MULTIVAC PSG инжекционна емулсия за свине

Authorised

- Haemophilus parasuis, strain IL-1-DEP, Inactivated
- Pasteurella multocida, strain 9, Inactivated
- Pasteurella multocida, strain 1231, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 371, Inactivated
- Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 370, Inactivated

Product identification

Medicine name:
MULTIVAC PSG инжекционна емулсия за свине

Active substance:
Haemophilus parasuis, strain IL-1-DEP, Inactivated

Pasteurella multocida, strain 9, Inactivated

Pasteurella multocida, strain 1231, Inactivated

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 371, Inactivated

Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 370, Inactivated

Target species:
Pig

Route of administration:
Injection

Product details

Active substance and strength:

Haemophilus parasuis, strain IL-1-DEP, Inactivated
2.00 billion organisms / 1.00 Dose

Pasteurella multocida, strain 9, Inactivated
2.00 billion organisms / 1.00 Dose

Pasteurella multocida, strain 1231, Inactivated
2.00 billion organisms / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 371, Inactivated
2.00 billion organisms / 1.00 Dose

Salmonella enterica, subsp. enterica, serovar Choleraesuis, strain 370, Inactivated
2.00 billion organisms / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in Bulgarian
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:
Mintech Co EOOD

Marketing authorisation date:
8/10/2019

Manufacturing sites for batch release:
Mintech Co EOOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2926

Date of authorisation status change:
17/01/2023

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 and Labelling

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