Dexmedocord 0.5 mg/ml solution for injection for dogs and cats

Authorised

- Dexmedetomidine hydrochloride

Product identification

Medicine name:
Dexmedocord 0.5 mg/ml solution for injection for dogs and cats
Dexmedocord 0,5 mg/ml šķīdums injekcijām suņiem un kaķiem

Active substance:
Dexmedetomidine hydrochloride

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Dexmedetomidine hydrochloride
0.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM18

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Type I clear glass vial with a fluoropolymer coated chlorobutyl rubber stopper and an aluminium overseal. Cardboard box containing one 10 ml vial
Type I clear glass vial with a fluoropolymer coated chlorobutyl rubber stopper and an aluminium overseal. Cardboard box containing ten 10 ml vials

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Accord Healthcare B.V.

Marketing authorisation date:
12/01/2026

Manufacturing sites for batch release:
Fundacio Privada Dau

Responsible authority:

Food And Veterinary Service

Authorisation number:
V/SRP/26/0002

Date of authorisation status change:
12/01/2026

Reference member state:
Ireland

Procedure number:
IE/V/0888/001

Concerned member states:
Belgium Bulgaria Estonia France Germany Italy Latvia Lithuania Netherlands Poland Romania Slovakia Spain

United Kingdom (Northern Ireland)

Generic of:
600000003548

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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