Bolfo, 2.5mg/g, Kožní sprej, roztok

Authorised

- Propoxur

Product identification

Medicine name:
Bolfo, 2.5mg/g, Kožní sprej, roztok

Active substance:
Propoxur

Target species:
Dog
Cat

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Propoxur
2.50 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous spray, solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AE02

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Czechia

Available in:
Czechia

Package description:
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Elanco Animal Health GmbH

Marketing authorisation date:
29/05/2009

Manufacturing sites for batch release:
Tosvar S.r.l.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
99/017/09-C

Date of authorisation status change:
29/05/2009

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.