

# 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

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**Active substance:**

Photobacterium damsela, subsp. piscicida, strain DI21, Inactivated

Vibrio anguillarum, serotype O1, Inactivated

Vibrio anguillarum, serotype O2b, Inactivated

Vibrio anguillarum, serotype O2a, 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:**

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

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### **Pharmaceutical form:**

Emulsion for injection

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### **Withdrawal period by route of administration:**

#### **Intraperitoneal use:**

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#### **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

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### **Authorisation status:**

Valid

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### **Authorised in:**

Italy

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### **Package description:**

500 ml bottle

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New 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:**

27/06/2017

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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:**

105037

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**Date of authorisation status change:**

27/06/2017

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**Reference member state:**

France

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**Procedure number:**

FR/V/0314/001

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**Concerned member states:**

Croatia Cyprus Greece Italy Portugal Spain

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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)

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