Source URL: https://medicines.health.europa.eu/veterinary/en/600000064337

Magniject 250 mg/ml Solution for Injection

Authorised

• Magnesium sulfate heptahydrate

Product identification

Medicine name:

Magniject 250 mg/ml Solution for Injection

Active substance:

Magnesium sulfate heptahydrate

Target species:

Cattle

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Magnesium sulfate heptahydrate 25.00 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration: Subcutaneous use:

Cattle

- Meat and offal. 0 day
- Milk. 0 hour

•

Sheep

- Meat and offal. 0 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA12CC02

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Magniject, a clear colourless solution, is marketed in 400 ml Amber Type III glass vial sealed with black rubber wad and aluminium screw cap.

Magniject, a clear colourless solution, is marketed in 400 ml polypropylene container sealed with bromobutyl bung and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation: Complete application (stand-alone) - Council Directive 81/851/EEC
Marketing authorisation holder: Norbrook Laboratories (Ireland) Limited
Marketing authorisation date: 1/10/1991
Manufacturing sites for batch release: Norbrook Laboratories Limited Norbrook Manufacturing Limited
Responsible authority: Health Products Regulatory Authority
Authorisation number: VPA22664/030/001
Date of authorisation status change: 1/10/1991
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics