

Dihydrostreptomycin 25% Solution for Injection

Authorised

- Dihydrostreptomycin sulfate

Product identification

Medicine name:

Dihydrostreptomycin 25% Solution for Injection

Active substance:

Dihydrostreptomycin sulfate

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Dihydrostreptomycin sulfate

292.02 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

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Cattle

- Meat and offal. 30 day
- Milk. 60 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01GA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

A limpid, slightly yellow coloured, solution for injection packed in 100 ml vial, amber glass Type II, with butyl rubber stopper and non-reusable aluminium closure.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:

Alfasan Nederland B.V.

Marketing authorisation date:

1/10/1988

Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10980/004/001

Date of authorisation status change:

1/10/1988

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

Documents

Summary of Product Characteristics