DILPHES BG suspension for injection for cattle, sheep and goats

Authorised

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

Product identification

Medicine name:
DILPHES BG suspension for injection for cattle, sheep and goats

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

Target species:
Cattle
Sheep

Goat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Mannheimia haemolytica, serotype A2, strain CECT 924, Inactivated
41.08 enzyme-linked immunosorbent assay unit / 1.00 Dose

Mannheimia haemolytica, serotype A1, strain ATCC 33365, Inactivated
41.08 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pasteurella multocida, serotype 6B, strain CECT 962, Inactivated
100000.00 50% Protective Dose / 1.00 Dose

Pasteurella multocida, serogroup A, strain NCTC 12177, Inactivated
100000.00 50% Protective Dose / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB04
QI03AB
QI04AB02

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:

Available only in Bulgarian
Available only in Bulgarian

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:
Asklep-Pharma OOD

Marketing authorisation date:
14/05/2016

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

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2647

Date of authorisation status change:
19/10/2020

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

Documents

Package Leaflet

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Summary of Product Characteristics

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