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
ICTHIOVAC VR/PD, emulsione iniettabile per spigole
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

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

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Authorisation status:

Valid

Authorised in:

Italy

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:
27/06/2017

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

Responsible authority:
MdS

Authorisation number:
105037

Date of authorisation status change:
27/06/2017

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

Documents

Combined File of all Documents

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