### **IRISH MEDICINES BOARD ACT 1995**

#### EUROPEAN COMMUNITIES (ANIMAL REMEDIES) (No. 2) REGULATIONS 2007

(S.I. No. 786 of 2007)

VPA: **10545/030/001** Case No: 7005586

The Irish Medicines Board in exercise of the powers conferred on it by Animal Remedies (No. 2) Regulations (S.I. No. 786 of 2007) hereby grants to:

### Janssen Cilag Ltd.

### 50-100 Holmers Farm Way, High Wycombe, Buckinghamshire HP12 4EG, United Kingdom

an authorisation, subject to the provisions of the said Regulations and the general conditions of the attached authorisation, in respect of the Veterinary Medicinal Product:

#### Furexel Combi, oral paste

The particulars of which are set out in Part 1 and Part 2 of the said Schedule. The authorisation is also subject to any special conditions as may be specified in the said Schedule.

The authorisation, unless revoked, shall continue in force from 13/01/2009 until 22/11/2010.

Signed on behalf of the Irish Medicines Board

A person authorised in that behalf by the said Board.

(NOTE: This authorisation replaces any previous authorisation in respect of this product which is now null and void.)

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## Part II

# **Summary of Product Characteristics**

**4 CLINICAL PARTICULARS** 

5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES

6 PHARMACEUTICAL PARTICULARS

9 DATE OF THE FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

23rd November 2005

10 DATE OF REVISION OF THE TEXT

13th January 2009