

PARTICULARS TO APPEAR ON THE OUTER PACKAGE AND THE IMMEDIATE PACKAGE

PLASTIC BAG

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

ALPHA JECT micro 6 emulsion for injection for Atlantic salmon

2. STATEMENT OF ACTIVE SUBSTANCES

1 dose (0.05 ml):

Formaldehyde inactivated cultures of:

<i>Aeromonas salmonicida</i> subsp. <i>Salmonicida</i>	$\geq 12.6 \log_2$ ELISA units
<i>Listonella anguillarum</i> O1	RPS $\geq 75\%$
<i>Listonella anguillarum</i> O2a	RPS $\geq 75\%$
<i>Vibrio salmonicida</i>	RPS $\geq 90\%$
<i>Moritella viscosa</i>	$\geq 10.7 \log_2$ ELISA units
IPN virus Sp.	0.12-0.28 AU

Paraffin, light liquid: 23 mg.

3. PHARMACEUTICAL FORM

Emulsion for injection.

4. PACKAGE SIZE

250 ml
500 ml

5. TARGET SPECIES

Atlantic salmon

6. INDICATION(S)

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Intraperitoneal use.
Read the package leaflet before use.

8. WITHDRAWAL PERIOD(S)

Withdrawal period: 0 degree days.

9. SPECIAL WARNING(S), IF NECESSARY

Read the package leaflet before use.
Accidental injection is dangerous.

10. EXPIRY DATE

EXP:
Once broached use within: 10 hours.

11. SPECIAL STORAGE CONDITIONS

Store and transport refrigerated.
Do not freeze.
Protect from light.

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: read package leaflet.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, IF APPLICABLE

For animal treatment only.
To be supplied only on veterinary prescription.

14. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

Keep out of the sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

PHARMAQ AS
Industrivegen 50
Skogmo Industriområde
7863 Overhalla
Norway

16. MARKETING AUTHORISATION NUMBER(S)

Vm 21714/3001

17. MANUFACTURER'S BATCH NUMBER

Batch:

PACKAGE LEAFLET

ALPHA JECT micro 6 emulsion for injection for Atlantic salmon

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Marketing authorisation holder and manufacturer responsible for batch release:

PHARMAQ AS
Industrivegen 50
Skogmo Industriområde
7863 Overhalla
Norway

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

ALPHA JECT micro 6 emulsion for injection for Atlantic salmon.

3. STATEMENT OF THE ACTIVE SUBSTANCES AND OTHER INGREDIENTS

1 dose (0.05 ml) contains:

Active substance:

Formaldehyde inactivated cultures of:

<i>Aeromonas salmonicida</i> subsp. <i>salmonicida</i>	≥ 12.6 log ₂ ELISA units
<i>Listonella anguillarum</i> * serotype O1	RPS ≥ 75 %
<i>Listonella anguillarum</i> * serotype O2a	RPS ≥ 75 %
<i>Vibrio salmonicida</i>	RPS ≥ 90 %
<i>Moritella viscosa</i>	≥ 10.7 log ₂ ELISA units
Infectious Pancreatic Necrosis Virus serotype Sp	0.12-0.28 AU

ELISA units: Serological response in Atlantic salmon,

RPS: Relative Percentage Survival in challenge studies on Atlantic salmon.

AU: Antigenicity Units (quantity of virus antigen measured in the final product).

* *Listonella anguillarum* is synonymous with *Vibrio anguillarum*.

Adjuvant:

Paraffin, light liquid (mineral oil): 23 mg.

Emulsion for injection. White to cream coloured.

4. INDICATIONS

For active immunisation of Atlantic salmon to reduce mortality caused by infections with *Aeromonas salmonicida* (furunculosis), *Vibrio salmonicida* (coldwater vibriosis),

Listonella anguillarum serotype O1 and O2a (vibriosis), *Moritella viscosa* (winter sore) and IPNV (infectious pancreatic necrosis).

Onset of immunity: 520 degree days post vaccination for the bacterial antigens and 600 degree days post vaccination for IPNV.

Duration of immunity: 1 year for the bacterial antigens and 5.5 months for IPNV.

5. CONTRAINDICATIONS

None.

6. ADVERSE REACTIONS

The frequency of adverse reactions observed after vaccination in laboratory studies and field trials:

Very common: Melanisation in the abdominal cavity
Mild visceral adhesions (Speilberg score 1-2)
Common: Moderate visceral adhesions (Speilberg score 3)
Very rare: Serious visceral adhesions (Speilberg score ≥ 4)

The severity of adverse reactions may be influenced by different factors such as sanitation, vaccination technique, fish size at vaccination and water temperature during vaccination and in the first 6-12 weeks after vaccination. As a general precaution it is recommended to perform vaccination at water temperature of 15 °C or below.

Small fish and high water temperature may increase the severity of adverse reactions.

