DIGESTIA SUSPENSION ORAL PARA TERNEROS LACTANTES, LECHONES LACTANTES, CORDEROS LACTANTES, CABRITOS LACTANTES Y CACHORROS LACTANTES

Authorised

- Calcium hydrogen phosphate (anhydrous)
- Magnesium sulfate heptahydrate
- Calcium chloride dihydrate
- Potassium chloride
- Sodium chloride
- Hydrochloric acid
- Dimeticone
- PEPSIN

Product identification

Medicine name:
DIGESTIA SUSPENSION ORAL PARA TERNEROS LACTANTES, LECHONES LACTANTES, CORDEROS LACTANTES, CABRITOS LACTANTES Y CACHORROS LACTANTES

Active substance:
Calcium hydrogen phosphate (anhydrous)

Magnesium sulfate heptahydrate

Calcium chloride dihydrate

Potassium chloride

Sodium chloride

Hydrochloric acid

Dimeticone

PEPSIN

Target species:
Cattle (calf)
Sheep (lamb)
Pig (piglet)
Dog (puppy)
Goat (kid)

Route of administration:
Oral use

Product details

Active substance and strength:
Calcium hydrogen phosphate (anhydrous)
2.50 milligram(s) / 1.00 millilitre(s)

Magnesium sulfate heptahydrate
2.50 milligram(s) / 1.00 millilitre(s)

Calcium chloride dihydrate
0.05 milligram(s) / 1.00 millilitre(s)

Potassium chloride
0.70 milligram(s) / 1.00 millilitre(s)

Sodium chloride
2.50 milligram(s) / 1.00 millilitre(s)

Hydrochloric acid
16.76 milligram(s) / 1.00 millilitre(s)

Dimeticone
26.40 milligram(s) / 1.00 millilitre(s)

PEPSIN

6.70 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Cattle (calf)**
  - Meat and offal. 0 day
- **Sheep (lamb)**
  - Meat and offal. 0 day
- **Pig (piglet)**
  - Meat and offal. 0 day
- **Dog (puppy)**
  - Meat and offal. 0 day
- **Goat (kid)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA09AC01

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:

Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish

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:
CZ Vaccines S.A.U.

Marketing authorisation date:
9/02/1971

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
2936 ESP

Date of authorisation status change:
16/12/2013

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

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