

Term mining and term searching with applications used in the European Parliament

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 Part one – Tools for terminology search used by EP translators and new online tools

Part two – An introduction to term mining









 The Information Technology Support Unit (ITS DGTRAD) is the unit that provides technical and logistical support to Parliament's translation units so that they may concentrate on the job they do best – translating

Organisation:

- 5 teams: ServiceDesk, Project Administrators, Research and Development, Communication, Administration
- Some of our products, projects and services
 - TFlow support, DGTRAD IT Development plan, Translation Portal, Euramis/SPA, e-Parliament, Gepro+, AT4TRAD, FullDoc, CAT4TRAD, e-Dictionaries, CAT-Tools, Machine Translation







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CLS: Updated Forecasts - Now Also Per Policy Area

Posted by : pekka.kuusisto (Planning) on Wed 26/05/2010 at 16:08

You can now access the latest forecasts for the Committee Liaison Service which include the number of expected PRs, AMs, RRs and ADs for the coming two months. These forecasts are regular, updated and they show bot the number of documents according to their expects. Translation deadlines an the number of documents expected to be submitted for

In addition to the global forecasts containing procedures from now you can also have a closer look at different policy areas:

Economic and Scientific Policy

Structural and Cohesion Policy 970×847 Citizens' Rights and Constitutional Affairs

Budgetary Affairs and CRIS Special Committee

External Policies

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 Committee Liaison Service - A Closer Look Posted by : paivi.vainiomaki (Planning) on Wed 26/05/2010 at 12:08

You have probably heard of a new service in the Planning unit - the Committe Liaison Service. Since we are a new service, we would like to share with you some basic information about our objectives and activities.

Read more

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Yearly Screen-Mouse-Keyboard Cleaning Operation/Nettoyage Annuel Des Écrans-Claviers-Souris: 27 May - 3rd June

Posted by : paolo.panbianco (ITS) on Wed 26/05/2010 at 9:59

DG Itec informs us that they will perform the Screens, Keyboards and Mice yearly cleaning operation from the 27th of May until the 3rd of June. Click on "More..." to view the calendar,



normally.

MY DICTIONARIES

- * Brockhaus Enzyklopädie DE
- Grand dictionnaire terminologique FR
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- * MOT Collins French Dictionary FR<->EN
- Τριανταφυλλίδης (Triandaphillidis) On-Line Greek/grec EL
- MOT Oxford German dictionary 1.0 DE<->EN
- Oxford Language Dictionaries Online EN->DE
- Oxford English Dictionary EN
- Oxford Language Dictionaries Online DE->EN

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· Elise

	Into on othe	r tools
e.	FullDoc	Info
b.	Euramis	Info
×	EUR-Lex	Info
	IATE	Info
	FUSE	Info
	Gepro	Info
+	Shout!	Info
	Googlex	Info
	Profil	Info





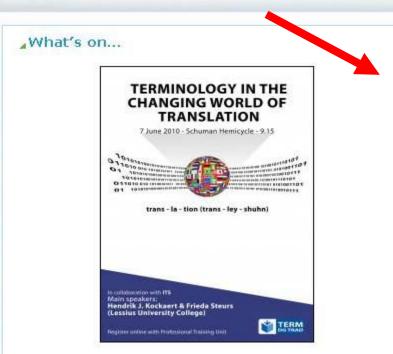
Info



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Home 🎇 28. May 2010 What's on... Terminology Coordination Terminology Material Proactive Terminology Language Specific Material Meetings IATE. Contacts TermCoord Projects Seminars and Conferences Did you know that ... Recommended reading Terminology Server Newsletter



Click the poster above to register, by entering your username and password in the dialogue-box that appears. Once you're logged-in, click on 'Courses for DG TRAD (Internal)', then click the 'Enrol' icon of 'Terminology in the Changing World of Translation', complete the details as requested, and click 'Submit'.

IATE	Info
Text Related Forum	Info
EP Est. Plan	Info
TOSCA	Info
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Elise	Info
Quest2	Info
CIRCA	Info

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31						
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DO Trad	Open Day Sth May
2010	Open Day - 8th May



Contact Us





Terminology Tools used in the EP Classification of tools





- Termbases, Glossaries, Dictionaries
 - IATE, E-dictionaries, Glossaries at the Term. Coord server, Glossaries created and hosted by the units
- Bilingual concordancers: search in context
 - FullDoc, Euramis concordance, TWB concordance, MyMemory, TAUS search, Linguee
- EU documentation and legal databases
 - Eur-Lex, Europarl, Council
- QUEST2 Meta-Search







Terminology Tools used in the EP Classification of tools

- Online Reference
 - Wikipedia, Google scholar
- Search Engines
 - Google, Yahoo, Microsoft ...
- Terminology management
 - Terminology macro
- Machine Translation







