ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS

Authorised

- Photobacterium damselae, subsp. piscicida, strain DI21, Inactivated
- Vibrio anguillarum, serotype O1, Inactivated
- Vibrio anguillarum, serotype O2b, Inactivated
- Vibrio anguillarum, serotype O2a, Inactivated

Product identification

Medicine name:
ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS

Active substance:
Photobacterium damselae, subsp. piscicida, strain DI21, Inactivated

Vibrio anguillarum, serotype O1, Inactivated

Vibrio anguillarum, serotype O2b, Inactivated

Vibrio anguillarum, serotype O2a, Inactivated

Target species:
Seabass

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:

Photobacterium damselae, subsp. piscicida, strain DI21, Inactivated
60.00 Relative Percentage Survival / 1.00 Dose

Vibrio anguillarum, serotype O1, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Vibrio anguillarum, serotype O2b, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Vibrio anguillarum, serotype O2a, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Seabass**
  - Fish meat. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10X

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

---

Authorisation status:

Valid

Authorised in:

France

Package description:

500 ml bottle

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
9/05/2017

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

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

Authorisation number:
FR/V/3638521 0/2017

Date of authorisation status change:
16/05/2022

Reference member state:
France

Procedure number:
FR/V/0314/001

Concerned member states:
Croatia Cyprus Greece Italy Portugal Spain

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
Download

Package Leaflet and Labelling

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

Summary of Product Characteristics

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

eu-puar-frv0314001-mr-rpe485-en.pdf