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Post Authorisation Assessments

Trimacare 80 Tablets for Dogs

Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
Submission of an updated certificate of suitability.
Change of distributor details.
Grouped variation to update European Pharmacopoeia Certificates of Suitability for already approved active substance manufacturers.
Submission of an updated European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer.
Variation to update the product literature and outer packaging.
Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005.
Deletion of an active substance manufacturer.
Renewal.
Variation to change the legal category from POM to POM-V.
Submission of an updated European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer.
Submission of an updated European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer.
Identical changes to a number of products.
Renewal.
Change of active substance manufacturer.