



## Ferrosal

### 1. Identification of the veterinary drug

WERAVET 17 Ferrosal  
Liquid dilution for injection

For animals: : horses, cattle, pigs, sheep, goats, dogs, cats

### 2. Qualitative and quantitative composition

1 ampoule of 2 ml (2010.1 mg) contains:

Medically active ingredient:

Ferrum metallicum Dil. C 30

199,1 mg

Ferrum Phosphoricum Dil. C 30

199,1 mg

Other components:

Sodium chloride

Water for injection

### 3. Pharmaceutical form

Liquid dilution for injection

### 4. Pharmacological and toxicological characteristics and, insofar as it serves the purpose for therapeutic use, specifications on pharmacokinetics

#### 4.1 Pharmacodynamic characteristics (including specifications on the substance or indication group; if applicable, ATC vet. Code)

Homeopathic drug

#### 4.2 Pharmacokinetic characteristics

No specifications

#### 4.3 Toxicological characteristics

*Acute toxicity, subchronic and chronic toxicity*

No examination results are available for the acute, subchronic and chronic toxicity of Ferrum metallicum C 30 and Ferrum Phosphoricum C 30.

*Mutagenic and tumour-causing potential*

No examination results are available for the mutagenic and tumour-causing potential of Ferrum metallicum C 30 and Ferrum phosphoricum C 30.

*Reproductive toxicity*

No examination results are available for the reproductive toxicity of Ferrum metallicum C 30 and Ferrum phosphoricum C 30.

### 5. Clinical specifications

#### 5.1 Target animal species

horses, cattle, pigs, sheep, goats, dogs, cats

### 5.2 Indications

The fields of application correspond to the veterinary-homeopathic drug pattern, and include:

- gastro-enteritis
- febrile initial conditions in young animals
- constitutional medicine for weakly anaemic animals with pale translucent skin

Use of the drug should not take place without the advice of a veterinarian and does not replace other drugs and measures prescribed by the veterinarian in this regard.

If the following signs appear, you should consult the veterinarian, since an illness which requires a veterinarian's diagnosis may be present:

- high fever or fever that lasts more than 3 days
- cases of diarrhea lasting longer than 2 days or accompanied by admixtures of blood or black coloration of the stool
- hemorrhagic diathesis and all types of haemorrhages
- hereditary or acquired coagulation defects
- acute and strong inflammatory symptoms such as redness, heat, swelling, pain and dysfunction

The use of WERAVET 17 Ferrosal should take place with consideration of the homeopathic "simile principle". For this purpose, the consultation of a veterinarian for the purpose of determining the disease pattern (homeopathic anamnesis) is recommended.

In the following illnesses, WERAVET 17 Ferrosal may be used only as supporting treatment:

- conditions of insufficiency due to reduced absorption or availability of nutrients required for life
- illnesses which require surgical treatment.

Because of its low iron content, WERAVET 17 Ferrosal cannot be used for iron substitution in iron deficiency therapy and thus cannot replace any iron preparations

- illnesses which require surgical treatment, such as haemorrhages, extensive necroses, large-area burns and tumours.

Prior to the commencement of treatment, the causes of the complaints should therefore be largely clarified.

### 5.3 Contraindications

None known

### 5.4 Side effects (type, frequency and degree of severity)

Note:

In treatment with a homeopathic drug, the existing complaints may temporarily worsen (initial worsening). In this case, you should terminate the use of the drug and consult the veterinarian.

### 5.5 Special precautionary measures for use

Due to the risk of a parenteral application of a drug, the administration of a drug with the same method of acting which is licensed for oral or rectal application is to be preferred.

Indications for intravenous injections are:

- immediate occurrence of the effect
- local irritations of the tissues in subcutaneous or intramuscular injection, due to the components and/or amount of the preparation.

### 5.6 Use during gestation and lactation

No examination results are available regarding the safety of using WERAVET 17 Ferrosal during gestation and lactation.

Like all drugs, homeopathic drugs should only be used during gestation and lactation after consultation with the veterinarian.

### 5.7 Interactions with other drugs and other interactions

The simultaneous administration of an incompatible drug which may negate the effect of WERAVET 17 Ferrosal should be avoided. Please consult your veterinarian.

### 5.8 Dosing with individual and daily specifications, type and duration of use

For intravenous, subcutaneous and intramuscular injection.

horses	3-4 ml
cattle	3-4 ml
pigs	2-3 ml
sheep	2-3 ml
goats	2-3 ml
dogs	1-2 ml
cats	1-2 ml



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In acute conditions, the injection should be repeated after 1 – 2 days; otherwise, repetition takes place after 4 – 8 days.

Administration of the drug should not take place longer than to the complete healing of the animal: A homeopathic drug can trigger a disease pattern which corresponds to the homeopathic drug pattern (symptomatics) in healthy animals.

**5.9 Overdose (symptoms, emergency measures and antidote)**  
No specifications

**5.10 Special warnings for every target animal species**  
No specifications

### 5.11 Waiting period

horses	edible tissues:	0 days
	milk:	0 days
cattle:	edible tissues:	0 days
	milk:	0 days
sheep	edible tissues:	0 days
	milk:	0 days
goats	edible tissues:	0 days
	milk:	0 days
pigs	edible tissues:	0 days

**5.12 Special precautionary measures for the user**  
No specifications

## 6. Pharmaceutical specifications

**6.1 Incompatibilities**  
No specifications

### 6.2 Duration of shelf life

**6.2.1 of the finished drug in the undamaged container**  
5 years

**6.2.2 of the finished drug after the container is opened**  
Use immediately after opening. Opened ampoules must be discarded.

**6.2.3 after mixing the preparation, ready for use**  
Not applicable

**6.3 Special notes regarding storage**  
Do not store above 25 °C.

**6.4 Type and content of the container (package sizes)**  
Original packaging with 10 and 40 ampoules à 2 ml (clear glass)

**6.5 Special precautionary measures for the disposal of unused drugs or other special precautionary measures to avoid dangers to the environment**  
Not applicable

## 7. Name or company and address of the pharmaceutical entrepreneur

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**8. License number**  
6157931.00.00

**9. Date of licensing / renewal**  
05.05.2003

**10. Prescription status / prescription drug in Germany**  
Non prescription drug

**11. Date of the information**  
July 2003