



Dermalis

1. Identification of the veterinary drug

WERAVET 11 Dermalis
Liquid dilution for injection

Active substance: sulfur 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:
Sulfur 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 Sulfur C 30.

Mutagenic and tumour-causing potential

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

Reproductive toxicity

No examination results are available for the reproductive toxicity Sulfur 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:

- dry and weeping eczema
- for stimulation in cases of parasitic eczema
- for loss of hair

- constitutional medicine for all animals with strong, unpleasant body odour
- effectiveness after all infectious diseases as a regeneration remedy established.

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:

- fever
- intensified itching and severe discaling skin
- purulent skin diseases
- longer lasting skin alterations
- acute and strong signs of inflammation, such as redness, heat, swelling, pain and function disturbance
- lasting, unclear, periodic or newly appearing complaints.

In clinical pictures showing injuries or severe alterations of the skin surface the veterinary surgeon is to decide on the necessity of a local treatment.

The use of WERAVET 11 Dermalis 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 11 Dermalis 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 11 Dermalis 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 11 Dermalis 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	1-3 ml
cats	1-3 ml

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.



Dermisal

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, 40 and 100 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

6157888.00.00

9. Date of licensing / renewal

20.12.2002

10. Prescription status / prescription drug in Germany:

non prescription drug

11. Date of information

June 2003