Termbases, Glossaries, Dictionaries

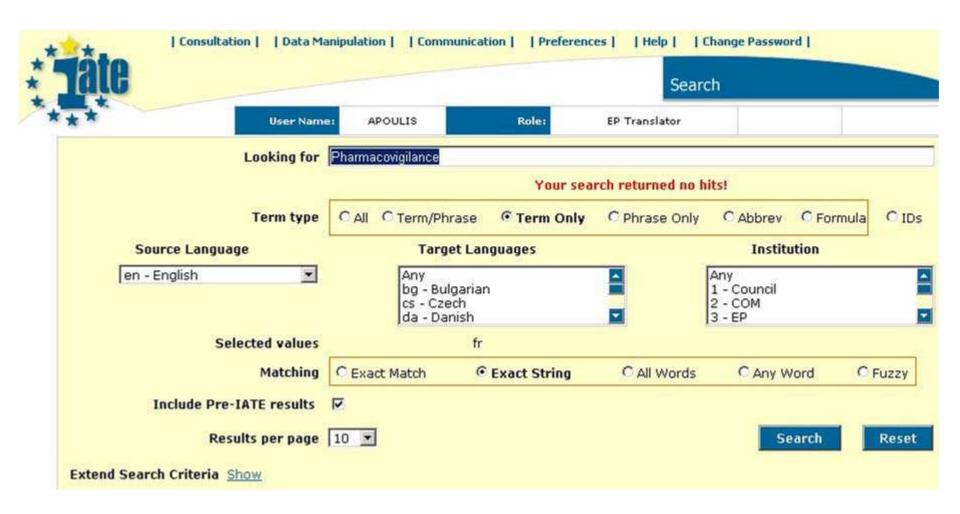
- The IATE (InterActive Terminology for Europe) database is a dynamic base designed, principally, to support the multilingual drafting of EU texts, legal texts in particular.
- IATE best practices: <u>http://tradunit2/units/terminology/wp-content/uploads/2008/12/best-practice-forterminologists-20080515-rev1.doc</u>
- General input criteria: Added value, relevance, avoidance of duplicates, Accuracy of data, Single concept, Minimum information (including definition and/context), Intellectual Property Rights







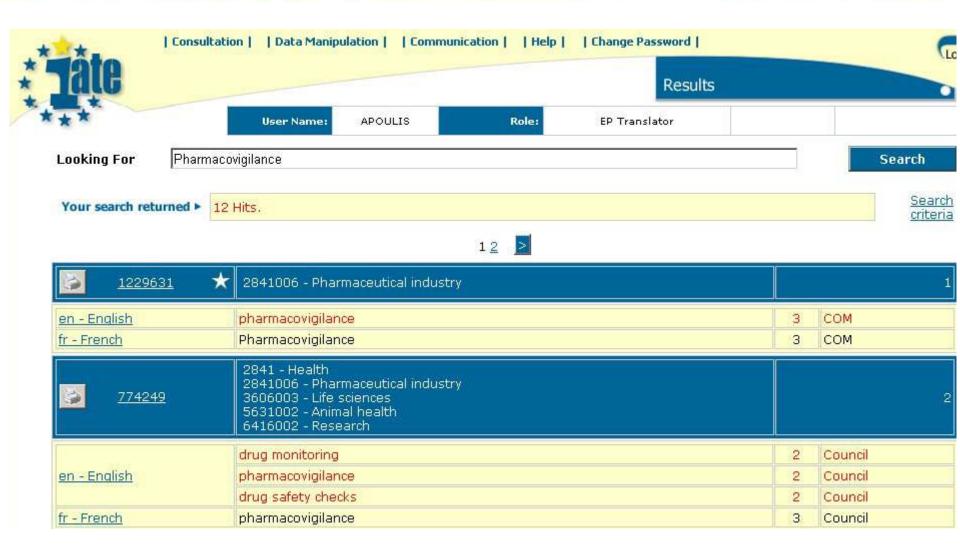
Termbases, Glossaries, Dictionaries







Termbases, Glossaries, Dictionaries





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Termbases, Glossaries, Dictionaries

- Most units have large collections of glossaries assembled in-house or retrieved from other resources (e.g. company web-sites, universities etc.)
- Glossaries offered on the Terminology Service's website
- The HU unit uses the Eurovoc thesaurus for the classification and organisation of their glossaries
 - Eurovoc is a multilingual thesaurus covering the fields in which the European Communities are active;
 - http://europa.eu/eurovoc/







Termbases, Glossaries, Dictionaries



Proactive Terminology



One of the most important goals of our team is to simplify and streamline Units workflow. To realize this purpose, we intend to provide Units with real time terminology about specific fields in every language, in order to improve the quality and the efficiency of the translations.

We intend to provide real time terminology in three ways:

- Anticipating workflow from the Commission.
- Getting bilingual lists of terms on specific domains.
- Working with automatic extractions tools for getting term lists.

List of expected topics to be discussed in Parliament in 2010 - last update 27 May 2010

Click to open an Excel file with information (sorted by month) about expected topics to be discussed this year.

RECENTLY ADDED GLOSSARIES in the following sections:

<u>Economics</u>, <u>Finance etc.</u> - added Apr 07 - EL, EN.

One bilingual glossary has been added. XRIMA is a Greek-English dictionary of modern business.



