

## CONSUMER PACKAGE LEAFLET FOR A VETERINARY PREPARATION

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

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

Sulfamethoprim Veterinary  
Solution per os (oral) for administration  
in drinking water

### 2. ACTIVE INGREDIENTS

Each 1 ml contains:  
Sulfadiazine sodium 86.4 mg  
Trimethoprim 16 mg

For the inactive ingredients see section 13.

### 3. WHAT IS THE MEDICINE INTENDED FOR?

An antibacterial preparation for treatment of  
infections caused by bacteria susceptible  
to the active ingredients in chickens only.

#### Therapeutic group

Sulfadiazine – A sulfonamide antibiotic  
Trimethoprim – An antibacterial substance

### 4. CONTRAINDICATIONS

None known.

### 5. SIDE EFFECTS

Side effects can be reported to the  
Ministry of Health by clicking on the link  
“Report Side Effects of Drug Treatment”  
found on the Ministry of Health homepage  
([www.health.gov.il](http://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 SPECIES

Chickens

### 7. METHOD OF ADMINISTRATION AND DOSAGE

1-2 ml of Sulfamethoprim Veterinary  
per liter of drinking water, for 3 to 5  
consecutive days (1 ml Sulfamethoprim  
Veterinary is intended for treatment of  
10 kg bodyweight).

On hot days the concentration should be  
reduced due to increased water uptake.  
Dilute the contents of the container with  
drinking water until a uniform solution is  
obtained.

### 8. HOW TO USE THE PREPARATION

Administer to the flock as a sole source of  
drinking water during treatment.

### 9. WITHDRAWAL PERIODS

Treatment should be stopped 8 days  
before slaughter. Do not use in laying hens  
producing eggs for human consumption.

### 10. WARNINGS

Special warnings relating to the safety of  
the person handling the preparation

- Sulfonamides may cause hypersensitivity following ingestion, inhalation, contact or on injection. In order to avoid sensitivity from direct contact with the preparation, the person handling the preparation should wear rubber gloves and a mask. Remove clothes that have come in contact with the preparation, and immediately wash eyes and skin should they have come in contact with the preparation.
- If there is known hypersensitivity to one of the ingredients of the preparation, do not come into contact with the preparation. If effects such as a rash develop after exposure to the preparation, see a doctor and show this leaflet.

### 11. INSTRUCTIONS FOR STORAGE

- Avoid poisoning! This medicine, and all other medicines, should be kept in a safe place out of the reach and sight 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 bottle. 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 unused veterinary medicine or waste material derived from the use a veterinary medicine, must be disposed of as toxic waste. Do not discard into the sewage.

### 13. FURTHER INFORMATION

- In addition to the active ingredients, the medicine also contains:  
Saccharine sodium, Propylene glycol, Brown shade\*, Purified water.  
Brown shade: Carmoisine F red no.3, Yellow FDC no.6, Edicol supra red, Ponceau4R.
- What the medicine looks like and the package contents: A plastic bottle containing a clear red-orange liquid.
- Package sizes: 1 liter.
- **Manufacturer and license holder:**  
Abic Veterinary Products Ltd., P.O.B. 489, Bet Shemesh Industrial Estate.

**This leaflet was checked and approved  
by the Ministry of Health in  
May 2019**

**Registration number of the medicine  
in the National Drug Registry of the  
Ministry of Health: 081-05-91497**