

The European Patent Office

Protecting your biotechnological inventions in Europe





Our mission



As the patent office for Europe, we support innovation, competitiveness and economic growth across Europe through a commitment to high quality and efficient services delivered under the European Patent Convention.



Structure of the European Patent Organisation





Autonomy



- Second largest intergovernmental institution in Europe
- Not an EU institution
- Financially independent
- Self-financing, i.e. revenue from fees covers operating and capital expenditure



38 member states

Albania • Austria • Belgium • Bulgaria Croatia • Cyprus • Czech Republic • Denmark • Estonia • Finland • France • Germany • Greece • Hungary • Iceland • Ireland • Italy • Latvia • Liechtenstein • Lithuania • Luxembourg • Former Yugoslav Republic of Macedonia • Malta • Monaco • Netherlands • Norway Poland • Portugal • Romania • San Marino • Serbia • Slovakia • harket of overs Slovenia • Spain • Sweden • Switzerland Turkey • United Kingdom

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European patent applications and patents can also be extended at the applicant's request to the following states:

Bosnia-Herzegovina · Montenegro



Locations

Staff (2010)



Munich	3 714
The Hague	2 623
Berlin	278
Vienna	107
Brussels	4
Total	6 726



The European Patent Convention

- The European Patent Convention (EPC)
 - provides the legal framework for the granting of European patents via a centralised procedure
 - establishes the European Patent Organisation
- 1973 Diplomatic Conference in Munich ► signature of the EPC by 16 countries
- 1977 Entry into force of the EPC in 7 countries marked as follows





Our role in the European patent system

 We provide patent protection in up to 40 European countries based on a single application in one of the three official languages (German, English, French)

European patent applications can be filed:

- direct with the EPO
- via the national patent offices of the contracting states
- based on an international (PCT) application

We are also responsible for

- limitation and revocation proceedings by patentees
- opposition proceedings by third parties
- appeal proceedings before the Boards of Appeal
- We will also be in charge of granting and administering the future Unitary Patent of the EU





Advantages of a European Patent

convenient

one application one language (EN, FR or DE) common procedure and law (EPC) patent protection in up to 38 countries a market of 600 million customers, ca. twice as in the US

cost-effective

costs less than three separate national patents major costs (translation) delayed after grant!

strong patent

thorough search stringent examination sound legal protection = very high assumption of validity central opposition and appeal



Biotechnological inventions



What is patentable at the EPO?

Art. 52(1) EPC

(1) European patents shall be granted for any *inventions*, in all fields of technology, provided that they are *new*, involve an *inventive step* and are susceptible of *industrial application*.



What is patentable in biotechnology?

All aspects of biotech: white/grey biotech	enzymes, industrial processes	100
red biotech	medical	
green biotech	agriculture and environment	
blue biotech	products derived from marine life	Contraction of the second seco
black biotech	energy production	





What does it really mean?

Products:

polypeptides	(enzymes, antibodies, etc.),
nucleic acids	(genes even human genes, promoters, vectors, antisense molecules, siRNAs, ribozymes, SNPs, etc.),
chemicals	(polymers, antibiotics, etc.)

Living organisms:

(non-human) animals, plants, cells, bacteria, viruses

Methods:

transformation, purification, production, *in silico* or *in vitro* screening, etc...

Medical uses:

"Compound X for the treatment of disease Y"





What about human genes?

Article 52(2)(a) EPC and Rule 29(2) EPC: an element isolated from the human body or otherwise produced by means of a technical process, including the sequence or partial sequence of a gene, may constitute **a patentable invention**, even if the structure of that element is identical to that of a natural element

Guidelines, C-IV, 2.3.1: to find a substance freely occurring in nature is mere discovery and therefore not patentable; if a substance found in nature is first to be isolated from its surroundings and a process for obtaining it is developed, that process is patentable. Moreover, if this substance can be properly characterised by its structure and it is new in the absolute sense of having no previously recognised existence, then the substance *per se* may be patentable.



Patent law versus bioethics



1992

The conflict between patent law and bioethics hits the headlines with for the first time the grant by the EPO of the first ever patent on a mammal (EP 169672).

Implanted with a human cancer gene, the so-called oncomouse has an increased disposition for developing tumours.



Opposition to stem cells



1999

The granting of the university of Edinburgh patent, which relates among other things to human embryonic stem cells, leads to wide-spread political debate about the boundaries of patent protection.



Decision of the Enlarged Board of Appeals



2009

The Enlarged Board of Appeals rules in its decision G2/06 that patents applications relating to methods or uses necessarily involving the destruction of human embryos are not patentable under ethical and moral considerations governed by Art. 53(a) EPC.

2012

Decision of the European Court of Justice substantially in line with the decision G2/06.