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Termbases, Glossaries, Dictionaries



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- Oxford Language
- Dictionaries Online DE->EN

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- FUSE Gepro
- Info Shout! Info Googlex Info

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Info

Info

- Profil
- · Elise





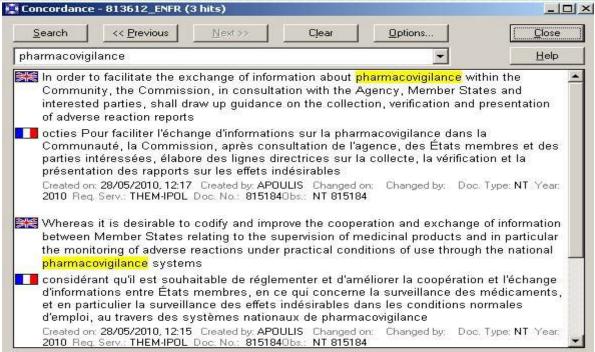


Bilingual Concordancers: TWB, Euramis, FullDoc



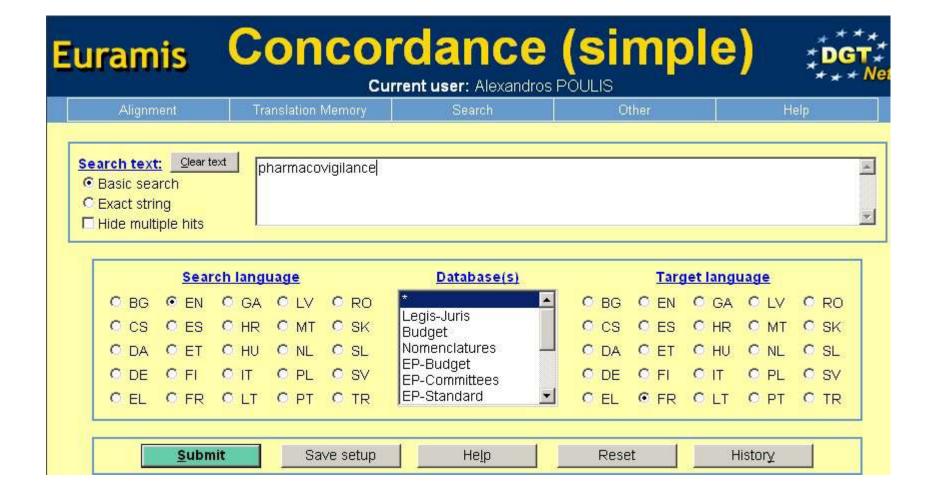
Whereas it is desirable to codify and improve the cooperation and exchange of information between Member States relating to the supervision of medicinal products and in particular the monitoring of adverse reactions under practical conditions of use through the national pharmacovigilance systems

considérant qu'il est souhaitable de réglementer et d'améliorer la coopération et l'échange d'informations entre États membres, en ce qui concerne la surveillance des médicaments, et en particulier la surveillance des effets indésirables dans les conditions normales d'emploi, au travers des systèmes nationaux de pharmacovigilance











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Translation Memory

Current user: Alexandros POULIS

Search text: pharmacovigilance

Database Name	Search Lng.	Target	Max. Results	Execution time
*	EN	FR	30	0 seconds

New Search

History

Page(s): 1 Search more...

Showing 30 hits / 30 (more results available)

TM: Legis-Juris Doc. No.: 22001D0153 Year: 2001 SL: EN (Direct) Trans: unknown Doc. Type: Instrument bodies agreements

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Chapter VIa "Pharmacovigilance" of Council Directive 81/851/EEC on the approximation of the laws of the Member States relating to veterinary medicinal products(2) is to be incorporated into the Agreement,

EN: Commission Directive 2000/37/EC of 5 June 2000 amending FR: La directive 2000/37/CE de la Commission du 5 juin 2000 modifiant le chapitre VI bis de la directive 81/851/CEE du Conseil concernant le rapprochement des législations des États membres relatives aux médicaments vétérinaires(2) doit être intégrée à l'accord,

TM: Legis-Juris Doc. No.: 31993L0039 Year: 1993 SL: EN (Direct) Trans: unknown Doc. Type: Directive

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EN: Whereas it is desirable to codify and improve the cooperation and exchange of information between Member States relating to the supervision of medicinal products and in particular the monitoring of adverse reactions under practical conditions of use through the national pharmacovigilance systems,

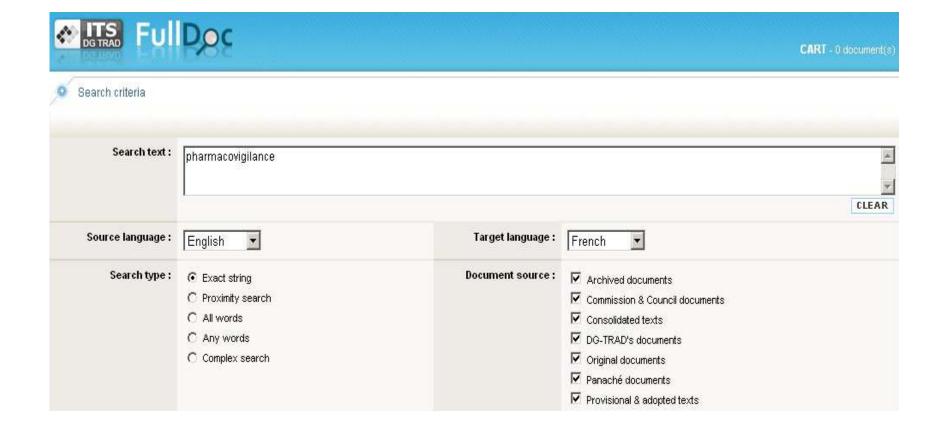
EN: Authorization shall be valid for five years and shall be renewable for five-year periods, on application by the holder at least three months before the expiry date and after consideration moins trois mois avant la date d'expiration, après examen par by the competent authority of a dossier containing in particular details of the data on pharmacovigilance and other information des données de la pharmacovigilance et les autres informations relevant to the monitoring of the medicinial product.

