

Alvegesic vet. 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen

Authorised

- Butorphanol tartrate

Product identification

Medicine name:

Alvegesic vet. 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen

Alvegesic vet. 10 mg/ml šķīdums injekcijām zirgiem, suņiem un kaķiem

Active substance:

Butorphanol tartrate

Target species:

Dog

Cat

Horse

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

Product details

Active substance and strength:

Butorphanol tartrate

14.58 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Dog

-

Cat

Intravenous use:

-

Dog

-

Horse

- Meat and offal. 0 day

- Milk. 0 hour

-

Cat

Subcutaneous use:

-

Dog

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

Cardboard box with 1 glass (Type II) vial of 10 ml with a bromobutyl rubber stopper and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

V.M.D.

Marketing authorisation date:

27/07/2011

Manufacturing sites for batch release:

V.M.D.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/11/0044

Date of authorisation status change:

27/07/2011

Reference member state:

Austria

Procedure number:

AT/V/0003/001

Concerned member states:

Belgium Bulgaria Czechia Estonia Finland France Germany Hungary Iceland

Ireland Italy Latvia Lithuania Netherlands Norway Portugal Romania
Slovakia Slovenia Spain 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

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