

Related Tool Ref: RT_VS&PV_01b

ADVERSE EVENT REPORT FORM

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EVEN	Т ТҮРЕ		NAME AND	ADRESS OF S	ENDER		SERIOU	SNESS			
Adverse reaction Lack of efficacy Off label use Environmental problems Infectious agent Residues				harmacist	□ Death □ Life threatening □ Significant disability □ Congenital anomaly □ Other important medical condition □ Not serious						
PATIENT(S)	Animal (s) \square Humain being(s) \square (for human beings, fill only age and sex bellow)										
Species	Breed	Sex	Statuts	Age/DOB	Weight (Kg)	Health status before start of Masivet					
		F □ M □	Neutered Pregnant Intact				Good air oor ritical Inknown				
Medical history/Risk factors											
SUSPECT DRUG (Masivet@)											
Start date	Route		Dosage/Frequency	End date	Indication		Batch number- Expiry date				
Masivet @ administered by: Veterinarian											
VETERINARY MEDICINAL PRODUCTS ADMINISTERED BEFORE THE ADVERSE EVENT (if more products are administered concurrently than the number of boxes available, please duplicate this form)											
Medicinal Veterinary Product (VPM)	Route		Dosage/Frequency	Start date	Administered by: Veterinian/Animal owner/Other		End date	Indication			
Has the Health Authorities (HA) been informed?											

SUSPECTED ADVERSE REACTION DATE	Time between first administration and		Number treated Number reacted		Ouration of adverse reaction						
Evolution: Death Date:/ Euthanasia Recovered with sequelae Recovered without sequelae Ongoing Date:/											
DESCRIPTION OF THE EVENT – Indicate also if the reaction has been treated, how and with what was the result?											
ı											
Causality assessment related to adverse reaction	o Masivet® and	Probable Possible Unclassified Unconclusive Unlikely									
Necropsie File number :	Investiga	ations 🗌 :									
OTHER RELEVANT DATA available) copy of medical repo				carried out,	copy of necropsy report (if						
HUMAN EXPOSURE CASE ((if the reported case	refers to huma	n being, please also co	omplete the	details of exposure below.)						
Contact with treated animal											
Oral ingestion											
Topical expositionn											
• Others											
Exposure dose:											
Date:	Place:	Name a	nd signature of sender	:							
Contact point (phone) (if differ	rent from number giv	ven on page 1)									

(*)AB Science, represented by its Chief Executive Officer, is responsible for personal data, intended to record and evaluate event reports of adverse events within the scope of veterinary pharmacovigilance defined in Articles R.5141-94 to R.5141-10 of the French Public Health Code). In accordance with the provisions of Regulation (EU) 2016/679 on the protection of individuals with regard to the processing of personal data and the data flow (GDPR) and the Data Privacy Protection and Rights - Act of 6 January 1978 amended, you have a right of access, rectification, limitation and, in certain cases, data erasure your private information. You may also, for legitimate reasons, forbid your private data processing.