

Ichthiovac ERM Concentrate for dip suspension

Authorised

- Yersinia ruckeri, serotype O1, biotype 1, strain 8363, Inactivated
- Yersinia ruckeri, serotype O1, biotype 2, strain 8302, Inactivated
- Yersinia ruckeri, serotype O2, biotype 1, strain 8365, Inactivated

Product identification

Medicine name:

Ichthiovac ERM Concentrate for dip suspension

Active substance:

Yersinia ruckeri, serotype O1, biotype 1, strain 8363, Inactivated

Yersinia ruckeri, serotype O1, biotype 2, strain 8302, Inactivated

Yersinia ruckeri, serotype O2, biotype 1, strain 8365, Inactivated

Target species:

Atlantic salmon

Route of administration:

Dipping

Product details

Active substance and strength:

Yersinia ruckeri, serotype O1, biotype 1, strain 8363, Inactivated

Presentation_strength: ≥ 10.19 log₁₀ BDC/ml Reference: In House Index: 0

Yersinia ruckeri, serotype O1, biotype 2, strain 8302, Inactivated

Presentation_strength: ≥ 9.91 log₁₀ BDC/ml Reference: In House Index: 1

Yersinia ruckeri, serotype O2, biotype 1, strain 8365, Inactivated

Presentation_strength: ≥ 10.07 log₁₀ BDC/ml Reference: In House Index: 2

Pharmaceutical form:

Concentrate for dip suspension

Withdrawal period by route of administration:

Dipping:

-

Atlantic salmon

- Fish meat. 0 day
Zero degree days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

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)

Package description:

Packaging: Bottle (PP), Package_size: 1 bottle, Content: 1000 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Laboratorios Hipra, S.A.

Marketing authorisation date:

23/01/2025

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

23/01/2025

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

Documents

Combined File of all Documents

English (PDF)

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