GENTA VIT

Authorised

- Biotin
- Folic acid
- Ascorbic acid
- VITAMIN B12
- VITAMIN B6
- D-PANTOTHENIC ACID
- Riboflavin
- Thiamine
- Nicotinic acid
- Menadione
- TOCOPHERYL ACETATE
- Colecalciferol
- Retinol
- Gentamicin sulfate

Product identification

Medicine name:
GENTA VIT

Active substance:
Biotin

Folic acid

Ascorbic acid

VITAMIN B12

VITAMIN B6

D-PANTOTHENIC ACID

Riboflavin

Thiamine

Nicotinic acid

Menadione

TOCOPHERYL ACETATE

Colecalciferol

Retinol

Gentamicin sulfate

Target species:
Pig

Route of administration:
In drinking water use

Product details

Active substance and strength:
Biotin
0.50 milligram(s) / 100.00 gram(s)

Folic acid
1.50 milligram(s) / 100.00 gram(s)

Ascorbic acid
375.00 milligram(s) / 100.00 gram(s)

VITAMIN B12
0.04 milligram(s) / 100.00 gram(s)

VITAMIN B6
15.00 milligram(s) / 100.00 gram(s)

D-PANTOTHENIC ACID
38.00 milligram(s) / 100.00 gram(s)

Riboflavin
37.00 milligram(s) / 100.00 gram(s)

Thiamine

10.00 milligram(s) / 100.00 gram(s)

Nicotinic acid
190.00 milligram(s) / 100.00 gram(s)

Menadione
15.00 milligram(s) / 100.00 gram(s)

TOCOPHERYL ACETATE
10.00 gram(s) / 100.00 gram(s)

Colecalciferol
7500.00 international unit(s) / 100.00 gram(s)

Retinol
75000.00 international unit(s) / 100.00 gram(s)

Gentamicin sulfate
1.60 gram(s) / 100.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:
In drinking water use:

- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA91

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

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Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

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:
Farma Vet OOD

Marketing authorisation date:
7/06/2011

Manufacturing sites for batch release:
Farma Vet OOD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1587

Date of authorisation status change:
2/06/2016

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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