

AquaVac PD3 Emulsion for Injection, for Atlantic Salmon

Authorised

- Salmon pancreas disease virus, strain F93-125, Inactivated
- Infectious pancreatic necrosis virus, Inactivated
- *Aeromonas salmonicida*, subsp. *salmonicida*, Inactivated

Product identification

Medicine name:

AquaVac PD3 Emulsion for Injection, for Atlantic Salmon

Active substance:

Salmon pancreas disease virus, strain F93-125, Inactivated

Infectious pancreatic necrosis virus, Inactivated

Aeromonas salmonicida, subsp. *salmonicida*, Inactivated

Target species:

Atlantic salmon

Route of administration:

Intraperitoneal use

Product details

Active substance and strength:

Salmon pancreas disease virus, strain F93-125, Inactivated

Infectious pancreatic necrosis virus, Inactivated
1.50 enzyme-linked immunosorbent assay unit / 1.00 Dose
Aeromonas salmonicida, subsp. salmonicida, Inactivated
80.00 Relative Percentage Survival / 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:

QI10AL

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

Bottles of polyethylene terephthalate (PET) closed with a rubber stopper and aluminium cap. Package 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:

Intervet International B.V.

Marketing authorisation date:

26/03/2015

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 06376/3056

Date of authorisation status change:

14/11/2023

Reference member state:

Ireland

Procedure number:

IE/V/0339/001

Concerned member states:

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents