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Post Authorisation Assessments

Fiprofile 402 mg/3600 mg Spot-on Solution for Very Large Dogs

Change in the contact details of the QPPV of an existing
pharmacovigilance system as described in the DDPS.
Minor change in the manufacturing process of the
finished product.
Change in the name of the manufacturer of the finished
product.
Changes in the qualitative and quantitative composition
of the immediate packaging of the finished product.
Increase in the shelf-life of the finished product as
packaged for sale, from 2 to 3 years.
Change in the specification limits of the finished product.
Change in the specification limits of the finished product.
Change in RMS from UK to IT
Change of specifications of a former non
Pharmacopoeial active substance to comply with the Ph.
Eur. monograph 2869.