

# ICTHIOVAC VR/PD, EMULSION FOR INJECTION FOR SEA BASS

Authorised

- Photobacterium damsela, 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 damsela, 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 damsela, 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

---

### **Authorisation status:**

Valid

---

### **Authorised in:**

Spain

---

### **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:**

2/06/2017

---

**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

---

**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

---

**Authorisation number:**

3548 ESP

---

**Date of authorisation status change:**

6/06/2017

---

**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](http://www.adrreports.eu/vet)

## Documents

### 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.

### Package Leaflet

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