

IPAR



Publicly Available Assessment Report for a Veterinary Medicinal Product

HyperCard 10 mg Coated Tablets for Cats

PRODUCT SUMMARY

EU Procedure Number	IE/V/0506/001 (formerly UK/V/0174/001)
Name, Strength, Pharmaceutical Form	HyperCard 10 mg Coated Tablets for Cats
Active Substances(s)	Diltiazem
Applicant	Dechra Regulatory B.V., Handelsweg 25, 5531 AE Bladel, Netherlands
Legal Basis of Application	Not Currently Available
Target Species	Cats
Indication For Use	
ATC Code	QC08DB01
	10 November 2008 (IE)

PUBLIC ASSESSMENT REPORT

The public assessment report reflects the scientific conclusion reached by the HPRA at the end of the evaluation process and provides a summary of the grounds for approval of the marketing authorisation for the specific veterinary medicinal product. It is made available by the HPRA for information to the public, after the deletion of commercially confidential information. The legal basis for its creation and availability is contained in Article 25.4 of EC Directive 2001/82/EC as amended by Directive 2004/28/EC for veterinary medicinal products. It is a concise document which highlights the main parts of the documentation submitted by the applicant and the scientific evaluation carried out by the HPRA leading to the approval of the product for marketing in Ireland.

The Summary of Product Characteristics (SPC) for this product is available on the HPRA's website.