

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 SUSPENSIÓN PARA BAÑO/SUSPENSIÓN
INYECTABLE PARA LUBINA Y RODABALLO

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:

Spain

Package description:

Available only in Spanish

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:

18/07/2002

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

1467 ESP

Date of authorisation status change:

18/07/2002

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

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Combined File of all Documents

Labelling

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Package Leaflet

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