FR: considérant qu'il est souhaitable de réglementer et d'améliorer la coopération et l'échange d'informations entre États membres, en ce qui concerne la surveillance des médicaments, et en particulier la surveillance des effets indésirables dans les conditions normales d'emploi, au travers des systèmes nationaux de pharmacovigilance,

FR: L'autorisation est valable pour cinq ans et renouvelable par périodes de cinq ans, sur demande introduite par le titulaire au l'autorité compétente d'un dossier reprenant notamment l'état pertinentes pour la surveillance du médicament.













Back to search page | Take CART - 0 document(s

Displaying 1 to 10 (of 312 occurrence(s))

Go to next DualDisplay - Pages: | 2 3 4 5 6 7 8 9 10 >>

Number	<u>Dual</u> display	Source document	Target document	Doc. type	Creation date	<u>Last</u> update	Extract	Euramis align
Ť	♦	797090.doc	797090FR.doc	PA	21/01/2010	11/01/2010	urité alimentaire sur la proposition de règlement du Parlement européen et du Conseil modifiant, en ce qui concerne la pharmacovigilance des médicaments à usage humain, le règlement (CE) n° 726/2004 établissant des procédures communautaires pour l'autorisation et la surveillance en ce qui concerne les médicaments à usage humain et à usage vétérinaire, et instituant une Agence européenne des médicaments	П
2	\$	SEC(2008) 2671.doc	SEC(2008) 2671.doc		10/12/2008	17/12/2008	ORKING DOCUMENT Accompanying document to the Proposal for a REGULATION OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL amending, as regards pharmacovigilance of medicinal products for human use, Regulation (EC) No 726/2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency and the Proposal for a DIRECTIVE OF THE EUROP	П
3	*	798819EN.doc	798819FR.doc	PR	14/01/2010	08/03/2010	TitreType>DRAFT REPORT on the proposal for a regulation of the European Parliament and of the Council amending, as regards pharmacovigilance of medicinal products for human use, Regulation (EC) No 726/2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency (COM(2008)0664 – C6 0515/2008 –	П
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> Back to search pa

"pharmacovigilance"

(occurrence 1/27)

> Go to next occurrence

> Go to previous occurrence

(To find a specific text Press Ctrl + F)

d'euros dans l'UE.

l'UE et le coût total des effets indésirables pour la société s'élève à 79 milliards

> Back to result page

Medicines regulation and particularly the system of authorising medicines are La réglementation en matière de médicaments, et plus particulièrement le critical for ensuring that only high quality, efficacious medicines of acceptable système d'autorisation des médicaments, sont essentiels pour garantir que seuls safety are allowed on the EU market. However, the established limitations of drug des médicaments de grande qualité, efficaces et d'un niveau acceptable de development, particularly the limitations of clinical trials in terms of their size, sécurité soient autorisés sur le marché communautaire. Toutefois, en raison des duration and controlled conditions means that some side effects will only be limites imposées à la mise au point des médicaments, et notamment aux essais detected after a medicine has been authorised and entered the market.

Pharmacovigilance is the science and activities relating to the detection, La pharmacovigilance est une science qui englobe les activités relatives à la assessment, understanding and prevention of adverse effects of drugs and comprises: collecting and managing data on the safety of medicines, scrutinising data and making decisions with regard to safety issues, acting to protect public health (including regulatory action), and, communicating with stakeholders about the safety of medicines.

Key stakeholders involved in pharmacovigilance are:

carers of patients,

all healthcare professionals but particularly:

doctors (to advise patients, prescribe medicines and report ADRs)

pharmacists (to advise patients, dispense medicines and report ADRs)

nurses (to advise patients, administer medicines and report ADRs)

cliniques pour ce qui de leur taille, de leur durée et des conditions contrôlées, certains effets secondaires ne seront détectés qu'une fois le médicament

autorisé et mis sur le marché.

détection, l'évaluation, la compréhension et la prévention des effets indésirables des médicaments: la collecte et la gestion d'informations sur la sécurité des at the data to detect 'signals' (any new or changing safety issues), evaluating the médicaments, l'examen des données afin de détecter les «signaux» (toute question de sécurité, nouvelle ou en évolution), l'évaluation des données et la prise de décisions dans des questions de sécurité, l'adoption de mesures visant à protéger la santé publique (notamment des mesures réglementaires) et la communication avec les parties concernées au sujet de la sécurité des médicaments.

