File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600001874710

RESPIVAC aMPV (--) - Lyophilisate for oculonasal suspension/use in drinking water

Authorised

- Turkey rhinotracheitis virus, strain 1062, Live

Product identification

Medicine name:
RESPIVAC aMPV (--) - Lyophilisate for oculonasal suspension/use in drinking water

Active substance:
Turkey rhinotracheitis virus, strain 1062, Live

Target species:
Chicken
Turkey

Route of administration:
Oculonasal use
In drinking water use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain 1062, Live

Presentation_strength:$10^{2.4}$ – $10^{5.4}$ $CCID_{50}$/dose Reference:Ph. Eur. Index:0

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Oculonasal use:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Turkey**
  - Not applicable. 0 day Zero days

In drinking water use:

- **Chicken**
  - Not applicable. 0 day Zero days
- **Turkey**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD01

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:Vial (glass), Package_size:1 vial, Content:1000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:1000 doses
Packaging:Vial (glass), Package_size:1 vial, Content:2000 doses
Packaging:Vial (glass), Package_size:1 vial, Content:5000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:5000 doses
Packaging:Vial (glass), Package_size:1 vial, Content:10000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:2000 doses
Packaging:Vial (glass), Package_size:10 vials, Content:10000 doses

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
30/05/2024

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:
30/05/2024

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

Documents

Combined File of all Documents

English (PDF)
Published on: 9/10/2026
Download

ema-puar-v6160-respivacampv-vra0000309778-en.pdf

ema-puar-v6160-respivacampv-former-respivactrt-en.pdf