of stem cell research
- ethical aspects of national authorities *granting IP*
 - e.g. ethics of a patent office granting patents on life forms
- ethical aspects of an individual *seeking exclusive IP*
 - e.g. ethics of a public funded agency patenting research)

Bioethics and Biosafety

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BIOETHICS AND BIOSAFETY IN BIOTECHNOLOGY

display, spoken description, or any other way in which information is made available to the public.

- Involve an inventive step, as well as being new, the invention must not be obvious from the state of the art. Obviousness is from the viewpoint of a person skilled in the area of technology that the invention is in.
- Be industrially applicable. This condition requires that the invention can be made or used in any kind of industry.

A patented invention is recorded in a patent document. A patent document must have

- description of the invention, possibly with drawings, with enough details for a person skilled in the area of technology to perform the invention.
- claims to define the scope of the protection. The description is taken into account while interpreting the claims.

The original patent document of a patent application is published by a patent office. The application then adds to the state of the art for later applications and anyone can comment on the application. Often the patent document needs altering or amending to meet the conditions above before a patent can be granted. The final version of the granted patent document is then republished. If more information about the state of the art is discovered after grant, the patent document can be amended and republished again.

Patent rights are territorial; a UK patent does not give rights outside of the UK. Patent rights last for up to 20 years in the UK. Some patents, such as those for medicinal products, may be eligible for a further 5 years protection with a Supplementary Protection Certificate.

A patent can be of value to an inventor—as well as protecting his business, patents can be bought, sold, mortgaged, or licenced to others. They also benefit people other than the inventor since large amounts of information can be learnt from other peoples patents — they can stop you from reinventing things or you can monitor what your competitors are doing. Patents also spur you or others on to develop your idea further, and once the term of the patent expires it can be freely performed by anyone which benefits the public and the economy.

TRADEMARK

A trademark is any sign which can distinguish the goods and services of one trader from those of another. A sign includes words, logos, colours, slogans, three-dimensional shapes and sometimes sounds and gestures. A trademark is therefore a “badge” of trade origin. It is used as a marketing tool so that customers can recognize the product of a particular trader. To be registrable in the UK it must also be capable of being represented graphically, that is, in words and/or pictures.

DESIGN

A design refers to the appearance of the whole or a part of a product resulting from the features of, in particular, the lines, contours, colours, shape, texture or materials of the product or its ornamentation.

In the United Kingdom, designs are protected by three legal rights:

(a) Registered designs rights

- gives the owner a monopoly on their product design.
- brings the right to take legal action against others who might be infringing the design and to claim damages.

4. Statement

5. Description of Materials and Test Methods

(a) Description of methods and materials used

- ## **(b) Reference to OECD Test Guidelines or other test guidelines.**

(a) A sum

- (b) All information and data required in the study plan;
(c) A presentation of the results, including calculations and statistical methods;
(d) An evaluation and discussion of the results and, where appropriate, conclusions.

7. Storage

The location

Storage and Retention of Records and Material

Storage and retrieval

Storage and retrieval

1. Archives should be designed and equipped for the accommodation and the secure storage of:

- (a) the study plans;
 - (b) the raw data;
 - (c) the final reports;
 - (d) the reports of laboratory inspections and study audits performed according to the Quality Assurance Programme;
 - (e) samples and specimens.

2. Material retained in the archives should be indexed so as to facilitate orderly storage and rapid retrieval.

Retention
Movement of material in and out of the archives should be properly recorded.

1. The following should be retained for the period specified by the appropriate authorities:

(g) The study

(b) Records of all inspections and audits performed by the Quality Assurance Programme

- (b) Records of all inspections and audits performed by the quality assurance programme
 - (c) Summary of qualifications, training, experience and job description of personnel
 - (d) Records and reports of the maintenance and calibration of equipment
 - (e) The historical file of Standard Operating Procedures

2. Samples and specimens should be retained only as long as the quality of preparation

3. If a test facility or an archival contracting facility goes out of business and has no local

3. If a test facility or an archive contracting facility goes out of business and has no legal successor, the archive should be transferred to the archives of the sponsor(s) of the study(s).



SEMESTER 5 / MEI 2017

INDIVIDUAL ASSIGNMENT

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SUBJECT : IPR, Bioethics, Biosafety
SUBJECT CODE : SBB 3194
LECTURER'S NAME : Prof. Dr Arokiaraj ^/l Pappusamy & Mr. P. Kandiah
SUBMISSION DATE : 02 August 2017

