ICTHIOVAC VNN

Authorised

- Redspotted grouper nervous necrosis virus, strain 1103, Inactivated

Product identification

Medicine name:
ICTHIOVAC VNN
ICTHIOVAC VNN, emulzija za injekciju, za lubine

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
1.00 Relative Percentage Survival / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Seabass**
  - All relevant tissues. 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:
Croatia

Package description:
Vial of 5000 doses (5000 doses)

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:
29/03/2019

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

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/19-01/221

Date of authorisation status change:
12/03/2025

Reference member state:
France

Procedure number:
FR/V/0349/001

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.
