

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. no withdrawal period
Withdrawal period is 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10X

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Cyprus

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:

14/04/2019

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Authorisation number:

CY00736V

Date of authorisation status change:

14/04/2019

Reference member state:

France

Procedure number:

FR/V/0349/001/DC

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.

Combined File of all Documents