BUPROXAN MULTIDOSE 0.3 MG/ML SOLUTION FOR INJECTION FOR DOGS, CATS AND HORSES

Authorised

- Buprenorphine hydrochloride

Product identification

Medicine name:
BUPROXAN MULTIDOSE 0.3 MG/ML SOLUTION FOR INJECTION FOR DOGS, CATS AND HORSES

Active substance:
Buprenorphine hydrochloride

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

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:

Buprenorphine hydrochloride
0.32 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Horse (non food-producing)**
  - Not applicable. no withdrawal period

Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AE01

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Clear colourless glass vial, 10 ml, Type I

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
9/04/2026

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 134875

Date of authorisation status change:
19/02/2026

Reference member state:
France

Procedure number:
FR/V/0520/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

Generic of:
600000035317

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

Documents

Combined File of all Documents

English (PDF)
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