



**Veterinary
Medicines
Directorate**

**United Kingdom
Veterinary Medicines Directorate
Woodham Lane
New Haw
Addlestone
Surrey KT15 3LS**

National Procedure

**PUBLICLY AVAILABLE ASSESSMENT REPORT FOR A VETERINARY
MEDICINAL PRODUCT**

Dicyclanil Elanco 5% Pour-on Suspension

Date created: December 2021

MODULE 1**PRODUCT SUMMARY**

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| Name, strength and pharmaceutical form | Dicyclanil Elanco 5% Pour-on Suspension for Sheep |
| Applicant | Elanco Europe Ltd Form 2, Bartley Way Bartley Wood Business Park Hook RG27 9XA United Kingdom |
| Active substance | Dicyclanil (ISO) |
| ATC Vetcode | QP53AX24 |
| Target species | Sheep |
| Indication for use | Prevention of blowfly strike due to <i>Lucilia sericata</i> on sheep. Prevention of blowfly strike due to <i>Wohlfahrtia magnifica</i> on sheep |

MODULE 2

The Summary of Product Characteristics (SPC) for this product is available on the Product Information Database of the Veterinary Medicines Directorate.

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

MODULE 3**PUBLIC ASSESSMENT REPORT**

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| Legal basis of original application | Informed consent application in accordance with Article 13 (c) of Directive 2001/82/EC as amended. |
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I. SCIENTIFIC OVERVIEW

The quality / safety / efficacy aspects of these products are identical to those of CliK 5% Pour-on Solution. The initial application for CliK 5% Pour-on Solution 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 of the product(s) is favourable.

MODULE 4**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.

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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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