



Post Authorisation Assessments

Tri-Solfen Cutaneous Solution for Pigs

Vm 50406/3028

08 April 2025	Change in the shelf-life or storage conditions of the finished product. Change in the storage conditions of the finished product. Change in the specification parameters and/or limits of the finished product. Change in the specification parameters and/or limits of the finished product.
08 April 2025	Change in legal entity of MA holder from Dechra Limited, Snay gill Industrial Estate, Keighley Road, Skipton, North Yorkshire, BD23 2RW, United Kingdom to Dechra Regulatory B.V., Handelsweg 25, 5531 AE Bladel, The Netherlands.
18 May 2024	Change in the shelf-life of the finished product. Change in storage conditions of the finished product or the diluted/reconstituted product.
20 September 2023	Update to Ph. Eur. CEP for the active substance Adrenaline.
07 November 2022	Updated certificate of suitability from an already approved manufacturer.
01 November 2022	Change in address of a microbiological testing site.
01 November 2022	Editorial changes to part 2 of the dossier.
25 May 2022	Deletion of a non-significant parameter of an active substance.
10 May 2022	Change in the colour of the of the polypropylene screw and spigot caps of the finished product packaging from orange/ turquoise to Rhodamine Red C.
10 May 2022	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
21 April 2022	Variation to add secondary packaging component to keep all primary packaging components together.
20 April 2022	Change in distributor details from Dechra Limited, Snaygill Industrial Estate, Keighley Road, Skipton, North Yorkshire, BD23 2RW, UK to Dechra Veterinary Products Ltd, Sansaw Business Park, Hadnall, Shrewsbury, Shropshire, SY4 4AS, UK.
25 February 2022	Replacement to a test procedure for the finished product.