ICTHIOVAC VNN, EMULSION FOR INJECTION FOR SEABASS

Authorised

- Redspotted grouper nervous necrosis virus, strain 1103, Inactivated

Product identification

Medicine name:
ICTHIOVAC VNN, EMULSION FOR INJECTION FOR SEABASS
ICTHIOVAC VNN ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ

Active substance:
Redspotted grouper nervous necrosis virus, strain 1103, Inactivated

Target species:
Seabass

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:
Redspotted grouper nervous necrosis virus, strain 1103, Inactivated
1.30 relative potency / 0.10 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intraperitoneal use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10X

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
500 ml (5000 doses) high density polyethylene bottles closed with nitrile-chlorobutyl rubber stoppers and aluminium caps.

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:
28/12/2022

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
144684/29-12-2022/K-0235402

Date of authorisation status change:
28/12/2022

Reference member state:
France

Procedure number:
FR/V/0349/002

Concerned member states:
Croatia Cyprus Greece Italy Portugal Spain

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

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