ICTHIOVAC VNN, EMULSION FOR INJECTION FOR SEA BASS

Authorised

- Redspotted grouper nervous necrosis virus, strain 1103, Inactivated

Product identification

Medicine name:
ICTHIOVAC VNN, EMULSION FOR INJECTION FOR SEA BASS
ICTHIOVAC VNN, ενέσιμο γαλάκτωμα για λαβράκι

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 / 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:
Cyprus

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:
18/11/2022

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

Responsible authority:

Ministry Of Agriculture Rural Development And Environment

Authorisation number:
CY00902V

Date of authorisation status change:
18/11/2022

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

Documents

Combined File of all Documents

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