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LABIDROSOL B SOLUCION INYECTABLE

Authorised

- Pyridoxine hydrochloride
- Dexpanthenol
- Cyanocobalamin
- Riboflavin sodium phosphate hydrate
- Nicotinamide
- Thiamine hydrochloride

Product identification

Medicine name:
LABIDROSOL B SOLUCION INYECTABLE

Active substance:
Pyridoxine hydrochloride
Dexpanthenol
Cyanocobalamin
Riboflavin sodium phosphate hydrate
Nicotinamide
Thiamine hydrochloride

Target species:
Cattle
Sheep
Goat
Horse

Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Pyridoxine hydrochloride
5.00 milligram(s) / 1.00 millilitre(s)

Dexpanthenol
20.00 milligram(s) / 1.00 millilitre(s)

Cyanocobalamin
0.10 milligram(s) / 1.00 millilitre(s)

Riboflavin sodium phosphate hydrate
2.00 milligram(s) / 1.00 millilitre(s)

Nicotinamide
40.00 milligram(s) / 1.00 millilitre(s)

Thiamine hydrochloride
12.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

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

- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11EA

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
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:
Labiana Life Sciences S.A.

Marketing authorisation date:
30/11/1970

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3717 ESP

Date of authorisation status change:
11/01/2019

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

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