

AMOXIPRO 10 % POUDRE POUR ADMINISTRATION DANS L'EAU DE BOISSON/ LE LAIT VEAU PORC POULET CANARD DINDE

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

AMOXIPRO 10 % POUDRE POUR ADMINISTRATION DANS L'EAU DE BOISSON/ LE LAIT VEAU PORC POULET CANARD DINDE

Active substance:

Amoxicillin trihydrate

Target species:

Turkey

Chicken (pullet for egg production, future layer)

Pig

Cattle (calf)

Duck

Chicken (for reproduction)

Chicken (broiler)

Chicken (layer hen)

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate

114.80 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water/milk

Withdrawal period by route of administration:

Oral use:

-

Turkey

- Meat and offal. 2 day

-

Chicken (pullet for egg production, future layer)

- Meat and offal. 2 day

-

Pig

- Meat and offal. 2 day

-

Cattle (calf)

- Meat and offal. 2 day

-

Duck

- Meat and offal. 2 day

-

Chicken (for reproduction)

- Meat and offal. 2 day

-

Chicken (broiler)

- Meat and offal. 2 day

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Chicken (layer hen)

- Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les espèces pondeuses et productrices d'œufs de consommation, 4 semaines avant le démarrage de la ponte et pendant celle-ci.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

16/09/1997

Manufacturing sites for batch release:

Ceva Sante Animale

Ceva Sante Animale

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/7044514 5/1997

Date of authorisation status change:

16/09/2012

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.