Bactidiaryl Oral Powder

Authorised

- Tetracycline hydrochloride
- NEOMYCIN SULFATE

Product identification

Medicine name:
Bactidiaryl Oral Powder

Active substance:
Tetracycline hydrochloride

NEOMYCIN SULFATE

Target species:
Cattle

Route of administration:
Oral use

Product details

Active substance and strength:
Tetracycline hydrochloride
0.25 gram(s) / 100.00 gram(s)

NEOMYCIN SULFATE
500000.00 international unit(s) / 100.00 gram(s)

Pharmaceutical form:

Oral powder

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01RA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Aluminium foil pack containing 100g of an oral powder, 50 sachets per box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:

Vetoquinol Ireland Limited

Marketing authorisation date:

1/10/1988

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.
Vetoquinol S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10983/006/001

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