Les principales parties concernées par la pharmacovigilance sont:

patients as the users of medicines and families and carers as advisers and les patients, en tant que consommateurs de médicaments, ainsi que les familles et les soignants, en tant que personnes leur prodiquant conseils et soins;

tous les professionnels de la santé, mais plus particulièrement:

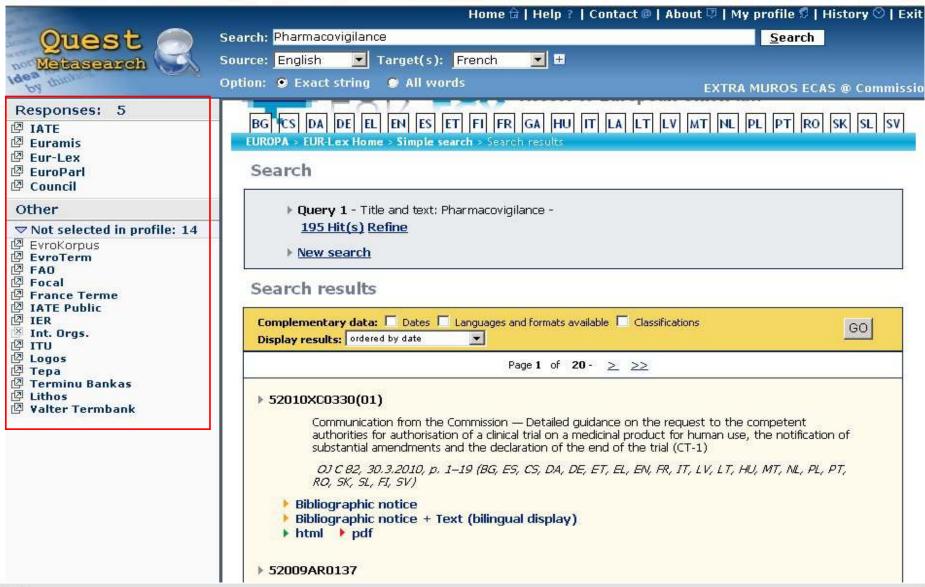
les médecins (pour conseiller les patients, prescrire des médicaments et notifier les effets indésirables)

les pharmaciens (pour conseiller les patients, délivrer des médicaments et notifier les effets indésirables)

les infirmières (pour conseiller les patients, administrer des médicaments et notifier les effets indésirables)













Bilingual Concordancers: MyMemory, TAUS search, Linguee

- New generation online concordancers
- Based on crowdsourcing and webcrawling techniques
- They also offer integration to CAT-Tools
- Examples: MyMemory, Linguee, TAUS widgets







Bilingual Concordancers: MyMemory, TAUS search, Linguee



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How to contribute? Delete wrong alignments, vote translations, add new ones.

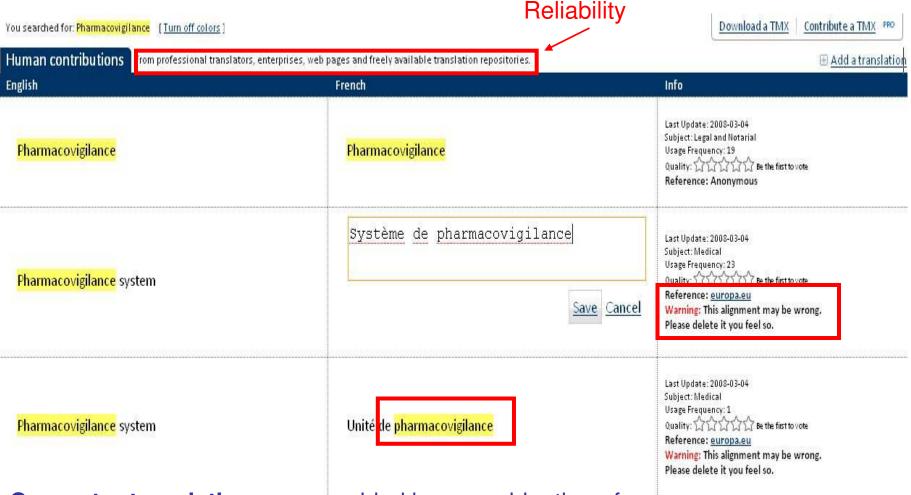
te tengo super abandono (Spanish>French) | arachidi (Catalan>Italian) | intergen (English>Swedish) | 的類取工 (>) |
paternalistic (English>Japanese) | kórea (Slovak) (Zech) | para confirmação, escrevo aqui novamente : (Portuguese>German) |
trade (English>Spanish) | la paix (French>Kirundi) | 的類取工 (Chinese Simplified>) | eu me stresso (Portuguese>German) |

ceroso (Portuguese>English) | vendedor técnico (Portuguese>German) | 的類取工 (Japanese>) | giudizio (Italian Spanish)





Bilingual Concordancers: MyMemory, TAUS search, Linguee



Computer translations are provided by a combination of MyMemory's statistical machine translator, Google, Systran and Worldlingo





corpus

Bilingual Concordancers: MyMemory, TAUS search, Linguee

TAUS SEARCH

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Computed translations(?))			
corpus (noun) corpus (n	oun) (56%), corps (noun) (15%), caverr	neux	(adjective) (9%)	
English (United Kingdom) Segment		French (France) Segment	
At mid-cycle, high levels of and ovulation.	of LH trigge corpus uteum fermation	n	En milieu de cycle, le pie de LH déclenche la formation du corps jaune et l'ovulation.	iX
stimulate their ovaries, to	vomen who have received treatment trigger ovulation (the release of egg elopment of a special structure on the that helps pregnancy.	s by	Ovitrelle est utilisé chez les femmes qui ont suivi un traitement de stimulation ovarienne, afin de déclencher l'ovulation (libération d'ovules par les ovaires) et la formation d'une structure spécifique autour de l'ovaire (16 corps) aune) qui favorise la grossesse.	iX
100 001	es its particular concern at the detent ures and the suspension of habeas		À cet égard, l'Union européenne est tout particulièrement préoccupéee par la détention sans jugem ent de personnalités politiques et la suspension de l'habeas corpus.	iX
where it causes the muscle in the spongy tissue of the penis (the			Durant la stimulation sexuelle normale, de la GMPc est produite dans le pénis, elle y provoque le relâchement du muscle lisse du tissu spongieux du pénis (corpora cavernosa), favorisant l'afflux sanguin dans le corps caverneux qui produit l'érection.	iΧ





