



PUBLIC HEALTH

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PHARMACEUTICALS - COMMUNITY REGISTER

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❖ Procedures for centrally authorised medicinal products¹

Community Register of medicinal products for human use²

Active	By EU number	Alphabetical
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Withdrawn, suspended, expired or not renewed	By EU number	Alphabetical
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Refused		Alphabetical
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Community Register of orphan medicinal products for human use³

Active	By EU number	Alphabetical
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Withdrawn or expired	By EU number	Alphabetical
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Refused		Alphabetical
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Community Register of veterinary medicinal products⁴

Active	By EU number	Alphabetical
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Withdrawn, suspended, expired or not renewed	By EU number	Alphabetical
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Refused		Alphabetical
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❖ Procedures for nationally authorised medicinal products

Medicinal products for human use		Alphabetical
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Veterinary medicinal products		Alphabetical
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Adopted Commission Decisions of the last six months

Human medicinal products by ATC
Veterinary medicinal products by ATC

General index on active ingredient
General index on brand name
General index on Marketing Authorisation Holders and Sponsors

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- 1 Variation procedures which do not affect the Commission Decision granting the marketing authorisation (including its annexes) are no longer reflected in the Community Register of medicinal products since 1 April 2011.
- 2 The Register of medicinal products for human use authorised by the EU under the centralised procedure. Published in accordance with Article 13 of Regulation (EC) No 726/2004 .
- 3 The Register of orphan medicinal products for human use designated and published by the EU in accordance with Article 5 of Regulation (EC) N141/2000.
- 4 The Register of veterinary medicinal products authorised by the EU under the centralised procedure. Published in accordance with Article 38 of Regulation (EE) 726/2004.
- 5 The list of Exceptional Marketing Authorisations is published in accordance with Article 126a of Directive 2001/83/EC of the European Parliament and of the Council of 6 november 2001 on the community code relating to medicinal products for human use

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