

# 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

ICTHIOVAC VR/PD, ενέσιμο γαλάκτωμα για λαβράκι

### 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

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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

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### Authorisation status:

Valid

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### Authorised in:

Cyprus

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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:**

23/01/2023

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**Manufacturing sites for batch release:**

Laboratorios Hipra S.A.

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**Responsible authority:**

Ministry Of Agriculture Rural Development And Environment

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**Authorisation number:**

CY00935V

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**Date of authorisation status change:**

23/01/2023

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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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