

SUMMARY OF THE PRODUCT OF CHARACTERISTICS

1. NAME OF VETERINARY MEDICINAL PRODUCT

Animec 10 mg/ml Solution for Injection for Cattle, Pigs and Sheep

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ml contains:

Active substance:

Ivermectin 10 mg

Excipients:

For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Solution for injection.

Clear colourless to slight, yellow-coloured solution with no visible particulates.

4. CLINICAL PARTICULARS

4.1 Target Species

Cattle (beef and non-lactating dairy cattle), Sheep and Pigs.

4.2 Indications for use, specifying the target species

Treatment of infections with the following parasites in beef and non-lactating dairy cattle, pigs and sheep:

Cattle:

Gastro-intestinal roundworms (adult and fourth stage larvae):

Ostertagia spp. (including inhibited *O.ostertagi*)

Haemonchus placei

Trichostrongylus axei

Trichostrongylus colubriformis

Cooperia spp.

Oesophagostomum radiatum

Nematodirus helvetianus (adult)

N. spathiger (adult)

Lungworms (adult and fourth stage larvae):

Dictyocaulus viviparus.

Warbles:

Hypoderma bovis

H. lineatum.

Mange mites:

Psoroptes bovis

Sarcoptes scabiei var. *bovis.*

Sucking lice:

Linognathus vituli

Haematopinus eurysternus

The use of the product in cattle should take into account geographical differences in the occurrence patterns of parasites.

Pigs:

Gastrointestinal roundworms: (adult and fourth stage larvae):

Ascaris suum

Hyostromylus rubidus

Oesophagostomum spp.

Strongyloides ransomi (adult only)

Lungworms:

Metastrongylus spp. (adult)

Lice:

Haematopinus suis

Mange mites:

Sarcoptes scabiei var. *suis*

Sheep:

Gastrointestinal roundworms (adult and fourth-stage larvae):

Teladorsagia circumcincta including inhibited larvae

T. trifurcata

Haemonchus contortus including inhibited larvae

Trichostrongylus axei (adults)

T. colubriformis and *T. vitrinus* (adults)

Cooperia curticei

Oesophagostomum columbianum

O. venulosum (adults)

Nematodirus filicollis

Chabertia ovina

Trichuris ovis (adults)

Lungworms:

Dictyocaulus filaria (adult and fourth-stage larvae)

Protostrongylus rufescens (adults)

Nasal bots (all larval stages) :

Oestrus ovis.

Mange mites:

Psoroptes ovis

4.3 Contraindications

Do not administer by the intravenous or intramuscular route.

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

Do not use in dogs or cats as severe adverse reactions may occur (see also section 4.5).

4.4 Special warnings for each target species

In sheep treatment of psoroptic mange (sheep scab) with one injection is not recommended because, although a clinical improvement may be seen, elimination of all mites may not occur.

Sheep scab (*Psoroptes ovis*) is an extremely contagious external parasite of sheep. To ensure complete control great care must be taken to avoid re-infestation, as mites may be viable for up to 15 days off the sheep. It is important that all sheep which have been in contact with infected sheep are treated. Contact between treated, infected and untreated flocks must be avoided until at least seven days after treatment.

Unnecessary use of antiparasitics or use deviating from the instructions given in the SPC may increase the resistance selection pressure and lead to reduced efficacy. The decision to use the product should be based on confirmation of the parasitic species and burden, or of the risk of infestation based on its epidemiological features, for each herd or flock.

Repeated use for an extended period, particularly when using the same class of substances, increases the risk of resistance development. Within a herd or flock, maintenance of susceptible refugia is essential to reduce that risk. Systematically applied interval-based treatment and treatment of a whole herd or flock should be avoided. Instead, if feasible, only selected individual animals or subgroups should be treated (targeted selective treatment). This should be combined with appropriate husbandry and pasture management measures. Guidance for each specific herd/flock should be sought from the responsible veterinarian.

Resistance to ivermectin has been reported in *Cooperia spp.* and in *Ostertagia ostertagi* in cattle. Resistance has also been reported in *Haemonchus contortus* in cattle outside the EU.

In sheep, resistance to ivermectin is widespread in *Teladorsagia circumcincta*, *Trichostrongylus spp.*, *Haemonchus contortus* and in other gastro-intestinal parasite species.

Multiple resistance was reported in *Teladorsagia circumcincta* to benzimidazoles, macrocyclic lactones and levamisole and in *Haemonchus contortus* to ivermectin and benzimidazoles.

Multiple resistance to macrocyclic lactones has also been reported in *Psoroptes ovis* scab mites in sheep and in cattle.

The use of this product should take into account local information about susceptibility of the target parasites, where available. It is recommended to further investigate cases of suspected resistance, using an appropriate diagnostic method (e.g. Faecal Egg Count Reduction Test). Confirmed resistance should be reported to the marketing authorisation holder or to the competent authorities.

