Uriphex 50 mg/ml, oral solution for dogs

Authorised

- Phenylpropanolamine

Product identification

Medicine name:
Uriphex Vet 50 mg/ml Mixtúra, lausn Handa hundum
Uriphex 50 mg/ml, oral solution for dogs

Active substance:
Phenylpropanolamine

Target species:
Dog (bitch)

Route of administration:
Oral use

Product details

Active substance and strength:
Phenylpropanolamine
40.28 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Dog (bitch)**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG04BX91

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

Authorisation status:
Valid

Authorised in:
Iceland

Package description:
HDPE bottle closed with white polypropylene child-resistant cap and LDPE syringe adaptor. A 1 mL HDPE/polypropylene graduated syringe is supplied with each bottle.
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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:
30/08/2023

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

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/23/012/01

Date of authorisation status change:
30/08/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0383/001

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

This information is not available for this product.

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

Documents

Combined File of all Documents