Torphasol 10 mg/ml solution for injection for horses

Authorised

- Butorphanol tartrate

Product identification

Medicine name:
Torphasol 10 mg/ml solution for injection for horses

Active substance:
Butorphanol tartrate

Target species:
Horse

Route of administration:
Intravenous use

Product details

Active substance and strength:
Butorphanol tartrate
14.70 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:
Portugal

Available in:
Portugal

Package description:
Cardboard box with 1 clear glass vial (type I) of 20 ml with a grey butyl rubber stopper and an aluminium cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
27/11/2011

Manufacturing sites for batch release:
aniMedica GmbH
Industrial Veterinaria S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
374/01/11RFVPT

Date of authorisation status change:
25/08/2025

Reference member state:
Ireland

Procedure number:
IE/V/0246/001

Concerned member states:
Austria Belgium Denmark Estonia Finland France Germany Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Spain 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

Combined File of all Documents

English (PDF)
Published on: 24/08/2025
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