ICTHIOVAC VNN

Not authorised

- Redspotted grouper nervous necrosis virus, strain 1103, Inactivated

Product identification

Medicine name:
ICTHIOVAC VNN

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:
Surrendered

Authorised in:
Spain

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:
17/04/2019

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

Responsible authority:

Spanish Agency For Medicines And Health Products

Authorisation number:
3775 ESP (Non Valid)

Date of authorisation status change:
17/04/2019

Reference member state:
France

Procedure number:
FR/V/0349/001

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.

Package Leaflet

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

Labelling

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

Combined File of all Documents