The frequency of adverse reactions is defined using the following convention:

- very common (more than 1 in 10 animals treated displaying adverse reaction(s))
- common (more than 1 but less than 10 animals in 100 animals treated)
- uncommon (more than 1 but less than 10 animals in 1,000 animals treated)
- rare (more than 1 but less than 10 animals in 10,000 animals treated)
- very rare (less than 1 animal in 10,000 animals treated, including isolated reports).

If you notice any side effects, even those not already listed in this package leaflet or you think that the medicine has not worked, please inform your veterinary surgeon/fish health biologist.

7. TARGET SPECIES

Atlantic salmon (*Salmo salar*) of a minimum weight of 25 g.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

Dosage

Administer a single dose of 0.05 ml per fish. Fish should not be vaccinated more than once.

Administration route

The vaccine should be administered by intraperitoneal (i.p) injection into the midline about one fin length anterior to the base of the pelvic fin.

It is recommended to starve the fish for a minimum of 48 hours before vaccination. The fish should be anaesthetised prior to injection.

9. ADVICE ON CORRECT ADMINISTRATION

To reduce the risk of adverse reactions, it is important to deposit the entire dose in the abdominal cavity. The injection needle used should have appropriate length to penetrate the abdominal wall and 1-2 mm into the abdominal cavity.

Let the vaccine slowly reach 15-20°C by keeping it at room temperature. Ensure a homogenous emulsion prior to use by squeezing and shaking the vaccine bag for approx. 2 minutes. Only administer the vaccine if it appears as a homogenous, white to cream coloured emulsion after shaking. The vaccine should not be used if the vaccine shows signs of a brownish water phase in the bottom of the container. Contact the distributor for further advice.

The injection devices used for vaccination, i.e. automatic vaccination machines or manual syringes, must be designed and suitable for administration of the recommended dose volume in the target species. The devices must be operated by trained personnel and should be calibrated according to the manufacturers' recommendation prior to use. Special care should be taken to ensure air is removed from the injection equipment (chambers and tubes) prior to vaccination. Regular dose controls are recommended.

The vaccination equipment should be thoroughly cleaned / sterilized before use.

10. WITHDRAWAL PERIOD

Zero degree days.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.

Store and transport refrigerated (2°C-8°C).

Do not freeze.

Protect from light.

Do not use this veterinary medicinal product after the expiry date which is stated on the label after EXP.

Shelf life after first opening the container: 10 hours.

12. SPECIAL WARNINGS

Special precautions for use in animals:

Vaccinate healthy fish only.

Vaccination should preferably be performed at water temperatures of 15°C or below.

Do not vaccinate at water temperatures below 3°C or above 18°C.

Avoid vaccination during smoltification.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

People with known hypersensitivity to fish vaccines should avoid contact with the veterinary medicinal product.

Protective equipment consisting of guarded needles should be used during manual vaccination.

Ensure that the method of fixation and handling of the fish minimises the risk of accidental self-injection. Repeated self-injections may aggravate the adverse effects or increase the risk of anaphylactic shock.

To the user:

This veterinary medicinal product contains mineral oil. Accidental self-injection may result in severe pain and swelling, particularly if injected into a joint or finger, and in rare cases could result in the loss of the affected finger if prompt medical attention is not given. If you are accidentally injected with this veterinary medicinal product, seek prompt medical advice even if only a very small amount is injected and take the package leaflet with you. If pain persists for more than 12 hours after medical examination, seek medical advice again.

To the physician:

This veterinary medicinal product contains mineral oil. Even if small amounts have been injected, accidental injection with this product can cause intense swelling, which may, for example, result in ischaemic necrosis and even the loss of a digit. Expert, PROMPT, surgical attention is required and may necessitate early incision and irrigation of the injected area, especially where there is involvement of finger pulp or tendon.

Fertility:

The potential effect of vaccination on spawning function has not been investigated. Vaccination of broodfish should only be done according to a benefit-risk assessment by the responsible veterinarian/fish health biologist.

Interaction with other medicinal products and other forms of interaction:

No information is available on the safety and efficacy of this vaccine when used with any other veterinary medicinal product. A decision to use this vaccine before or after any other veterinary medicinal product therefore needs to be made on a case by case basis.

Overdose (symptoms, emergency procedures, antidotes):
Following administration of 0.1 ml of the vaccine (double dose) no other adverse reactions than those described in section "Adverse reactions" were seen.

Incompatibilities:

Do not mix with any other veterinary medicinal product.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

Ask your veterinary surgeon or fish health biologist how to dispose of medicines no longer required. These measures should help to protect the environment.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

March 2023

15. OTHER INFORMATION

250 ml or 500 ml multilayer plastic foil bag. The giving port is closed with a sealed rubber stopper.

Pack sizes:

250 ml and 500 ml

Not all pack sizes may be marketed.

Approved 22 June 2023

