

Dolorex 10 mg/ml Solution for Injection for horse, dog and cat

Authorised

- Butorphanol tartrate

Product identification

Medicine name:

Dolorex 10 mg/ml Solution for Injection for horse, dog and cat

Active substance:

Butorphanol tartrate

Target species:

Dog

Horse

Cat

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Butorphanol tartrate

14.60 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intravenous use:**

-

Horse

- Meat and offal. 0 day

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

26/10/2007

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

24777

Date of authorisation status change:

26/10/2007

Reference member state:

Germany

Procedure number:

DE/V/0354/001

Concerned member states:

Belgium Czechia Denmark Finland France Greece Hungary Ireland
Luxembourg Netherlands Norway Portugal Slovakia Sweden
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 29/12/2024

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Package Leaflet

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Combined File of all Documents

English (PDF)

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Labelling

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