VPA22693/002/001

Qualimec 10 mg/ml Solution for Injection.

Variation	Summary	Date
Vet - A2	VNRA - Vet - A2 Vet - A2 Administrative changes -	11/11/25
	Change in the (invented) name of the veterinary medicinal	
	product	
Vet - B47 b)	VNRA - Vet - B47 b) Vet - B47 b) - Change to comply with	26/06/25
	Ph. Eur. or with a national pharmacopoeia of a Member State:	
	change to comply with an update of the relevant monograph of	
	the Ph. Eur. or national pharmacopoeia of a Member State	
Vet - B47 b)	VNRA - Vet - B47 b) - b) Change to comply with an update of	
	the relevant monograph of the Ph. Eur. or national	06/03/25
	pharmacopoeia of a Member State - B47 b) Changes to the	
	quality part of the dossier: Change to comply with Ph. Eur. or	
	with a national pharmacopoeia of a Member State: — change	
	to comply with an update of the relevant monograph of the Ph.	
	Eur. or national pharmacopoeia of a Member State	
Vet - A2	VNRA - Vet - A2 - Change in the (invented) name of the	03/03/25
	veterinary medicinal product - A2 Administratvie changes:	
	Change in the (invented) name of the veterinary medicinal	
	product	
	VNRA - Vet - A2 - Change in the (invented) name of the	10/01/25
Vet - A2	veterinary medicinal product - A2 Administratvie changes:	
, , , , , , , , , , , , , , , , , , , ,	Change in the (invented) name of the veterinary medicinal	
	product	
Vet - A2	VNRA - Vet - A2 - Change in the (invented) name of the	10/01/25
	veterinary medicinal product - A2 Administrative changes:	
	Change in the (invented) name of the veterinary medicinal	
	product VNIDA Vat. CC. Introduction of a summary of the DSME or	
Vet - C6	VNRA - Vet - C6 - 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 - C6	13/12/24
	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 - B3 e)	VNRA - Vet - B3 e) - e) Deletion of a test procedure - B3 e)	
	Changes to the quality part of the dossier: Deletion of a test	
	procedure — for the active substance or a starting material,	
	reagent or intermediate of the active substance; —for the	22/02/24
	immediate packaging of the active substance; — for an	
	excipient or the finished product; —for the immediate	
	packaging of the finished product	
Vet - B3 d)	VNRA - Vet - B3 d) - d) Deletion of a non-significant	22/02/24
	specification parameter (active substance, starting material,	
	intermediate - B3 d) Changes to the quality part of the dossier:	

	D1.: C :: C . : C	
	Deletion of a non-significant specification parameter (e.g.	
	deletion of an obsolete parameter) of — an active substance;	
	— a starting material; —an intermediate or reagent used in the	
	manufacturing process of the active substance	
Vet - C10 a)	VNRA - Vet - C10 a) - a) Administrative information	15/12/23
	concerning the holder's representative - C10 a) Changes to the	
	safety, efficacy and pharmacovigilance part of the dossier:	
	Changes to the labelling or the package leaflet which shall not	
	be connected with the SPC: — administrative information	
	concerning the holder's representative	
Vet - A2	VNRA - Vet - A2 - Change in the (invented) name of the	15/11/23
	veterinary medicinal product - A2 Administratvie changes:	
	Change in the (invented) name of the veterinary medicinal	
	product	
Vet - F.II.e.5 b)	VRA-S - Vet - F.II.e.5 b) - b) Change in the fill weight/fill	17/02/23
	volume of sterile multidose (or single-dose, partial use)	
	parenteral medicinal products, including	
	1 '	
	biological/immunological medicinal products F.II.e.5 b)	
	Quality Changes - Container closure system -Change in pack	
	size of the finished product - Change in the fill weight/fill	
	volume of sterile multidose (or single-dose, partial use)	
	parenteral medicinal products, including	
	biological/immunological medicinal products.	
Vet - F.II.e.1 b) 2.	VRA-S - Vet - F.II.e.1 b) 2 b) Change in type of container or	17/02/23
	addition of a new container 2. Sterile medicinal products and	
	biological/ immunological medicinal products - F.II.e.1 b) 2.	
	1	
	immediate packaging of the finished product - Change in type	
	of container or addition of a new container - Sterile medicinal	
	products and biological/immunological medicinal products	