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Poulvac IBMM + ARK lyophilisate for oculonasal suspension for chickens

Authorised

- Infectious bronchitis virus, type 793/B, strain Arkansas 3168, Live
- Infectious bronchitis virus, type Massachusetts, strain 1263, Live

Product identification

Medicine name:
Poulvac IBMM + ARK lyophilisate for oculonasal suspension for chickens

Active substance:
Infectious bronchitis virus, type 793/B, strain Arkansas 3168, Live

Infectious bronchitis virus, type Massachusetts, strain 1263, Live

Target species:
Chicken

Route of administration:
Oculonasal use

Product details

Active substance and strength:

Infectious bronchitis virus, type 793/B, strain Arkansas 3168, Live
3.30 log 10 50% embryo infective dose / 1.00 Dose

Infectious bronchitis virus, type Massachusetts, strain 1263, Live
3.30 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension

Withdrawal period by route of administration:
Oculonasal use:

- **Chicken**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD07

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Hydrolytic type I glass vials closed with butyl rubber stopper and sealed with an aluminium crimp cap. containing 10 x 10,000 doses (box of 10 vials).
Hydrolytic type I glass vials closed with butyl rubber stopper and sealed with an aluminium crimp cap. containing 10 x 5,000 doses (box of 10 vials).
Hydrolytic type I glass vials with a butyl rubber (Ph Eur) stopper and aluminium cap of 6 ml containing 1,000 doses (box of 10 vials).

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
2/02/2004

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V259375

Date of authorisation status change:
2/02/2004

Reference member state:
Netherlands

Procedure number:
NL/V/0258/001

Concerned member states:
Belgium Greece Ireland Italy United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents