SUMMARY OF PRODUCT CHARACTERISTICS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Dysect Sheep Pour-on solution 12.5 g/l.

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each litre contains:

Active substances:
Alphacypermethrin 12.5 g

Excipients:

Qualitative composition of excipients and other constituents	**Quantitative composition if that information is essential for proper administration of the veterinary medicinal product**
Copper Chlorophyll Complex (E141)	0.6 g
Phytorob 926/67	
Benzoic acid	

Clear dark green liquid free from solids.

3. CLINICAL INFORMATION

3.1 Target species

Sheep.

3.2 Indications for use for each target species

For the control and treatment of blow fly strike, on sheep and lambs and for the treatment of lice and ticks on sheep. Reduces the incidence of headfly strike in sheep and lambs for up to 6 weeks.

3.3 Contraindications

Do not treat lambs under one week old.
Do not treat animals in very hot weather.

3.4 Special warnings

None.

3.5 Special precautions for use

Special precautions for safe use in the target species:
For external use only.

Because alphacypermethrin is retained in the wool grease of the fleece, it is recommended that the product be applied to sheep with a minimum wool length of 1.0cm.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Personal protective equipment consisting of waterproof apron, boots, and impervious gloves should be worn when handling the veterinary medicinal product and recently treated animals.
Use the product in a well-ventilated area.
Wash any splashes from the skin and eyes immediately, using plenty of water.
Remove heavily contaminated clothing immediately and wash before re-use.
Wash hands and exposed skin with soap and water after treatment and before eating drinking and smoking.

Special precautions for the protection of the environment:
Not applicable.

3.6 Adverse events

Sheep:Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Fleece discolouration[1]

[1] Observed from approximately 6 to 8 weeks after treatment.

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 the 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 and lactation.

3.8 Interaction with other medicinal products and other forms of interaction

None known.

3.9 Administration routes and dosage

The veterinary medicinal product is applied at a rate of 40 ml for animals weighing 25 kg or more, and 25 ml for animals weighing less than 25 kg. The dose is applied using an adjustable dosing gun fitted with a T-bar with 3 or 4 outlets and is applied directly onto the wool in a band from the back of the head to the base of the tail. For the reduction of headfly strike 5ml per animal is applied between the horns, irrespective of size of the animal.

Blowfly - For the control of blowfly a single application will normally provide 8 to 10 weeks protection. Re-treatment may be necessary after this period. For the treatment of blowfly strike, a single treatment applied to the infected area will ensure blowfly larvae are killed.
Ticks - A single treatment will normally provide control of ticks for 8 to 12 weeks.

Headfly - A single treatment will reduce the incidence for up to 6 weeks.

Lice - A single treatment will normally kill all lice.

	DOSAGE (ml)	
PEST	**SHEEP (>25KG)**	**LAMBS (<25KG)**
Blowfly	40	25
Lice	40	
Ticks	40	
Headfly	5	5

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

No adverse reactions were recorded in double dose exposure trials.

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: 49 days
Milk: not permitted for use in animals producing milk for human consumption.

4. PHARMACOLOGICAL INFORMATION

4.1 ATCvet code: QP53AC08

4.2 Pharmacodynamics

Alphacypermethrin is a synthetic pyrethroid, which works as an insecticide and repellent by inhibiting neural transmission in insects.

4.3 Pharmacokinetics

Alphacypermethrin is lipophyllic and is retained in the wool grease of the sheep fleece to provide protection from blowflies for up to 8 to 10 weeks.

5. PHARMACEUTICAL PARTICULARS

5.1 Major incompatibilities

None known.

5.2 Shelf life

Shelf life of the veterinary medicinal product as packaged for sale: 2 years.

5.3 Special precautions for storage

Do not store above 25 °C.
Store in a dry place.
Store in the original container.
Keep the container tightly closed.
Protect from direct sunlight
Do not freeze.
If contents freeze, they must be thawed and thoroughly mixed before use.
Store away from food, drink, and animal feeding stuffs.

5.4 Nature and composition of immediate packaging

Container: Fluorinated HDPE backpack dispenser of 2.5 litre and 5 litre content.
Closure: Aluminium overseal with polyethylene foam wadding and a polypropylene cap.
Dosing device: Pour-on applicator with a glass filled nylon body and polypropylene barrel.

Not all pack sizes may be marketed.

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

Medicines should not be disposed of via wastewater.
DANGEROUS to fish and aquatic life. Do not contaminate ponds, waterways or ditches with the product or empty container.

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

Zoetis Belgium S.A.

7. MARKETING AUTHORISATION NUMBER

Vm 60021/3007

8. DATE OF FIRST AUTHORISATION

10 December 1999

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

January 2026

10. CLASSIFICATION OF VETERINARY MEDICINAL PRODUCT

Veterinary medicinal product subject to prescription.

Find more product information by searching for the ‘Product Information Database’ on www.gov.uk.

Gavin Hall

Approved: 18 March 2026