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DOXIPULVIS 500 MG/G POWDER FOR USE IN DRINKING WATER / MILK

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
DOXIPULVIS 500 MG/G POWDER FOR USE IN DRINKING WATER / MILK

Active substance:
Doxycycline hyclate

Target species:
Turkey
Pig
Cattle (calf)
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Doxycycline hyclate

577.10 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. 12 day
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 4 weeks before the start of the laying period.

- **Pig**
  - Meat and offal. 6 day

- **Cattle (calf)**
  - Meat and offal. 14 day

- **Chicken**
  - Meat and offal. 7 day
  - Eggs. no withdrawal period

Not for use in birds producing eggs for human consumption. Do not use within 4 weeks before the start of the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
S P Veterinaria S.A.

Marketing authorisation date:
18/08/2016

Manufacturing sites for batch release:
S P Veterinaria S.A.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402241.00.00

Date of authorisation status change:
13/01/2022

Reference member state:
France

Procedure number:

FR/V/0298/001

Concerned member states:
Bulgaria Cyprus Germany Greece Hungary Ireland Italy Malta Poland Portugal Romania Spain 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)
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