ICTHIOVAC-VR

Authorised

- Vibrio anguillarum, serotype O2b, strain RV-22, Inactivated
- Vibrio anguillarum, serotype O2a, strain RG-111, Inactivated
- Vibrio anguillarum, serotype O1, strain R-82, Inactivated

Product identification

Medicine name:
ICTHIOVAC-VR
Icthiovac-VR concentrado para suspensão de imersão/para injeção em robalo e pregado

Active substance:
Vibrio anguillarum, serotype O2b, strain RV-22, Inactivated
Vibrio anguillarum, serotype O2a, strain RG-111, Inactivated
Vibrio anguillarum, serotype O1, strain R-82, Inactivated

Target species:
Seabass

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:

Vibrio anguillarum, serotype O2b, strain RV-22, Inactivated
60.00 Relative Percentage Survival / 1.00 millilitre(s)

Vibrio anguillarum, serotype O2a, strain RG-111, Inactivated
60.00 Relative Percentage Survival / 1.00 millilitre(s)

Vibrio anguillarum, serotype O1, strain R-82, Inactivated
60.00 Relative Percentage Survival / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Seabass**
  - Meat. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10D
QI10X

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

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Botella of 1000 ml

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:
Laboratorios Hipra S.A.

Marketing authorisation date:
23/07/2020

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
989/01/20RIVPT

Date of authorisation status change:
13/10/2022

Reference member state:
Spain

Procedure number:
ES/V/0385/001

Concerned member states:
Cyprus France Greece Italy Portugal

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

Documents

Summary of Product Characteristics

Combined File of all Documents