

PATIENT PACKAGE INSERT FOR A VETERINARY PREPARATION

The medicine is dispensed with a veterinarian's prescription only
For use in animals only

1. NAME OF THE VETERINARY MEDICINE, ITS FORM AND STRENGTH

S.C.P. 47 Veterinary

2. ACTIVE INGREDIENT

Each 1 gram contains:

Sulfachloropyrazine sodium monohydrate 470 mg

For details of the inactive ingredients see section 13.

3. WHAT IS THE MEDICINE INTENDED FOR?

The medicine is intended for prevention and treatment of coccidiosis, and diseases caused by the bacteria *Salmonella* and *Pasteurella* in chickens.

Therapeutic group: An antibiotic of the sulfonamide group.

4. CONTRAINDICATIONS

Do not administer to breeding or egg-laying flocks.

Do not administer to animals that are sensitive to sulfonamides or to any of the ingredients of the medicine.

5. SIDE EFFECTS

On occasion, effects of hypersensitivity may appear.

Side effects can be reported to the Ministry of Health by clicking on the link "Reporting Side Effects of Drug Treatment" found on the Ministry of Health home page (www.health.gov.il) that directs you to the online form for reporting side effects, or by entering the link:

<https://sideeffects.health.gov.il>

6. TARGET ANIMALS

Chickens

7. METHOD OF ADMINISTRATION AND DOSAGE

64 grams for every 100 liters of drinking water over 3 consecutive days. In severe cases, treatment may be continued for an additional 3 days at half the dosage after consultation with the veterinarian. Should it be decided to repeat the treatment, this can be done after a two-day break.

8. HOW TO USE THE PREPARATION

Prepare fresh drinking water each day.

Administer to the flock as their sole source of drinking water.

9. WITHDRAWAL PERIOD

Stop the treatment 4 days before slaughter.

10. WARNINGS

• Special warnings regarding use in the target animal

The medicine should be used on the basis of a microbiological susceptibility test on bacteria isolated from the animal. If this is not possible, treatment must be based on regional epidemiological data.

Avoid overdosage: The quantity of water consumed is higher on hot days.

Avoid underdosage: The quantity of water consumed by sick animals is lower.

If there is no clinical improvement within 3 days, an alternative antibiotic treatment should be considered based on susceptibility testing.

• Special warnings regarding the safety of the use of the medicine in animals

At the end of the treatment thoroughly clean the drinking water systems so that no residue of the medicine remain in the system.

• Special warnings regarding the safety of the person handling the preparation

People with known hypersensitivity to sulfonamides or to any of the ingredients of the medicine must avoid contact with the preparation.

Avoid contact with the skin and mucous membranes. Wear protective clothing, safety glasses, gloves and a mask. Do not smoke, eat or drink while handling the preparation.

• Drug interactions and other forms of interactions

Not known

• Incompatibility

In the absence of compatibility studies, do not mix this preparation with other veterinary preparations.

11. STORAGE INSTRUCTIONS

• Avoid poisoning! This medicine and any other medicine should be kept in a closed place out of the reach of children and/or infants in order to avoid poisoning.

• Do not use the medicine after the expiry date (exp. date) that appears on the package. The expiry date refers to the last day of that month.

• Storage conditions: Store below 25°C.

12. INSTRUCTIONS REGARDING DISPOSAL OF THE PREPARATION/ REMNANTS OF THE PREPARATION AFTER USE

Any remnants of a veterinary preparation, or any waste obtained upon use of a veterinary preparation, must be discarded as toxic waste. Do not discard into the sewage system.

13. FURTHER INFORMATION

• In addition to the active ingredient, the medicine also contains dextrose.

• The medicine is packed in bags containing a cream-whitish-colored powder.

• Each package contains either 320 grams or 1 kg. Not all package sizes may be marketed.

• **Registration holder:** Abic Veterinary Products Ltd., P.O.B. 489 Beit Shemesh.

This leaflet was checked and approved by the Ministry of Health in: 12/2021.

Registration number of the medicine in the National Drug Registry of the Ministry of Health: 072-14-91672