agriculture, it promises to improve food security with crops with higher returns, greater nutritional value and resistance to pests and environmental hazards, while reducing dependence on fertilizers and herbicides. In 1994, the first meeting of the Conference of the Parties to the CBD (COP) established a group of experts on biotechnology. Both should adopt security measures to deal with possible hazards and risks for human health and the environment. Although Flavr Savr was not commercially viable, two crops marketed shortly after a worldwide debate: BT cotton, a genetically modified plant to produce an insecticide to combat a common plague; and Soya List Roundup, which had been modified to resist glyphosate, a broad-spectrum herbicide marketed as Roundup (Meyer, 2011)). Human beings have used biotechnology during millennia: farmers experienced selective breeding and cross-fertilization to improve the yield and resistance of their animal crops and races, while fermentation resulted in gastronomic miracles through cheese production and Alcoholic beverages. (Photo: Franz Dejon, IISD / ENB) The highly accused case became a symbol of the fight between countries with great interests in agricultural biotechnology and a skeptical public to GMOs. The ethical standards promote the essential values for collaborative work, such as trust, accountability, mutual respect and equity 11. (2003). In some situations, it may be difficult or little practical to work in biological safety cabinets; In this case, the personal protection team can constitute the main barrier between personnel and infectious materials. While the need to preserve the world's genetic resources was debated at the Stockholm Conference on the Human Environment in 1972, biotechnology has not yet been discussed at the Stockholm Conference on the Human Environment. 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In Zambia, the government carried out intensive research and then refused GM food aid ³ environmental and socio-economic reasons, Ä Ä to protect its agricultural exports to Europe, which have strict regulation³ regulation³ osu us odneyulcni ,aÄgoloncetoib al noc sadanoicaler senoitseuc ratart aÄrebed n³ÄicnevnoC al euq ne noreitsisni ollorrased ne sesÄap soL .9 51 .soviv somsinagro raretla ed dadicapac artseun ³Änoiculover)acit@Äneg aÄreinegni omoc adiconoc n@Äibmat(anredom aÄgoloncetoib al ,so±Äa atneucnic ecah ,ograbme niS .H ,reyeM fdp.ne-20-eruhcorb-sb/snoitacilbup/cod/tnidbc. www//:sptth .sogseir ed n³Äitseg y n³Äicaulave al radroba y secnava sol rasivrepus arap orof nu anoicroporp olocotorP le ,secep y sotcesni omoc ,satnlp sal ed sotnitsid somsinagro noc sadanoicaler s;Äm zev adac senoicacilpa y sodaborpa MG sovitluc s;Äm noc ,senoicagitsevni sal neugisorp euq adidem A .anamuh dulas al arap sogseir sol atneuc ne odneinet ,dadisrevidoib al erbos sosrevda sotcefe renet nadeup euq anredom aÄgoloncetoib al ed setnatluser)VMO(sodacifidom soviv somsinagro ed soruges n³Äicazilitu al y n³Äicalupinam al ,etropsnart le ne n³ÄicctorP ed odauceda levin nu a riubirthoc se olocotorP led ovitejbo lE .)BDC(acig³Äloib dadisrevid al erbos oinevnoC led ocrum le ne odaborpa ,0002 ed aÄgoloncetoib al ed dadirugeS erbos anegatraC ed olocotorP le se dadirugesoiB ed lanoicanretni nemig@Är led oelc@Än le ,dadilautca al nE .oicremoc le noc sadanoicaler senoitseuc ed eires anu erbos odreucu nu a ragell odup on orep ,aÄgoloncetoib al ed dadirugeS erbos olocotorP le aÄraborpa EPC al euq aÄnopus eS .etneibma oidem led n³Äicadarged al rineverp arap selbatner sadidem ed n³Äicpoda al razalpa arap n³Äazar omoc esracovni ;Ärdop on latot acifÄtneic erbmuditrec ed atlaf al ,selbisreverri o sevarg so±Äad ed sazanema natsixe odnauC .arto a eicepse anu ed NDA ed sarbeh odneirifsnart y odneyartxe saviv salul@Äc sal ed acit@Äneg arutcurtse al odalupinam nah socifÄtneic sol ,secnotne edseD .aduya al ratpeca arap sodanoiserc naÄev es euq norala±Äes sonreibog soirav lauc le etnarud ,etabed osnetni nu ³Älbatne es n³Äicaunitnoc A .R ,reppt ,J ,rarelredniK ,.A ,oicnecsA ,.D .ragul us ne and distribution of its benefits, as well as access to genetic resources, the raw material for biotechnological innovation. CBD Secretariat. Routledge. Although Challenges remain, the Cartagena Protocol on Biotechnology Security has contributed to the development of many national regulatory frameworks on biotechnology, particularly in developing countries. The World ³ Organization for Animal Health focuses on animal diseases, including through the development of health standards for international trade in animals and animal products. Needs and recommendations 7. However, the history of modern biotechnology shows that innovation efforts ³ targeted a limited number of crops suitable for the agricultural model and markets of developed countries, rather than developing countries. For example, the biologÄa sintÄ currently creates microorganisms that synthesize products for fuels, pharmaceuticals and chemicals. Mohammed Ph.D. Research Center ³ Science Faculty Duhok, Kurdistan, Iraq Common Biosafety and Biosecurity Issues 2. (2003b). The 1975 Asylum Conference on Recombinant DNA brought together professionals who agreed on the first safety guidelines for biotechnology, in an early ³ application of the precautionary principle³ 1 Majed H. The scientific and normative debate on agricultural applications of engineering reached its point in the 1990s and 2000s. The Cartagena Protocol on Biosafety: an analysis of the results. An agrÚfico agrÄcola begins a field research ³ a day on genetically modified soybeans. Cordonier Segger, et al. The complainants claimed that the EU had failed to fulfill its obligations under the WTO SPS Agreement because its measures were not based on scientific ³. The CBD, which radically changes the global governance of biodiversity to suit the needs and expectations of developing countries .otxet .otxet le ne sairasecen sacidÄruj senoiculos sal a norejudnec senoicaicogen sal ed apate amitl@Ä al etnarud levin otla ed n³Äicapicitrap al y ryaM nauJ ed aicnediserp arodaripsni aL .acisÄF .socim³Änoce y selaicos sotcepsa n@Äibmat onis ,dadisrevidoib al ed elbinetsos osu le y n³Äicavresnoc al noc sadanoicaler senoitseuc ol³Äs on acraba ,dadisrevidoib ne socir The international community responded with food aid delivered through the World Food Program, but then it was revealed that 75% of the MAÍA donated by the USA. It could be generically modified. cosbey, a. , & burgiel, s. discussions on specific references at the beginning of precaution were driven à €

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