LEAFLET FOR:
NICOPRON - 80 powder

NICOPRON - 80 pulvis

Powder for cattle and sheep

1. NAME AND ADDRESS OF THE AUTHORIZATION HOLDER AND OF THE PRODUCER (IF THEY DIFFERENT)

VETPROM AD, 2400, Radomir, Bulgaria, 26, Otets Paisiy Str.

Tel.: 0777/8-24-93, 8-02-68; Fax. 0777/8-23-91

E-mail: vetprom@abv.bg

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

NICOPRON – 80 powder

NICOPRON - 80 pulvis

3. ACTIVE SUBSTANCE AND EXCIPIENTS

Active Substance:

Sodium propionate 79.691 g.

Excipients:

Ferrous sulphate

Copper sulphate

Manganese sulphate

Cobalt sulphate

4. THERAPEUTIC INDICATIONS

In cattle and sheep, with acetonemia (ketosis), primary and secondary indigestion, inadequate nutrition, overfeeding with concentrated fodder, timpania, obstipation, surgical interventions, antibiotic and sulfonamide administration, etc. To better the absorption of the fodder and increasing growth of animals.

5. CONTRAINDICATIONS

Not to be used in animals with hypersensitivity towards the active substance and the excipients.

6. ADVERSE REACTIONS

Not known.

7. ANIMAL SPECIES, FOR WHICH THIS VMP IS DESIGNED

Cattle and sheep.

8. RECOMMENDED DOSES FOR EACH OF THE ANIMAL SPECIES, ADMINISTRATION ROUTE

Cows – 1 to 2 packs in 1 to 5 liters of water;

Sheep – 1 pack in 1 liter of water.

1 to 3 times daily for 2 to 3 days.

The prepared solution is no to be used a second time.

Administration:

Orally, the solution of the preparation is administered with the help of a bottle.

9. ADVICE FOR PROPER ADMINISTRATION

Always use **NICOPRON – 80 powder** as prescribed by your veterinarian. If in doubt, ask your veterinarian.

10. WITHDRAWAL PERIOD

Milk – zero hours.

Meat and internal organs – zero days.

11. SPECIAL PRECAUTIONS FOR STORAGE OF THE VMP

Keep the medication out of sight and reach of children!

Do not store above 25ºC.

Protect from light.

Keep in dry place.

This veterinary medicinal product should not be used after the expiry date on the indicated label.

12. SPECIAL PRECAUTIONS FOR USE

Interactions with other veterinary medicinal products and other forms of interactions:

In a powder form, the constituents are in the form of acids and they cannot interact. No interactions are known with other medicinal products.

Overdose (symptoms, emergency measures, antidotes) if necessary:

When higher doses are applied, an alkalosis may develop. The alkalosis subsides after the cessation of the administration.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED VETERINARY MEDICINAL PRODUCT OR WASTE MATERIALS DERIVED FROM THE USE OF SUCH PRODUCTS

The VMP should not be disposed of together with household wastes or sewage. Ask your veterinarian what to do with the no longer necessary VMPs. These measures would help environmental protection.

14. DATE OF THE LAST UPDATE OF THE TEXT

07/11/2014

15. ADDITIONAL INFORMATION

Please, contact the local representative of the authorization holder for any information concerning this veterinary medicinal product:

VETPROM AD, 2400, Radomir, Bulgaria, 26, Otets Paisiy Str.
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