Bovex 2,265 %, suspensie voor oraal gebruik voor runderen en schapen

Authorised

- Oxfendazole

Product identification

Medicine name:
Bovex 2,265 %, suspensie voor oraal gebruik voor runderen en schapen

Active substance:
Oxfendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Oxfendazole
22.65 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Milk. 6 day
  - Meat and offal. 16 day
- **Sheep**
  - Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
27/10/1997

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 8863

Date of authorisation status change:
9/09/2015

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

Documents

Combined File of all Documents

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