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TAbic IB VAR206, effervescent tablets for suspension for chickens

Authorised

- Infectious bronchitis virus, strain 2-06, Live

Product identification

Medicine name:
TAbic IB VAR206, effervescent tablets for suspension for chickens

Active substance:
Infectious bronchitis virus, strain 2-06, Live

Target species:
Chicken (hen)

Route of administration:
Nebulisation use

Product details

Active substance and strength:
Infectious bronchitis virus, strain 2-06, Live

Pharmaceutical form:
Effervescent tablet

Withdrawal period by route of administration:

Nebulisation use:

- **Chicken (hen)**
  - Meat and offal. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Tablets (500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.

Tablets (1000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.

Tablets (2500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.

Tablets (1000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.

Tablets (10 000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.

Tablets (5000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
Tablets (2000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.
Tablets (10 000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
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Tablets (5000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.
Tablets (1500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
Tablets (3000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
Tablets (3500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
Tablets (4000 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.
Tablets (4500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 1 blister.

Tablets (1500 doses) are packed in Aluminium – Aluminium blisters: Alum Soft Silver (PVC/PVDC) – Alum Silver (PVC). Each blister contains 10 tablets. Cardboard box contains 2 blisters.
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Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Phibro Animal Health (Poland) Sp. z o.o.

Marketing authorisation date:
18/05/2016

Manufacturing sites for batch release:
Synoptis Industrial Sp. z o.o.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2538

Date of authorisation status change:
28/01/2022

Reference member state:
Poland

Procedure number:
PL/V/0109/001

Concerned member states:
Bulgaria Czechia Latvia Lithuania Romania Slovakia Slovenia

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.

Labelling

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

Combined File of all Documents

Package Leaflet

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