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DOLOREX

Authorised

- Butorphanol tartrate

Product identification

Medicine name:
DOLOREX

Active substance:
Butorphanol tartrate

Target species:
Dog
Horse (non-food-producing)
Cat

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Butorphanol tartrate
14.57 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Dog**
  - Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

Intravenous use:

- **Dog**
  - Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

- **Horse (non-food-producing)**
  - Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

- **Cat**
  - Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

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

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Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
Intervet International B.V.

Marketing authorisation date:
22/03/1993

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
22/03/1993

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

Documents

Combined File of all Documents

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