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Deccox

Authorised

- Decoquinate

Product identification

Medicine name:
Deccox

Active substance:
Decoquinate

Target species:
Cattle
Sheep

Route of administration:
In-feed use

Product details

Active substance and strength:
Decoquinate
6.00 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

In-feed use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Sheep**
  - Meat and offal. 0 day
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51AX14

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
bag of 10 kg

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
15/12/2008

Manufacturing sites for batch release:
Phibro Medolla Manufacturing S.r.l.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1956 ESP

Date of authorisation status change:
4/02/2010

Reference member state:
Spain

Procedure number:
ES/V/0323/001

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/03/2026
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Package Leaflet

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Labelling

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