

ICTHIOVAC-STR ESTREPTOCOCOSIS RODABALLO Suspension for injection

Authorised

- Streptococcus parauberis, strain AZ-12.1, Inactivated
- Streptococcus parauberis, strain RA-99.1, Inactivated

Product identification

Medicine name:

ICTHIOVAC-STR ESTREPTOCOCOSIS RODABALLO Suspension for injection

Active substance:

Streptococcus parauberis, strain AZ-12.1, Inactivated

Streptococcus parauberis, strain RA-99.1, Inactivated

Target species:

Turbot

Route of administration:

Intraperitoneal use

Product details

Active substance and strength:

Streptococcus parauberis, strain AZ-12.1, Inactivated

75.00 Relative Percentage Survival / 0.10 millilitre(s)

Streptococcus parauberis, strain RA-99.1, Inactivated
75.00 Relative Percentage Survival / 0.10 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intraperitoneal use:

-

Turbot

- Meat. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10D

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

bottle of 500 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios Hipra S.A.

Marketing authorisation date:

4/08/2005

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

R752/05DGV

Date of authorisation status change:

13/10/2022

Reference member state:

Spain

Procedure number:

ES/V/0109/001

Concerned member states:

France Greece Portugal

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

Documents

Combined File of all Documents