IPAR

Publicly Available Assessment Report for a
Veterinary Medicinal Product

Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Swine

PRODUCT SUMMARY

EU Procedure number	(formerly FR/V/0174/001)
Name, strength and pharmaceutical form	Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Swine
Active substances(s)	Florfenicol
Applicant	Intervet Ireland Limited Magna Drive Magna Business Park, Citywest Road Dublin 24 Ireland
Legal basis of application	Full application (Article 12(3) of Directive No 2001/82/EC)
Date of Authorisation	31 August 2007 (IE)
Target species	Pigs
Indication for use	
ATCvet code	QJ01BA90
Concerned Member States	BG, CY, EL, IT, PT, SK, ES, UK(NI)

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.