

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

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Horse (non food-producing)

- Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

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Cat

- Unspecified. 0 day

La somministrazione del medicinale deve essere effettuata esclusivamente dal medico veterinario

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised 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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