



Osteosal

1. Identification of the veterinary drug

WERAVET 14 Osteosal

Liquid dilution for injection
Active substance: Calcium Carbonicum Hahnemanni C 30

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:
Calcium Carbonicum Hahnemanni 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 Calcium Carbonicum Hahnemanni C 30.

Mutagenic and tumour-causing potential

No examination results are available for the mutagenic and tumour-causing potential of Calcium Carbonicum Hahnemanni C 30.

Reproductive toxicity

No examination results are available for the reproductive toxicity of Calcium Carbonicum Hahnemanni 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:

- homeopathic constitution agents for heavy, coarse-boned animals and young animals
- homeopathic constitution agent for disturbances of the calcium metabolism
- growth and developmental disturbances in young animals
- prevention in the case of inability to rise in cows, due to hypocalcaemia.

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:

- increased muscle tension, muscle trembling and cramping
- behavioural changes (nervous to fearful)
- fever
- acute and strong signs of inflammation, such as redness, heat, swelling, pain and function disturbance
- severe reduction in general state
- lasting, unclear, periodic or newly appearing complaints.

In the case of the disease patterns "disturbances in calcium metabolism", "growth and development disturbances in young animals", and "prevention in the case of inability to rise in cows due to hypocalcaemia", the veterinarian must determine the necessity for allopathic treatment with vitamin and/or mineral substance substitution.

The use of WERAVET 14 Osteosal 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 14 Osteosal 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.

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 14 Osteosal 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 14 Osteosal 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 ampoules 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
6157807.00.00

9. Date of licensing / renewal
07.10.2003

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

11. Date of the information
October 2003