4.5 Special precautions for use

Special precautions for use in animals

In cattle, to avoid secondary reactions due to the death of Hypoderma larvae in the oesophagus or in the spine it is recommended to administer the product at the end of warble fly activity and before the larvae reach their resting sites. Consult your veterinarian on the correct timing of treatment.

Avermectins may not be well tolerated in non-target species. Cases of intolerance with fatal results are reported in dogs – especially Collies, Old English Sheepdogs and related breeds and crosses, and also in turtles/tortoises. In addition, care should be taken to avoid ingestion of spilled product or access to used containers by these other species.

Since ivermectin is highly bound to plasma proteins, special care should be taken in cases of sick animals or in nutritional conditions associated with low plasma protein levels.

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

'This product may cause eye and skin irritation. Avoid contact with skin or eyes. In case of skin or eye contact, wash exposed area with plenty of clean water. If symptoms persist, seek medical advice.

Take care to avoid accidental self-injection: the product may cause local irritation and/or pain at the site of injection. In case of accidental self-injection, seek immediate medical advice and show the information leaflet or the label to the physician.

Do not eat or smoke while handling the product.

Wash hands after use.

Other precautions

The product is very toxic to aquatic organisms and dung insects.

Long term effects on dung insects caused by continuous or repeated use cannot be excluded therefore repeated treatment of animals on a pasture with an ivermectin-containing product within a season should only be given in the absence of alternative treatments or approaches to maintain animal/flock health, as advised by a veterinarian.

Treated cattle should not have direct access to ponds, streams or ditches for 14 days after treatment.

4.6 Adverse reactions (frequency and seriousness)

Cattle

Transitory discomfort has been observed in some cattle. A low incidence of soft tissue swelling at the injection site has been observed. These reactions have disappeared without treatment.

Pigs

Mild and transient pain reactions may be seen in some pigs.
All these reactions disappeared without treatment.

Sheep

Immediately following subcutaneous injection, activity suggesting pain, sometimes intense but usually transient, has been observed in some sheep.

4.7 Use during pregnancy, lactation or lay

The product can be administered during pregnancy and lactation in cows, ewes and sows. The product does not affect fertility. It can be used in breeding cows and bulls, breeding ewes and rams, in sows and boars.

4.8 Interaction with other medicinal products and other forms of interaction

Do not combine with vaccination against lungworm. If vaccinated animals are to be treated, treatment should not be carried out within a period of 28 days before or after vaccination.

4.9 Amounts to be administered and administration route

For single administration only by subcutaneous injection. Each ml contains 10 mg of ivermectin sufficient to treat 50 kg of bodyweight of cattle and sheep, and 33 kg of bodyweight of pigs. Replace with a fresh sterile needle after every 10 to 12 animals. Massage the injection site after administration of the product. Injection of wet or dirty animals is not recommended.

Underdosing could result in ineffective use and may favour resistance development. To ensure a correct dosage, body weight should be determined as accurately as possible. If animals are to be treated collectively, reasonably homogeneous groups should be set up, and all animals of a group should be dosed at the rate corresponding to the heaviest one.

Accuracy of the dosing device should be thoroughly checked.

Cattle:

The product should be given only by subcutaneous injection at the recommended dosage level of 200 µg ivermectin per kg bodyweight under the loose skin in front of, or behind, the shoulder in cattle. This is equivalent to 1 ml per 50 kg bodyweight.

The volume administered per injection site should not exceed 10 ml.

Pigs:

In pigs, the recommended dosage level is 300 µg ivermectin per kg bodyweight. This is equivalent to 1 ml per 33 kg bodyweight. The recommended route of administration is by subcutaneous injection into the neck. The volume administered per injection site should not exceed 5ml.

Young Pigs:

In young pigs, especially those below 16 kg for which less than 0.5 ml of the product is indicated, dosing accurately is important. The use of a syringe that can accurately deliver as little as 0.1 ml is recommended.

Sheep

The recommended dose is 200 µg ivermectin per kg bodyweight (corresponding to 1 ml of the product per 50 kg bw) by subcutaneous injection over the neck.

The volume administered per injection site should not exceed 1ml.

For the treatment and control of sheep scab (*Psoroptes ovis*), two injections with a seven-day interval are required to treat clinical signs of scab and to eliminate mites.

In young lambs weighing less than 25 kg give 0.1 ml of the product per 5 kg. The use of a syringe that can deliver as little as 0.1 ml is recommended.

This product does not contain any antimicrobial preservative. Swab septum before removing each dose.

When using the 200, 250 or 500ml pack sizes, use only automatic syringe equipment. For the 50ml pack size, use of a multiple dose syringe is recommended. To refill the syringe, use of a draw-off needle is recommended to avoid excessive broaching of the stopper.

4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary

In the case of overdose a symptomatic treatment should be given.

Cattle

Single doses of 4.0 mg ivermectin per kg (20 x the use level) given subcutaneously resulted in ataxia and depression.

