

TAbic IB VAR206, effervescent tablets for suspension for chickens

Authorised

- Avian infectious bronchitis virus, strain 2-06, Live

Product identification

Medicine name:

TAbic IB VAR206, effervescent tablets for suspension for chickens

TABIC IB VAR206, comprimate efervescente pentru suspensie pentru găini

Active substance:

Avian infectious bronchitis virus, strain 2-06, Live

Target species:

Chicken (hen)

Route of administration:

Nebulisation use

Product details

Active substance and strength:

Avian 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

Authorisation status:

Valid

Authorised in:

Romania

Available in:

Romania

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:

28/11/2019

Manufacturing sites for batch release:

Synoptis Industrial Sp. z o.o.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

220048

Date of authorisation status change:

14/01/2025

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

Combined File of all Documents

English (PDF)

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