

Dolorex 10 mg/ml Solution for Injection for Horse, Dog and Cat

Not
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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

United Kingdom (Northern Ireland)

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:

MSD Animal Health UK Limited

Marketing authorisation date:

5/07/2007

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 01708/3041

Date of authorisation status change:

24/04/2025

Reference member state:

Germany

Procedure number:

DE/V/0354/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents

ie-puar-mr-iev0194001-dolorex-10-mgml-solution-for-injection-for-horse-d-en.pdf