Pigs

A dose of 30 mg ivermectin per kg (100 x the recommended dose of 0.3 mg per kg) injected subcutaneously to pigs caused lethargy, ataxia, bilateral mydriasis, intermittent tremors, laboured breathing and lateral recumbency.

Sheep

At dose levels up to 4 mg ivermectin per kg (20 x the use level) given subcutaneously resulted in ataxia and depression.

No antidote has been identified; however, symptomatic therapy may be beneficial.

4.11 Withdrawal Period(s)

Cattle

Meat and offal: 49 days.

Do not use in lactating cows producing milk for human consumption. Do not use in cattle producing milk for human consumption or in dairy cows within 60 days prior to calving.

Pigs

Meat and offal: 28 days.

Sheep

Meat and offal: 25 days

Do not use in lactating sheep producing milk for human consumption.

Do not use in sheep within 60 days of lambing where milk is to be used for human consumption

5. PHARMACOLOGICAL PROPERTIES

Pharmacotherapeutic Group: Endectocides
ATCvet code: QP54AA01.

5.1 Pharmacodynamic properties

Ivermectin is a member of the macrocyclic lactone class of endectocides which have a unique mode of action. Compounds of this class bind selectively and with high affinity to glutamate-gated chloride ion channels which occur in invertebrate nerve and muscle cells. This leads to an increase in the permeability of the cell membrane to chloride ions with hyperpolarization of the nerve or muscle cell, resulting in paralysis and death of the parasite. Compounds of this class may also interact with other ligand gated chloride channels, such as those gated by the neurotransmitter gamma-aminobutyric acid (GABA). The margin of safety for compounds of this class is attributable to the fact that mammals do not have glutamate-gated chloride channels, the macrocyclic lactones have a low affinity for other mammalian ligand-gated chloride channels and they do not readily cross the blood-brain barrier.

Multidrug-resistance to macrocyclic lactones is mediated by the efflux activity of ATP-binding cassette (ABC) transporters such as P-glycoproteins. Selection of resistant isolates with ivermectin leads to cross-resistance to eprinomectin and moxidectin depending upon the underlying mechanism of resistance. Molecular mechanisms of resistance involve mutations in several genetic loci associated with alterations of the glutamate-and GABA-gated chloride channels reducing the binding affinity of the molecules of macrocyclic lactones.

5.2 Pharmacokinetic properties

Cattle

Maximum plasma concentration

At a dose level of 200 µg ivermectin per kg a maximum plasma concentration of 35-50 ng/ml is reached in ± 2 days.

It is also established that ivermectin is carried mainly in the plasma (80%). This distribution between plasma and blood cells remains relatively constant.

Excretion: length of time and route

Only about 1-2% is excreted in the urine the remainder is excreted in the faeces, approximately 60% of which is excreted as unaltered drug. The remainder is excreted as metabolites or degradation products.

Pig

Maximum plasma concentration

During trials carried out at a dose level of 300 µg ivermectin per kg bodyweight, peak plasma concentrations were reached in 3 (±0.5) days.

Excretion: length of time and route

Biliary excretion is also the major route of ivermectin excretion in pigs.

Sheep

At a dose level of 200 µg ivermectin per kg bodyweight, a mean C_{max} of 10.976 ng/mL was reached at a mean T_{max} of 40 hours, and the mean elimination half-life was 73.9 hours.

The major route of excretion of ivermectin and its metabolites in sheep is faeces (99 %) with 1 % excreted in the urine.

Environmental properties

Like other macrocyclic lactones, Ivermectin has the potential to adversely affect non-target organisms. Following treatment, excretion of potentially toxic levels of Ivermectin may take place over a period of several weeks. Faeces containing Ivermectin excreted onto pasture by treated animals may reduce the abundance of dung feeding organisms which may impact on the dung degradation.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Glycerol
Glycerol Formal

6.2 Major Incompatibilities

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

6.3 Shelf-life

Shelf life of the veterinary medicinal product as packaged for sale: 3 years
Shelf life after first opening the immediate packaging: 28 days.

6.4 Special precautions for storage

This veterinary medicinal product does not require any special storage precautions.

6.5 Nature and composition of immediate packaging

Multidose polyethylene bottles of 50 ml, 250 ml and 500 ml sealed with bromobutyl seals and aluminium overseals.
Not all pack sizes may be marketed.

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

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.
The product should not enter water courses as this may be dangerous to fish and other aquatic organisms.

7. MARKETING AUTHORISATION HOLDER

Chanelle Pharmaceuticals Manufacturing Ltd
Loughrea
Co. Galway
H62 FH90
Ireland

8. MARKETING AUTHORISATION NUMBER

Vm 08749/3141

9. DATE OF THE FIRST AUTHORISATION

21 September 2000

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

September 2025

Gavin Hall
Approved: 11 November 2025