Norvax Compact PD Emulsion for Injection for Atlantic Salmon

Not authorised

- Salmon pancreas disease virus, strain F93-125, Inactivated

Product identification

Medicine name:
Norvax Compact PD Emulsion for Injection for Atlantic Salmon

Active substance:
Salmon pancreas disease virus, strain F93-125, Inactivated

Target species:
Atlantic salmon

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:
Salmon pancreas disease virus, strain F93-125, Inactivated
50118700.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Atlantic salmon**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10AA01

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

Authorisation status:
Surrendered

Authorised in:
United Kingdom (Northern Ireland)

Package description:
Bottles of polyethylene terephthalate (PET) closed with a rubber stopper and aluminium cap. Pack size: 250 ml (2,500 doses).
Bottles of polyethylene terephthalate (PET) closed with a rubber stopper and aluminium cap. Pack size: 500 ml (5,000 doses).

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
MSD Animal Health UK Limited

Marketing authorisation date:
10/08/2011

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 01708/3037

Date of authorisation status change:
20/08/2024

Reference member state:
Ireland

Procedure number:
IE/V/0257/001

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

Documents

Summary of Product Characteristics