File downloaded on 2026-11-01
Source URL: https://medicines.health.europa.eu/veterinary/en/600000045832

HATCHPAK IB H120 NEO EFFERVESCENT TABLET FOR OCULONASAL SUSPENSION FOR CHICKENS

Authorised

- Infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:
HATCHPAK IB H120 NEO EFFERVESCENT TABLET FOR OCULONASAL SUSPENSION FOR CHICKENS

Active substance:
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:

Infectious bronchitis virus, type Massachusetts, strain H120, Live
3.70 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Effervescent tablet

Withdrawal period by route of administration:
Oculonasal use:

- **Chicken (one day-old chick)**
  - Meat and offal. 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

Available in:
Netherlands

Package description:
Box of 1 blister of 10 tablets of 1,000 doses
Box of 10 blisters of 10 tablets of 2,000 doses
Box of 10 blisters of 10 tablets of 1,000 doses
Box of 1 blister of 10 tablets of 2,000 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:
25/11/2016

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 119555

Date of authorisation status change:
20/06/2022

Reference member state:
France

Procedure number:
FR/V/0171/002

Concerned member states:
Belgium Cyprus Denmark Finland Germany Greece Iceland Ireland Italy Netherlands Slovenia Spain Sweden 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

English (PDF)
Published on: 1/04/2026
Download

eu-puar-frv0171002-mr-rpe682-en.pdf