

United Kingdom
Veterinary Medicines Directorate
Woodham Lane
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# **NATIONAL PROCEDURE**

# PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY MEDICINAL PRODUCT

Molemec Injection for Cattle and Sheep 10 mg/ml Solution for Injection



# **PRODUCT SUMMARY**

Name, strength and pharmaceutical form	Molemec Injection for Cattl Injection	e and S	sheep 10	mg/ml	Solution	for
Applicant	Merial Animal Health Ltd					
Active substance(s)	Ivermectin					
ATC Vetcode	QP54AA01					
Target species	Cattle and sheep					
Indication for use	Indicated for the effective treatment and control of the following parasites of cattle and sheep:  CATTLE					
	PARASITE Gastrointestinal Roundworms	Adult		L4	Inhibite L4	ed
	Ostertagia lyrata	•		•		
	Ostertagia ostertagi	•		•	•	
	Cooperia oncophora	•		•		
	Cooperia pectinata	•		•		
	Cooperia punctata	•		•		
	Haemonchus placei	•		•		
	Trichostrongylus axei	•		•		
	Trichostrongylus colubriformis	•		•		
	Bunostomum	_		_		
	phlebotomum	•		•		
	Oesophagostomum	•		•		
	radiatum					
	Strongyloides papillosus	•				
	Nematodirus helvetianus	•				
	Nematodirus spathiger	•				
	Trichuris spp.	•				
	Lungworms					
	Dictyocaulus viviparus	•		•		
	Eye Worms					
	Thelazia spp	•				

#### Warbles

Hypoderma bovis H. lineatum

# Mange Mites

Psoroptes ovis Sarcoptes scabei var. bovis

### **Sucking Lice**

Linognathus vituli Haematopinus eurysternus Solenopotes capillatus

The product may also be used as an aid in the control of biting lice (*Damalinia bovis*) and the mange mite *Chorioptes bovis*, but complete elimination may not occur.

#### **Persistent Activity**

Given at the recommended dosage of 1ml per 50kg bodyweight, the product

controls re-infection with the following nematodes up to the duration shown:

Parasite	No. of Days After Treatment
Barbers pole worm - Haemonchus placei	14
Small intestinal worm - Cooperia spp.	14
Hairworm – Trichostrongylus axei	14
Brown stomach worm - Ostertagia ostertagi	21
Nodular worm - Oesophagostomum radiatum	21
Lungworm – Dictyocaulus viviparus	28

To obtain optimal benefit from the persistent activity for grazing animals, it is recommended that calves which are set-stocked in the first grazing season

should be treated 3, 8 and 13 weeks after the day of turn-out. This can protect the animals from parasitic gastroenteritis and lungworm disease throughout the grazing season, provided they are set-stocked, all the calves are included in the programme and that no untreated cattle are added to the pasture.

Treated animals should continue to be monitored according to good husbandry practices.

<u>SHEEP</u>			
DADACITE	A al14	1.4	Inhibited
PARASITE	Adult	L4	L4
Gastrointestinal			
Roundworms			
Ostertagia circumcincta	•	•	•
O. trifurcata	•	•	
Haemonchus contortus	•	•	•
Trichostrongylus axei	•		
T. colubriformis	•	•	
T. vitrinus	•		
Cooperia curticei	•	•	
Oesophagostomum	•	•	
columbianum			
O. venulosum	•		
Nematodirus filicollis	•	•	
Chabertia ovina	•	•	
Trichuris ovis	•		
L			
Lungworms			
Dictyocaulus filaria	•	•	
Protostrongylus rufescens	•		

# **Nasal Bots**

Oestrus ovis

# **Mange Mites**

Psoroptes ovis\*

Benzimidazole-resistant strains of *Haemonchus contortus* and *Ostertagia circumcincta* are also controlled.

# **MODULE 2**

The Summary of Product Characteristics (SPC) for this product is available on the Veterinary Medicines Directorate website (<a href="www.vmd.defra.gov.uk">www.vmd.defra.gov.uk</a>)



#### **PUBLIC ASSESSMENT REPORT**

	Informed Consent application in accordance
application	with Article 13c of Directive 2001/82/EC as amended.

### I. SCIENTIFIC OVERVIEW

The quality / safety / efficacy aspects of this product is/are identical to Ivomec Classic Injection for Cattle and Sheep (Ivermectin). The initial application for Ivomec Classic Injection for Cattle and Sheep (Ivermectin) was assessed before there was a requirement to have a public assessment report, therefore no details in this section are available.

#### II. OVERALL CONCLUSION AND BENEFIT- RISK ASSESSMENT

The data submitted in the dossier demonstrate that when the product is used in accordance with the Summary of Product Characteristics, the benefit/risk profile for the target species is favourable and the quality and safety of the product for humans and the environment is acceptable.



#### POST-AUTHORISATION ASSESSMENTS

The SPC and package leaflet may be updated to include new information on the quality, safety and efficacy of the veterinary medicinal product. The current SPC is available on the Product Information Database of the Veterinary Medicines Directorate website.

(www.gov.uk/check-animal-medicine-licensed)

The post-authorisation assessment (PAA) contains information on significant changes which have been made after the original procedure which are important for the quality, safety or efficacy of the product.

The PAA for this product is available on the Product Information Database of the Veterinary Medicines Directorate website.

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