

Salmabic emulsija injekcijām vistām

Not
authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT B3, Inactivated
- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT C8, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain PT 2(4+), Inactivated

Product identification

Medicine name:

Salmabic emulsija injekcijām vistām

Active substance:

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT B3, Inactivated
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT C8, Inactivated
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain PT 2(4+),
Inactivated

Target species:

Chicken (layer hen)
Chicken (for reproduction)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT B3, Inactivated
100000000.00 Colony forming unit / 1.00 unit(s)

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT C8, Inactivated
100000000.00 Colony forming unit / 1.00 unit(s)

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain PT 2(4+),
Inactivated
100000000.00 Colony forming unit / 1.00 unit(s)

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Latvia

Package description:

Available only in [Latvian](#)

Available only in [Latvian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Abic Polska Sp. z o.o.

Marketing authorisation date:

7/04/2011

Manufacturing sites for batch release:

Abic Polska Sp. z o.o.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/NRP/11/0018

Date of authorisation status change:

23/02/2022

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.