# 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 INJECTABLE POUR BARS
ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS
ICTHIOVAC VR/PD, emulsione iniettabile per spigole

**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

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intraperitoneal use:**

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

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

QI10X

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**Legal status of supply:**

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

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Package description:**

500 ml bottle

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## 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: 19/02/2024
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Package Leaflet and Labelling

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Summary of Product Characteristics

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