Bovilis Huskvac oral suspension for cattle

Authorised

- Dictyocaulus viviparus, third stage larvae, Live

Product identification

Medicine name:
Bovilis Huskvac oral suspension for cattle

Active substance:
Dictyocaulus viviparus, third stage larvae, Live

Target species:
Cattle

Route of administration:
Oral use

Product details

Active substance and strength:
Dictyocaulus viviparus, third stage larvae, Live
1000.00 Organisms / 1.00 Dose

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AN01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
25 ml larval suspension in 30 ml glass hydrolytic class type III bottles, closed with a metal screwcap with a PEP facedinlay.Each package presentation consists of 12 x 25 ml (1 dose) bottles.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet (Ireland) Limited

Marketing authorisation date:
22/12/2003

Manufacturing sites for batch release:
MSD Animal Health UK Limited
Intervet International B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/081/001

Date of authorisation status change:
22/12/2003

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

Documents

Summary of Product Characteristics