

HATCHPAK IB H120 FROZEN SUSPENSION FOR OCULONASAL USE

Authorised

- Avian infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:

HATCHPAK IB H120 FROZEN SUSPENSION FOR OCULONASAL USE
Hatchpak IB H120, diepgevroren suspensie voor vernevelsuspensie

Active substance:

Avian infectious bronchitis virus, type Massachusetts, strain H120, Live

Target species:

Chicken (one day-old chick)

Route of administration:

Oculonasal use

Product details

Active substance and strength:

Avian infectious bronchitis virus, type Massachusetts, strain H120, Live
3.70 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:

Concentrate for nebuliser solution

Withdrawal period by route of administration:

Oculonasal use:

-

Chicken (one day-old chick)

- 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:

Netherlands

Package description:

Container of liquid nitrogen of canisters of 4 yellow ampoules of 10,000 doses

Container of liquid nitrogen of canisters of 4 yellow ampoules of 15,000 doses

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:

Boehringer Ingelheim Animal Health Netherlands B.V.

Marketing authorisation date:

24/10/2013

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10459

Date of authorisation status change:

24/08/2022

Reference member state:

France

Procedure number:

FR/V/0171/001

Concerned member states:

Belgium Bulgaria Cyprus Czechia Germany Greece Hungary Italy Latvia
Lithuania Netherlands Poland Romania Slovakia Slovenia
United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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