Need more information?



www.epo.org info@epo.org Tel. + 49 (0)89 2399 - 4636

Or better yet, ask us on booth 2727, we are happy to help you further!



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Our role in the international (PCT) system

- We process international patent applications
 - we act as a receiving office for international applications (PCT)
 - We carry out more than 40% of all international search and preliminary examination procedures





PCT: Why should you chose the EPO as International Search Authority?



International applications - the PCT

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Source: Wipo



PCT International Application Filings from 1990 to 2011



Note: The figures given for PCT applications filed in 2011 are WIPO estimates.

Source: WIPO Statistics Database, March 2012



US is the world's major user of the PCT procedure



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Source: WIPO Statistics Database, March 2012



The EPO as PCT authority

EPO as a Receiving Office for international applications (PCT)

EPO as International Search Authority

also for US applications

in any field of technology, including biotechnology

without restriction in terms of numbers

EPO as International Preliminary Examination Authority





The EPO, the most popular International Search Authority



% of total filings

Source: WIPO aggregated data



Why?

high quality

timeliness

reduced cost



Quality

Same quality as for the European searches background and expertise of the examiners access to 600 millions records, more than 7,000 journals largest patent database worldwide constant investment in search tools machine translation tools sequence searches in-house software development



EPO best patent office in the world according to the users!



"excellent" or "very good", while a further 37% considered it to be "good". Direct

Recent posts

Why trademarks rock

Links



Timeliness

Under the current framework, the EPO does not receive many search reports from the other IP5 Offices on time to be reused by EPO examiners.





Timeliness in transmitting ISRs



Note: Timeliness is calculated as the time elapsed between the priority date and the date on which the ISA transmits the ISR to the International Bureau. Source: WIPO Statistics Database, March 2012



Reduced cost

The International Search Report issued by the EPO as ISA will be used as European search report

Upon entry into the regional phase (European phase), no additional fee for a supplementary European search report

Saving: 1.165 EUR



The Unitary Patent


Key facts about the unitary patent

Basic principles

- a European patent granted under the EPC
- unitary effect for the territories of the 25 EU member states currently participating, at the applicant's request
- co-existence with the existing European patent and national patents
- validated in one single administrative step by the EPO for all the participating states in the language in which it was granted
- language regime being finalised; transition measures foreseen

Objective

European Council Presidency and EU Commission intend to have the first unitary patent granted in 2014





Advantages

For inventors

- protection in one single step for the 25 states currently participating
- significant cost savings (translation, validation, administration)
- **simplified validation procedure** (instead of up to 25 different procedures)
- simplified and more cost-efficient renewal procedure
- increased legal certainty due to uniform litigation system

For Europe

- optimal protection in the participating states as a whole
- better framework conditions for innovative companies and organisations
- simplified European protection mechanism for companies from outside Europe
- improved competitiveness of the European patent system



Overview of European patent grant procedure (I)





Overview of European patent grant procedure (II)





The unitary patent as a European patent





Overview of European patent grant procedure (I)





Overview of European patent grant procedure (II)







Cooperative Patent Classification (CPC)





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Cooperative Patent Classification -CPC

• The USPTO and the EPO agree to co-operate on a joint classification system based on ECLA (October 2010).



USPTO and EPO Work Toward Joint Classification System

"In view of the significant benefit to stakeholders of developing a transparent and harmonized approach to a global classification system for patent documents; in order to make the search process more effective; and in the belief that cooperation between their two offices will facilitate progress in undertaking classification harmonization projects under the IPS Common Hybrid Classification initiative, the USPTO and the EPO have agreed together to work toward the formation of a partnership to explore the development of a joint classification system based on the European Classification system (ECLA) that will incorporate the best classification practices of the two offices. This system would be aligned with the World Intellectual Property Organization (WIPO) classification standards and the International Patent Classification aspects of such a partnership.

The IP5 partner offices will be continually apprised of progress at appropriate IP5 forums. Stakeholders will receive regular updates on the substance and progress of classification partnership discussions between the two offices."

David J. Kappos

Benoît Battistell

October 25, 2010

Classification



Cooperative Patent Classification -CPC

The USPTO and the EPO agree on a **Joint Patent classification system based on ECLA**

For the EPO:

- Improve file and document routing
- Saving resources on (re-)classification of US documents in the future
- Common base for future classification revisions
- Renumbering of ECLA

For the USPTO:

Moving to an IPC-based classification system Enhanced access to non-US documentation



GLOBAL PATENT INDEX

EPO's patent information service for experts



Worldwide coverage (DOCDB)

→ advanced prior art search

Weekly update

□ advanced patent watch

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TRY GPI ON THE EPO STAND!



Grant rate in biotech vs. EPO overall





Withdrawals after European search

Biotechnology vs EPO overall





Numer of filed applications

Biotech vs. EPO overall

