# 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:**
Cyprus

**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:**
14/09/2020

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

**Responsible authority:**
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

**Authorisation number:**
CY00798V

**Date of authorisation status change:**
14/09/2020

**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 |

Combined File of all Documents

English (PDF)
Published on: 24/07/2025
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