

PORCILIS LAWSONIA ID LYOPHILISATE AND SOLVENT FOR EMULSION FOR INJECTION FOR PIGS

Authorised

- Lawsonia intracellularis, strain SPAH-08, Inactivated

Product identification

Medicine name:

PORCILIS LAWSONIA ID LYOPHILISATE AND SOLVENT FOR EMULSION FOR INJECTION FOR PIGS

Active substance:

Lawsonia intracellularis, strain SPAH-08, Inactivated

Target species:

Pig

Route of administration:

Intradermal use

Product details

Active substance and strength:

Lawsonia intracellularis, strain SPAH-08, Inactivated

5323.00 unit(s) / 1.00 unit(s)

Pharmaceutical form:

Lyophilisate and solvent for emulsion for injection

Withdrawal period by route of administration:

Intradermal use:

•

Pig

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB18

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

Cardboard box with 1 x 200 doses of vaccine and cardboard box with 1 x 40 ml solvent

Cardboard box with 10 x 200 doses of vaccine and cardboard box with 10 x 40 ml solvent

Cardboard box with 1 x 50 doses of vaccine and cardboard box with 1 x 10 ml solvent

Cardboard box with 10 x 50 doses of vaccine and cardboard box with 10 x 10 ml solvent

Cardboard box with 1 x 100 doses of vaccine and cardboard box with 1 x 20 ml solvent

Cardboard box with 10 x 100 doses of vaccine and cardboard box with 10 x 20 ml solvent

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:

Intervet International B.V.

Marketing authorisation date:

8/04/2021

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 817/21/04/2121

Date of authorisation status change:

8/04/2021

Reference member state:

France

Procedure number:

FR/V/0424/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

eu-puar-frv0424001-mr-rpe878-en.pdf