corpus

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TAUS SEARCH

_	English (United Kingdom) >	French (France) Search			
	Industry:	gal Services			
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	Part of speech:	Any			
Computed translations(?)					
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	es its particular concern at the detenti ures and the suspension of habeas	on À cet égard, l'Union européenne est tout particulièrement préoccupéee par la détention sans jugement de personnalités politiques et la suspension de l'habeas corpus .	X		
of them - voted for the Wa regret the absence of the they felt a strong commitr	n the European Free Alliance - a majo tson report; not that they did not also habea <mark>s corpus amendment, but tha</mark> nent in conscience, as we all do, to cial space going forward, and though	- une majorité d'entre eux - ont voté en faveur du rapport Watson; t non pas qu'il n'aient pas regretté, eux aussi, l'absence de l' amendement relatir à l'habeas corpus, mais ils ont ressenti, en	i X		
corpus and President Bu	s now seeking to bar claims for habe sh and his henchmen trying to stifle th would ban inhuman treatment of U	the demandes d'habeas corpus et le président Bush et ses hommes	i X		





Bilingual Concordancers: MyMemory, TAUS search, Linguee

What Where does the text content come from?

with L expre: Our most important source is the bilingual web. Other valuable sources include EU documents and senter patent specifications.



pharmacovigilance

Search

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[...] expiry date and after consideration by the competent authority of a [...] der Unterlagen, die insbesondere eine Übersicht über den Stand der Angaben zur Pharmakovigilanz und die übrigen für die dossier containing in particular details of the data on pharmacovigilance and other information relevant to the monitoring of the medicinal product. Arzneimittelüberwachung maßgebenden Informationen enthalten. E+ English: eur-lex.europa.eu/LexUriServ/LexUrl...0083:EN:HTML G+ German: eur-lex.europa.eu/LexUriServ/LexUri...0083:DE:HTML All marketing authorisation holders must designate a qualified person Jeder Inhaber einer Genehmigung für das Inverkehrbringen eines Arzneimittels benennt eine für die Arzneimittelüberwachung responsible for pharmacovigilance, who must establish and maintain a central information system concerning the side-effects of the medicinal verantwortliche Person, die ein zentrales Informationssystem für die product in question. Nebenwirkungen des Arzneimittels einrichtet. English: europa.eu/scadplus/leg/en/lvb/l21230.htm □ German: europa.eu/scadplus/leg/de/lvb/l21230.htm. Thirdly, the technical aspects of the proposed directive must be Dritte Bemerkung: Unerläßlich ist ferner eine Vereinheitlichung der harmonised with documents already issued or currently being drawn up, technischen Aspekte des vorgeschlagenen Dokuments mit den bereits and in particular those relating to the ICH, those on GCP and those on verabschiedeten oder in der Erarbeitung befindlichen Dokumenten, pharmacovigilance. insbesondere betreffend die ICH, die GCP und die Et English: www.europarl.europa.eu/ Arzneimittelüberwachung. German: www.europarl.europa.eu/ Show more examples | Suggest translation Other translations of 'pharmacovigilance': pharmacovigilance - and the (unsolved) labelling problem. Pharmakovigilanz und das (ungelöste) Kennzeichnungsproblem. en/region/ulm/magazin/03428/index.html ☐ German: bio-pro.de/de/region/ulm/magazin/03428/index.html Pharmakovigilanz, der Effizienz von Arzneimitteln, dem pharmacovigilance, drug efficacy, knowledge management, education Wissensmanagement und der Ausbildung. and training. English: bio-pro.de/en/life/magazin/04250/index.html 🗁 German: bio-pro.de/de/life/magazin/04250/index.html The expedient assignment of the EMEA does not have an effect on its Die sinnvolle Zuordnung der EMEA ist für deren koordinative Arbeit in der coordinative role in pharmacovigilance since the criteria are based on Pharmakovigilanz folgenlos, da die Kriterien auf internationalen international medical guidelines. medizinischen Richtlinien basieren. English: bio-pro.defenfregion/ulm/magazin/01240/index.html □ German: bio-pro.de/de/region/ulm/magazin/01240/index.html







Online References

Wikipedia, Google scholar

Article Discussion



WIKIPEDIA The Free Encyclopedia

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Tiếng Việt

Definition

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Pharmacovigilance

From Wikipedia, the free encyclopedia

Pharmacovigilance (sometimes abbreviated PV or PVG) is the pharmacological science relating to the detection, assessment, understanding and prevention of adverse effects, particularly long term and short term side effects of medicines. [1] Generally speaking, pharmacovigilance is the science of collecting, monitoring, researching, assessing and evaluating information from healthcare providers and patients on the adverse effects of medications, biological products, herbalism and traditional medicines with a view to:

- · identifying new information about hazards associated with medicines
- preventing harm to patients.

