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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

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

Package Leaflet

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

Labelling

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