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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 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID2) 50 millilitre(s): unspecified outer container with 1 Vial (Glass) with 50 millilitre(s)
(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (Glass) with 10 millilitre(s)

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 Deutschland GmbH

Marketing authorisation date:
23/03/2007

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400992.00.00

Date of authorisation status change:
7/10/2011

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

Combined File of all Documents

English (PDF)

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