

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

CLiK Extra 65 mg/ml Pour On Suspension for Sheep

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Active substance:

Dicyclanil: 65 mg/ml

Excipients:

Qualitative composition of excipients and other constituents	Quantitative composition if that information is essential for proper administration of the veterinary medicinal product
Methyl parahydroxybenzoate (E218)	1.50 mg
Propyl parahydroxybenzoate	3.00 mg
Butylated hydroxytoluene	0.50 mg
Ponceau 4R	0.05 mg
Polysorbate 20	
Acrylic acid copolymer	
Disodium edetate	
Distilled monoglycerides	
Triglycerides, medium-chain	
Propylene glycol	
Sodium hydroxide	
Purified water	

Pink coloured pour-on suspension.

3. CLINICAL INFORMATION

3.1 Target species

Sheep

3.2 Indications for use for each target species

Prevention of blowfly strike on sheep caused by *Lucilia sericata* or *Wohlfahrtia magnifica*.

3.3 Contraindications

Do not use in cases of hypersensitivity to the active substance or to any of the excipients.

3.4 Special warnings

The veterinary medicinal product is best applied before an anticipated blowfly challenge, or when a blowfly challenge is identified on, or in the vicinity of, the farm. Established strikes may require a separate treatment with a knockdown insecticide. It is recommended that animals with dirty back-ends are daggged or crutched prior to application. If daggging or crutching is undertaken in the weeks following application, these animals should be re-treated, otherwise protection could be lost. Do not apply during heavy rainfall, or when such conditions are expected. The resulting wash out may reduce the protection period.

3.5 Special precautions for use

Special precautions for safe use in the target species:

Not applicable.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

Redness and irritation may develop after skin or eye contact with the veterinary medicinal product.

Contact with skin and eyes should be avoided.

Personal protective equipment consisting of synthetic rubber gloves and PVC trousers should be worn when handling the veterinary medicinal product.

In case of skin contact remove contaminated clothing and thoroughly wash the affected parts of the body with soap and water.

In case of eye contact wash immediately with clean water.

Always wash hands and exposed skin with soap and water after work.

Do not eat, drink or smoke whilst using the veterinary medicinal product.

Residues remain on the fleece for some time after treatment, therefore, it is good agricultural practice to minimise handling of sheep after treatment. If you need to handle sheep within 3 months after treatment, wear synthetic rubber gloves and long trousers or coveralls. If sheep are wet wear waterproof trousers.

Do not shear sheep in the 3 months after treatment.

Special precautions for the protection of the environment:

Treated sheep must be kept away from watercourses for at least one hour after treatment. There is a serious risk to aquatic life if this advice is not followed.

3.6 Adverse events

Target species: Sheep.

None known.

Reporting adverse events is important. It allows continuous safety monitoring of a veterinary medicinal product. Reports should be sent, preferably via a veterinarian, to either the marketing authorisation holder or the national competent authority via the national reporting system. See package leaflet for respective contact details.

3.7 Use during pregnancy, lactation or lay

Pregnancy and lactation:

The safety of the veterinary medicinal product has not been established during pregnancy or lactation. Use only according to the benefit/ risk assessment by the responsible veterinarian.

Laboratory studies in rats and rabbits have not produced any evidence of teratogenic, foetotoxic or maternotoxic effects.

3.8 Interaction with other medicinal products and other forms of interaction

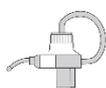
None known.

3.9 Administration routes and dosage

For external use only.

For the 0.8 L, 2.5 L and 5 L containers, unscrew the blue cap from the container and replace it with the white spigot cap provided. If using a manual gun, attach the tube from the application device to the white spigot cap. If using an automatic drenching-gun, connect the device directly to the spigot cap provided. After use, remove the white spigot cap and close the container with the blue cap.

For the 20 L container, if using a manual drenching gun with a spigot cap and a tube, unscrew the white cap from the container and replace it with a spigot cap. Attach the extended tubing to provide flexible extension between the container and applicator gun. If using an automatic drenching gun, connect the device directly to a spigot cap. After use, remove the white spigot cap and close the container with the white cap.



The veterinary medicinal product is applied according to the following recommendations:

Bodyweight (kg)	Dose Volume (ml)
10-20	20
21-30	24
31-50	30
>50	36

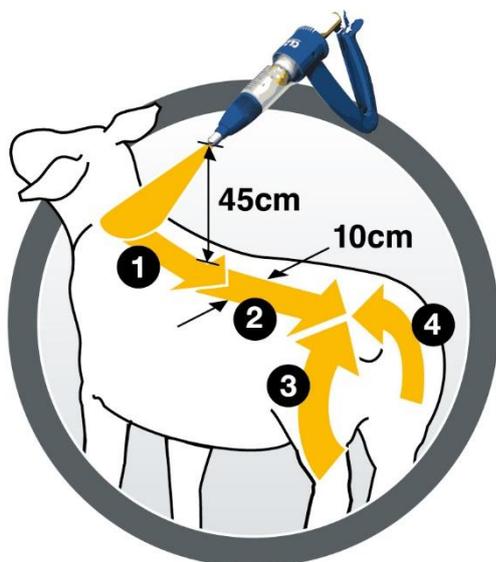
Guide dose volumes correspond to 0.6 – 2 ml [39 – 130 mg dicyclanil] per kg bodyweight.

