



EUROPEAN COMMISSION

Brussels, 10.8.2011  
C(2011)5945 final

**COMMISSION IMPLEMENTING DECISION**

**of 10.8.2011**

**withdrawing, at the holder's request, the marketing authorisation for "Yarvitan - Mitratapide", a medicinal product for veterinary use, granted by Decision C(2006)5549**

(Text with EEA relevance)

(ONLY THE DUTCH TEXT IS AUTHENTIC)

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THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency<sup>1</sup>,

Having regard to the application submitted by Janssen Pharmaceutica N.V. on 21 April 2011 with a view to the withdrawal of the marketing authorisation for the medicinal product "Yarvitan - Mitratapide"

Whereas:

- (1) The medicinal product "Yarvitan - Mitratapide", entered in the Community register of medicinal products under numbers EU/2/06/063/001-003 is authorised by Commission Decision C(2006)5549 of 14 November 2006.
- (2) Following the holder's request, that authorisation should be withdrawn,

HAS ADOPTED THIS DECISION:

### *Article 1*

At the holder's request, the marketing authorisation for "Yarvitan - Mitratapide", a medicinal product for veterinary use granted by Decision C(2006)5549 of 14 November 2006, is withdrawn.

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<sup>1</sup> OJ L 136, 30.4.2004, p. 1.

*Article 2*

This Decision is addressed to Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse, België.

Done at Brussels, on 10.8.2011.

*For the Commission*  
*Paola TESTORI COGGI*  
*Director-General*