Advantix Spot-On (400 mg + 2000 mg)/4 ml; roztwór do nakrapiania

Authorised

This information is not available for this product.

Product identification

Medicine name:
Advantix Spot-On (400 mg + 2000 mg)/4 ml; roztwór do nakrapiania

Active substance:
This information is not available for this product.

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:
Spot-on solution

Withdrawal period by route of administration:

Cutaneous use:

- **Dog**

- All relevant tissues. no withdrawal period

The withdrawal period does not apply.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC54

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

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:
25/02/2004

Manufacturing sites for batch release:
KVP Pharma+Veterinär Produkte GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1414

Date of authorisation status change:
25/02/2004

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

Documents

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.

Summary of Product Characteristics

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