VPA10774/027/003

Rycarfa 100 mg tablets for dogs

Variation	Summary	Date
Vet - A2	VNRA - Vet - A2 Vet - A2 Administrative changes -	30/09/25
	Change in the (invented) name of the veterinary medicinal	
	product VRA-S - Vet - G.I.18 Vet - G.I.18 - One-off alignment	
Vet - G.I.18	of the product information with version 9.0* of the QRD	08/08/25
	templates i.e. major update of the QRD templates in	
	accordance with Regulation (EU) 2019/6, for veterinary	
	medicinal products authorised in accordance with	
	Directive 2001/82/EC or Regulation (EC) No 726/2004	
Vet - C6	VNRA - Vet - C6 - Introduction of a summary of the	01/12/23
	PSMF or changes to the summary of the PSMF not already covered elsewhere in the Annex to Regulation	
	(EU) 2021/17 - C6 Changes to the safety, efficacy and	
	pharmacovigilance part of the dossier: Introduction of a	
	summary of the PSMF or changes to the summary of the	
	PSMF not already covered elsewhere in the Annex to	
	Regulation (EU) 2021/17	
Vet - B47 d)	VNRA - Vet - B47 d) - d) To reflect compliance with the	06/03/23
	Ph. Eur. by removing reference to the internal test method	
	and test method number - B47d) Changes to the quality	
	part of the dossier: Change to comply with Ph. Eur. or with a national pharmacopoeia of a Member State: — to	
	reflect compliance with the Ph. Eur. by removing	
	reference to the internal test method and test method	
	number	
Vet - B44(Do not use)	VNRA - Vet - B44 - Submission of a new or updated Ph.	07/12/22
	Eur. CEP from an already approved manufacturer for a	
	non-sterile active substance, starting material, reagent or	
	intermediate, excipient - B44 Changes to the quality part	
	of the dossier: Submission of a new or updated Ph. Eur.	
	CEP from an already approved manufacturer for a non-sterile: — active substance; — starting material,	
	reagent or intermediate used in the manufacturing process	
	of the active substance; — excipient	
B.I.b.2.a	IA - B.I.b.2.a - a) Minor changes to an approved test	03/03/22
	procedure - B.I.b.2.a - QUALITY CHANGES - ACTIVE	
	SUBSTANCE - Control of active substance - Change in	
	test procedure for active substance or starting	
	material/reagent/intermediate used in the manufacturing	
	process of the active substance - Minor changes to an approved test procedure	
	approved test procedure	