The etymological roots are: *pharmakon* (Greek), "drug;" and *vigilare* (Latin), "to keep awake or alert, to keep watch."

Pharmacovigilance is particularly concerned with adverse drug reactions, or ADRs, which are officially described as: "A response to a drug which is noxious and unintended, and which occurs at doses normally used... for the prophylaxis, diagnosis or therapy of disease, or for the modification of physiological function." [2]

Contents [hide]

- 1 Risks of medical treatment
- 2 Terms commonly used in drug safety
- 3 Finding the risks of drugs
- 4 Spontaneous reporting
- 5 Other reporting methods
- 6 International collaboration
- 7 Pharmacovigilance by region
 - 7.1 Europe
 - 7.2 United States
- 8 See also
- 9 References
- 10 External links

Related Terms

Pharmacovigilance is gaining importance for doctors and scientists as the number of stories in the mass media of drug recalls increases. [citation needed]

Because clinical trials involve several thousand patients at most; less entire months and ADRs are often unknown at the time a drug enters the market. Even very severe ADRs set that liver damage are often undetected because study populations are small. Postmarketing pharmacovigilane these tools such as data mining and investigation of case reports to identify the relationships between drugs and these.

Pharmacoenvironmentology: A branch of environmental pharmacology and a form of pharmacovigilance which deals entry of chemicals or drugs into the environment after elimination from humans and animals post-therapy. It deals specifically with those pharmacological agents that have impact on the environment via elimination through living



M



Wikipedia, Google scholar





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Pharmacovigilance

En France, « La pharmacovigilance a pour objet la surveillance du risque d'effet indésirable résultant de l'utilisation des médicaments et produits à usage humain » (article R. 5121-150 🗗 du code de la santé publique.)

Elle vise à garantir la sécurité d'emploi des médicaments. Elle repose sur le signalement des effets indésirables par les professionnels de santé (obligation leur est faite selon le code de la santé publique de signaler tout effet indésirable médicamenteux grave ou inattendu) et les industriels.

Les professionnels de santé déclarent les effets indésirables aux 31 centres régionaux de pharmacovigilance 🗗 (CRPV), situés dans des structures hospitalières, et qui ont chacun leur propre compétence géographique. Les évènements sont évalués et enregistrés, puis transmis à l'Afssaps (Agence française de sécurité sanitaire des produits de santé), qui coordonne l'ensemble du système.

Si un effet indésirable est transmis au laboratoire pharmaceutique propriétaire du médicament, il doit le déclarer directement à l'Afssaps sous 15 jours si l'effet indésirable est grave (décès, mise en jeu du pronostic vital, hospitalisation ou prolongation d'hospitalisation, invalidité, anomalie ou malformation congénitale, autre critère médicalement important). Les cas de Pharmacovigilance ne répondant pas à ces critères sont dit "non graves" et seront transmis aux Autorités Compétentes par le biais des rapports périodiques.

Les différentes agences nationales en charge du médicament ont organisé un réseau d'échange d'informations sur la pharmacovigilance. Un effet indésirable grave constaté pour un même médicament en Allemagne et en Grande-Bretagne est donc transmis en France et des recoupements sont faits. Cela peut amener à modifier les conditions d'utilisation du produit, à les restreindre voire à retirer le produit du marché. Les professionnels de santé peuvent être avertis de manière rapide par une Lettre aux prescripteurs, ou Dear Doctor Letter (DDL), envoyée par le laboratoire.

Les études cliniques de Phase IV, qui ont lieu après la mise sur le marché d'un médicament, sont une sorte de pharmacovigilance.

Les données de pharmacovigilance pour un médicament sont rassemblées et analysées par le laboratoire dans un document appelé Periodic Safety Update Report (PSUR), qui est fourni régulièrement aux autorités de santé (l'AFSSAPS pour la France) : tous les 6 mois pour un médicament nouveau, puis à terme tous les 3 ans (la sécurité d'utilisation du produit étant vérifiée sur de larges populations).







Pharmacovigilance Search Advanced Scholar Search Scholar Preferences

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Accelerating Preclinical CMC The SMi Group December 8

IP Drug, W Successful, P Risk, D ... - Biotechnology Law ..., 2009 - liebertonline.com Biopharma Executives 5635 Fishers Lane JW Marriott Hotel Rockville, MD 20852 1331 Pennsylvania Avenue Tel: (800) 332-2181 Washington, DC Web: www.windhover.com/FDA-CMS ... Intellectual Property and The SMi Group December 8 ... Generics Great Guildford Business Square ... All 2 versions

Smith v. PFIZER INC.

Dist. Court, MD Tennessee, Nashville Div., 2010 - Google Scholar

... The defendants have offered Weiss Smith as an expert in epidemiology, pharmacoepidemiology, and pharmacovigilance. ... Furthermore, the defendants do not plan to have Weiss Smith testify regarding the adequacy of defendants' pharmacovigilance practices or conduct. ...

The European Agency for the Evaluation of Medicines and European regulation of pharmaceuticals

JS Gardner - European Law Journal, 2009 - interscience, wiley, com-

... The applicant must apply at least three months before the end of authorisation, and the Agency reviews a dossier 'containing up-to-date information on pharmacovigilance' (reactions to the drug in actual use). ... C Pharmacovigilance ... Cited by 20 - Related articles

Quaid v. Baxter Healthcare Corp.

