Hexasol LA Solution for Injection

Authorised

- Flunixin meglumine
- Oxytetracycline dihydrate

Product identification

Medicine name:
Hexasol LA Solution for Injection

Active substance:
Flunixin meglumine

Oxytetracycline dihydrate

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Flunixin meglumine
33.18 milligram(s) / 1.00 millilitre(s)

Oxytetracycline dihydrate
323.52 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA56

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Available in:

Austria

Package description:

Supplied in 500 ml Type I/II, amber glass vial, sealed with bromobutyl rubber bung and aluminium cap.
Supplied in 250 ml Type I/II, amber glass vial, sealed with bromobutyl rubber bung and aluminium cap.
Supplied in 100 ml Type I/II, amber glass vial, sealed with bromobutyl rubber bung and aluminium cap.
Supplied in 50 ml Type I/II, amber glass vial, sealed with bromobutyl rubber bung and aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
7/10/2004

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00628

Date of authorisation status change:
7/10/2004

Reference member state:
Ireland

Procedure number:
IE/V/0151/001

Concerned member states:
Austria France Portugal

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

Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 24/11/2024
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Labelling

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Combined File of all Documents