Authorised

GABBROVET 140 MG/ML SOLUTION FOR USE IN DRINKING WATER, MILK OR MILK REPLACER FOR PRE-RUMINANT CATTLE AND PIGS

This information is not available for this product.

Product identification

Medicine name:

GABBROVET 140 MG/ML SOLUTION FOR USE IN DRINKING WATER, MILK OR MILK REPLACER FOR PRE-RUMINANT CATTLE AND PIGS

Gabbrovet 140 mg/mL, otopina za primjenu u pitkoj vodi, mlijeku ili mliječnoj zamjenici, za goveda prije uspostavljanja funkcije predželudaca i svinje

Active substance:

This information is not available for this product.

Target species:

Cattle

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

This information is not available for this product.

Pharmaceutical form:

Solution for use in drinking water

Withdrawal period by route of administration:

Oral use:

- . Cattle
 - Meat and offal. 20 day
- . Pig
 - Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Package description:

Box of 1 vial of 125 mL

Box of 1 vial of 250 mL

Box of 1 vial of 500 mL

Box of 1 vial of 1000 mL

Vial of 125 mL

Vial of 250 mL

Vial of 500 mL

Vial of 1000 mL

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:

Ceva Sante Animale

Marketing authorisation date:

21/03/2018

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/18-01/93

Date of authorisation status change:

21/03/2018

Reference member state:

France

Procedure number:

FR/V/0317/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Source URL: https://medicines.health.europa.eu/veterinary/700000029383