910 NE 2d 1236 - III: Appellate Court, 1st Dist., 3rd Div., 2009 - Google Scholar

... The last known addresses of the people significantly involved in pharmacovigilance and adverse events related to the subject Heparin products before March 2006 are in Pennsylvania and Park Ridge, Illinois. The Baxter employees ... Related articles - All 2 versions





Terminology Management Terminology Toolbar



 Permitting to copy and paste any new term and its translation into a Word table while translating in Word for later import into IATE after validation by the units' terminologists

 Repository files used for gathering, sorting and validating new terms for IATE







Terminology Management Terminology Toolbar





The aim of the Commission proposal is to limit the exposure of consumers to pharmacologically active substances intended for use in veterinary medicinal products for food producing animals as well as residues thereof in foodstuffs of animal origin.

Durch den Kommissionsvorschlag soll die Exposition der Verbraucher gegenüber pharmakologisch wirksamen Stoffen, die für die Verwendung in Tierarzneimitteln für zur Nahrungsmittelerzeugung genutzte Tiere vorgesehen sind, ebenso beschränkt werden wie das Auftreten von Rückständen dieser Stoffe in Nahrungsmitteln tierischen Ursprungs.

This includes foodstuffs imported from third countries.

Dies schließt aus Drittländern eingeführte Nahrungsmittel ein.







Repository Table

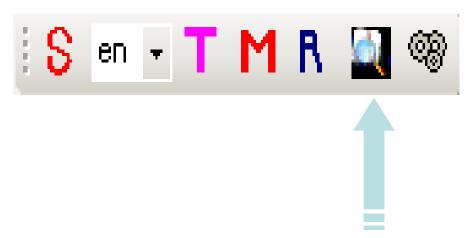
Source language	Source term	Target term	Author	Target file name	Date/time
en	exposure	Exposition	tdourda	R:\Terminology\presentations\macro.doc	22/09/2009 10:18:32
en	foodstuffs	Nahrungsmittel	tdourda	R:\Terminology\presentations\macro.doc	22/09/2009 10:18:33
en	compromise	Kompromiss	tdourda	R:\Terminology\presentations\macro.doc	22/09/2009 10:18:33
en	readability	Lesbarkeit	tdourda	R:\Terminology\presentations\macro.doc	22/09/2009 10:18:33







 Direct IATE search in WORD by simple selection of words



Example:

The industrial revolution in Brazil is leading to rapid deforestation of the Amazon basin.







Terminology Management Terminology Toolbar





Search results for: deforestation (en>de)

Source Term	Target Term	Inst.	Domains	Definition	Reference
deforestation	Abholzung	сом	AGRICULTURE, FORESTRY AND FISHERIES, Building and public works		BTP,Term Raumordnung und Städtebaus,1978
deforestation	Entwaldung	сом	AGRICULTURE, FORESTRY AND FISHERIES, Building and public works		BTP, Term Raumordnung und Städtebaus, 1978
deforestation	Abforstung	сом	AGRICULTURE, FORESTRY AND FISHERIES, Building and public works		Landmaschinen und landwirtschaftliche Ausrüstungen,CEMAGREF,19
deforestation	Kahlschlag	сом	AGRICULTURE, FORESTRY AND FISHERIES, Building and public works		Landmaschinen und landwirtschaftliche Ausrüstungen,CEMAGREF,19
deforestation	Rodung	сом	AGRICULTURE, FORESTRY AND FISHERIES	Dauemde oder vorübergehende Zweckentfremdung des Waldareals,auch wenn es sich um nicht bestockte Flächen handelt.	Duc,Glossar LFI,1998





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Translation

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Pharmacotherapy is the treatment of disease through the administration of drugs.[1] As such, it is considered part of the larger category of therapy.

Translate from:	English	-
Translate into:	French	•

English to French translation

La pharmacothérapie est le traitement de la maladie par l'administration de médicaments. [1] En tant que tel, il est considéré comme faisant partie de la catégorie plus large de la thérapie.

E Contribute a better translation

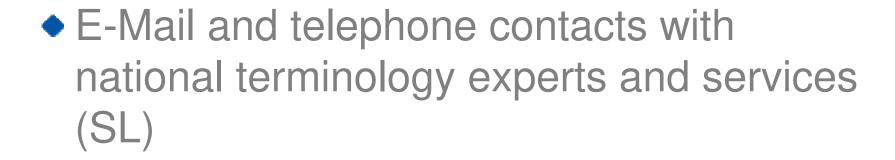
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- The goal of term mining is to automatically extract relevant terms from a document or collection of documents (corpus)
- Terminology extraction algorithms are based on statistical, linguistic or combined approaches.
- The Terminology Coordination service tests various terminology extraction tools
 - Xerox, Synchroterm, Multicorpora





- Terminology extraction tools are often provided by CAT Tools vendors for better integration to the translation workflow:
 - Alchemy term extraction API, SDL MultiTerm Extract 2009, Heartsome term extraction service and Araya bilingual term extraction tool etc.
- Online terminology extraction tools
 - Yahoo Term Extraction Web Service







THANK YOU



