

# Hexasol LA Solution for Injection

Authorised

- Flunixin meglumine
- Oxytetracycline dihydrate

## Product identification

**Medicine name:**

Hexasol LA Solution for Injection

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**Active substance:**

Flunixin meglumine

Oxytetracycline dihydrate

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**Target species:**

Cattle

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**Route of administration:**

Intramuscular use

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## 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)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. 28 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01AA56

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Available in:**

Ireland

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**Package description:**

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

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Fixed combination application (Article 13b of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

5/02/1999

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA22664/049/001

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**Date of authorisation status change:**

5/02/1999

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0151/001

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**Concerned member states:**

Austria France Portugal

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents