

# 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

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**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

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**Target species:**

Seabass

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**Route of administration:**

Intraperitoneal use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intraperitoneal use:**

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**Seabass**

- Meat. 0 degree day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI10D

QI10X

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Package description:**

Botella of 1000 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Laboratorios Hipra S.A.

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**Marketing authorisation date:**

8/09/2020

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**Manufacturing sites for batch release:**

Laboratorios Hipra, S.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

105515

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**Date of authorisation status change:**

8/09/2020

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0385/001

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**Concerned member states:**

Cyprus France Greece Italy Portugal

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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Summary of Product Characteristics