DILPHES 4

Authorised

- Pasteurella multocida, serogroup A, strain NCTC 12177, Inactivated
- Pasteurella multocida, serotype 6B, strain CECT 962, Inactivated
- Mannheimia haemolytica, serotype A1, strain ATCC 33365, Inactivated
- Mannheimia haemolytica, serotype A2, strain CECT 924, Inactivated

Product identification

Medicine name:
DILPHES 4

Active substance:
Pasteurella multocida, serogroup A, strain NCTC 12177, Inactivated
Pasteurella multocida, serotype 6B, strain CECT 962, Inactivated
Mannheimia haemolytica, serotype A1, strain ATCC 33365, Inactivated
Mannheimia haemolytica, serotype A2, strain CECT 924, Inactivated

Target species:
Cattle
Sheep
Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Pasteurella multocida, serogroup A, strain NCTC 12177, Inactivated
4.00 50% Protective Dose / 2.00 millilitre(s)

Pasteurella multocida, serotype 6B, strain CECT 962, Inactivated
4.00 50% Protective Dose / 2.00 millilitre(s)

Mannheimia haemolytica, serotype A1, strain ATCC 33365, Inactivated
41.08 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Mannheimia haemolytica, serotype A2, strain CECT 924, Inactivated
41.08 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Spain

Available in:
Spain

Package description:
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:
1/09/1987

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

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
3177 ESP

Date of authorisation status change:
6/02/2015

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.