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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

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:
France

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:
20/07/2020

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7704200 8/2020

Date of authorisation status change:
20/07/2020

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

English (PDF)

Published on: 22/10/2025
Updated on: 13/03/2026
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Package Leaflet and Labelling

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

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