European Commission
Public Health

News and updates on pharmaceuticals

EudraLex - Volume 5 - Pharmaceutical legislation Medicinal Products for veterinary use

Volume 5 of the publications "The rules governing medicinal products in the European Union" compiles the body of European Union legislation in the pharmaceutical sector for medicinal products for veterinary use.

Directives

Table Eudralex

2009/53/EC	Directive 2009/53/EC of the European Parliament and of the Council of 18 June 2009 amending Directive 2001/82/EC and Directive 2001/83/EC, as regards variations to the terms of marketing authorisations for medicinal products. (Official Journal L 168, 30/6/2009, p. 33 - 34).
2009/9/EC	Commission Directive 2009/9/EC of 10 February 2009 amending Directive 2001/82/EC of the European Parliament and of the Council on the Community code relating to medicinal products for veterinary use (Official Journal L 44, 14/2/2009 p. 10 - 61).
2006/130/EC	Commission Directive 2006/130/EC of 11 December 2006 implementing Directive 2001/82/EC of the European Parliament and of the Council as regards the establishment of criteria for exempting certain veterinary medicinal products for food -producing animals from the requirement of a veterinary prescription was published in the Official Journal on 12 December 2006.
2001/82/EC	Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products (Official Journal L 311, 28/11/2001 p. 1 - 66). (consolidated version : 18/7/2009)
2004/28/EC	Amended by Directive 2004/28/EC of the European Parliament and of the Council of 31 March 2004 amending Directive 2001/82/EC on the Community code relating to veterinary medicinal products (Official Journal L 136, 30/4/2004 p. 58 - 84).
Consolidated Directive	Consolidated Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products as amended by Directive 2004/28/EC.
91/412/EEC	Commission Directive 91/412/EEC of 23 July 1991 laying down the principles and guidelines of good manufacturing practice for veterinary medicinal products (Official Journal L 228, 17/8/1991 p. 70 - 73).
90/167/EEC	Council Directive 90/167/EEC of 26 March 1990 laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Community (Official Journal L 92, 7/4/1990).