To ensure administration of a correct dose, bodyweight should be determined as accurately as possible; accuracy of the dosing device should be checked.

If animals are to be treated collectively rather than individually, they should be grouped according to their bodyweight and dosed accordingly, in order to avoid under- or overdosing.

Recommended for treatment in sheep with any wool length, including off-shears. Shake the container well before use. For shaking heavier 20 L containers, the container should be placed on a horizontal surface, and shaken, or rolled, sideways, backwards and forwards on the horizontal surface. Do not dilute with water.

The veterinary medicinal product must be applied with a manual or automatic dosing gun, fitted with a spray nozzle, which guarantees the correct spreading of the product on the fleece. Best results will be achieved by holding the gun approximately 45 cm from the sheep during application. Apply as a fan spray using a 4 stroke method as shown in the pictogram along the spine of the animal in bands at least 10 cm wide from the middle of the shoulders and in an arc around the crutch and tail. Half the dose should be applied along the spine with the remainder over the tail and crutch area.



The veterinary medicinal product should be applied before, or at the start of, predicted fly activity. The veterinary medicinal product will protect against fly strike for 19 weeks. It is good agricultural practice to check animals regularly for fly strike.

3.10 Symptoms of overdose (and where applicable, emergency procedures and antidotes)

An overdose of up to at least 4 times the recommended dose does not lead to any signs of local or systemic intolerance. No antidote is known.

3.11 Special restrictions for use and special conditions for use, including restrictions on the use of antimicrobial and antiparasitic veterinary medicinal products in order to limit the risk of development of resistance.

Not applicable.

3.12 Withdrawal periods

Meat and offal: 40 days.

Not authorised for use in animals producing milk for human consumption.

4. PHARMACOLOGICAL INFORMATION

4.1 ATCvet code: QP53AX24.

4.2 Pharmacodynamics

Dicyclanil prevents the moult from the first to the second larval instar of *Lucilia sericata* and *Wohlfahrtia magnifica*. It is less effective against later larval stages and does not have any adulticidal action. The mode of action of dicyclanil is believed to be similar to that of the triazine compounds.

4.3 Pharmacokinetics

In studies with the 50 mg/ml formulation, after 7 days post dosing, approximately 5 % of the dose was absorbed and eliminated in urine and faeces. Peak blood levels were observed between 12 and 48 hours post dose, accounting for < 0.025 mg dicyclanil equivalents/kg.

In experimental metabolism studies, absorbed drug was widely distributed throughout the body. Longest half-lives were found in liver and kidney being 13 and 10 days respectively. In muscle, fat

and wool, unchanged dicyclanil was found to be the major residue, whereas in liver and kidney the desacyclopropyl dicyclanil was found to be the major residue together with unchanged dicyclanil.

Environmental properties

Dicyclanil has the potential to cause harmful effects on aquatic invertebrates and dung fly larvae. Following use of this veterinary medicinal product, levels of dicyclanil potentially harmful to dung fly larvae have been shown to be excreted in faeces for approximately 4 weeks. Faeces from treated animals may temporarily reduce the abundance of dung fly larvae which may impact on dung degradation.

5. PHARMACEUTICAL PARTICULARS

5.1 Major incompatibilities

In the absence of compatibility studies, this veterinary medicinal product must not be mixed with other veterinary medicinal products.

5.2 Shelf life

Shelf life of the veterinary medicinal product as packaged for sale: 3 years.

Shelf life after first opening the immediate packaging: 1 year.

5.3 Special precautions for storage

Protect from frost.

Store in the original container.

Keep the container tightly closed, away from food, drink and animal feedstuffs.

Protect from direct sunlight.

5.4 Nature and composition of immediate packaging

1 L, 2.5 L and 5 L white opaque polyethylene back pack containers and 20 L white opaque polyethylene container filled with 0.8 L, 2.2 L, 5 L and 20 L respectively. The 2.5 L and 5 L containers are closed with a polypropylene screw cap and the 1 L and 20 L with a polyethylene screw cap.

Not all pack sizes may be marketed.

5.5 Special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products

Medicines should not be disposed of via wastewater.

The veterinary medicinal product should not enter water courses as dicyclanil is dangerous for aquatic organisms.

Do not contaminate ponds or other waterways with product or empty containers.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any national collection systems applicable to the veterinary medicinal product concerned.

6. NAME OF THE MARKETING AUTHORISATION HOLDER

Elanco GmbH

7. MARKETING AUTHORISATION NUMBER(S)

VPA22020/022/001

8. DATE OF FIRST AUTHORISATION

25/11/2016

9. DATE OF THE LAST REVISION OF THE SUMMARY OF THE PRODUCT CHARACTERISTICS

03/11/2025

10. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCTS

Veterinary medicinal product subject to prescription.

Detailed information on this veterinary medicinal product is available in the Union Product Database (<https://medicines.health.europa.eu/veterinary>).