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