Veterinary Medicines Directorate

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Post Authorisation Assessments

MAQS Formic Acid 68.2g Beehive Strips for Honey Bees

•	15 June 2022	Change in the address of the marketing authorisation holder from NOD Apiary Ireland Ltd., 5 George's Dock, IFSC Dublin 1, D01 X8N7, Ireland to NOD Apiary Ireland Ltd., Tullow Industrial Estate, Tullow, Co Carlow, Ireland R93 W0D8. Addition of a manufacturer of a finished product responsible for importation. Addition of a manufacturer responsible for batch release. Addition of a secondary packaging site of a finished product.
•	16 March 2022	Addition of new distributor: Andermatt Biocontrol UK Limited, Unit 30, Mackley Industrial Estate, Henfield Road, Small Dole, West Sussex, BN5 9XR, UK.
•	08 January 2021	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product for solid pharmaceutical forms. Change in container closure system of the finished product. Change in the number of units (sachets) in a pack outside the range of the currently approved pack sizes of the finished product.
•	11 February 2020	Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	16 December 2019	Change in distributor details: addition of a distributor E.H. Throne (Beehives) Ltd, Beehive Business Park, Rand, Market Rasen, Lincolnshire, LN8 5NJ and removal of distributor BCW Agriculture Ltd, Unit 8, Burnside Business Park, Burnside Road, Market Drayton, Shropshire, TF9 3UX.
•	23 April 2019	Replacement of a manufacturer responsible for batch release.
•	22 January 2019	Change of MAH, from NOD Europe Ltd., 5 St Paul's Square, Old Hall Street, Liverpool, L3 9AE to NOD Apiary Ireland Ltd., 5 George's Dock, IFSC Dublin 1, D01 X8N7, Ireland.
•	22 August 2018	Change in RMS from UK to IE.
•	18 April 2018	Renewal – UK RMS
•	22 June 2017	Changes to the quality control testing arrangements for the active substance – addition of a site where batch control / testing takes place.

		Change in the manufacturer of an intermediate used in the manufacturing process of the active/ change in the manufacturer of the active where no Ph. Eur. Certificate of Suitability is part of the approved dossier.
•	08 December 2016	To add Bee-Equipment Ltd as an additional distributor.
•	31 August 2016	Changes to an existing pharmacovigilance system as described in the DDPS.
•	04 May 2016	Minor change in the manufacturing process of the finished product.
•	12 June 2013	Approval of mock ups for 2, 3 and 5 dose pack sizes.