

Oxytetracycline Injection 10%

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:

Oxytetracycline Injection 10%

Active substance:

Oxytetracycline hydrochloride

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Oxytetracycline hydrochloride
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 21 day
- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Amber type II glass 100 ml vial, sealed with a 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:

VPA10980/001/001

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