

# Clynav (--)- Solution for injection

Authorised

- Salmon pancreas disease virus, DNA plasmid pUK-SPDV-poly2#1 encoding structural polyprotein

## Product identification

**Medicine name:**

Clynav (--)- Solution for injection

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**Active substance:**

Salmon pancreas disease virus, DNA plasmid pUK-SPDV-poly2#1 encoding structural polyprotein

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**Target species:**

Atlantic salmon

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Salmon pancreas disease virus, DNA plasmid pUK-SPDV-poly2#1 encoding structural polyprotein

6.00 microgram(s) / 1.00 Bag

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Atlantic salmon**

- Not applicable. 0 day Zero degree days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI10AX

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Available in:**

Ireland , Netherlands , Norway

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**Package description:**

Packaging:Bag (ethyl vinyl acetate), Package\_size:1 bag, Content:250 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

INTERVET INTERNATIONAL B.V.

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**Marketing authorisation date:**

27/06/2017

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**Manufacturing sites for batch release:**

Merck Sharp & Dohme Animal Health S.L.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

27/06/2017

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 11/01/2024

Updated on: 8/06/2026

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ema-puar-clynav-v-2390-var-ii-0010-en.pdf

ema-puar-clynavv-2390-par-en.pdf