

Adverse event report form

Formulaire à faire parvenir à Pharmacovigilance@ab-science.com Fax: 33 (0)1 47 20 10 82 AB SCIENCE 3 avenue Georges V 75008 - PARIS	
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EVENT TYPE	NAME AND ADDRESS OF SENDER	SERIOUSNESS
<input type="checkbox"/> Adverse reaction <input type="checkbox"/> Lack of efficacy <input type="checkbox"/> Off label use <input type="checkbox"/> Environmental problems <input type="checkbox"/> Residues <input type="checkbox"/> Infectious agent	Veterinarian <input type="checkbox"/> Pharmacist <input type="checkbox"/> Other <input type="checkbox"/> _____ Phone _____ Fax _____ E-mail _____	<input type="checkbox"/> Death <input type="checkbox"/> Life threatening <input type="checkbox"/> Significant disability <input type="checkbox"/> Congenital anomaly <input type="checkbox"/> Other important medical condition <input type="checkbox"/> Not serious

PATIENT/S Animal (s) Human being(s) (for human beings, fill only age and sex below)

Species	Breed	Sex	Statut	Age (DOB)	Weight (kg)	Health status before start of masitinib
		F <input type="checkbox"/> M <input type="checkbox"/>	Neutered <input type="checkbox"/> Pregnant <input type="checkbox"/> Intact <input type="checkbox"/>			Good <input type="checkbox"/> Fair <input type="checkbox"/> Poor <input type="checkbox"/> Critical <input type="checkbox"/> Unknown <input type="checkbox"/>

Medical history/Risk factors _____

SUSPECT DRUG (Masivet@)

Start date	Route	Dasage/ Frequency	End date	Indication	Batch number-Expiry date

Masivet @ administered by Veterinarian Animal owner Other

VETERINARY MEDICINAL PRODUCTS ADMINISTERED BEFORE THE ADVERSE EVENT

Medicinal veterinary product (VMP)	Route	Dosage/ Frequency	Start date	Administered by Veterinarian/Animal owner/Other	End date	Indication

Has the Health Authorities (HA) been informed?

Yes No

SUSPECTED ADVERSE REACTION DATE ____/____/____	Time between first administration and event: _____	Number treated Number reacted	Duration of adverse reaction _____
Evolution : Death <input type="checkbox"/> Euthanasia <input type="checkbox"/> Recovered with sequelae <input type="checkbox"/> Recovered without sequelae <input type="checkbox"/> Ongoing <input type="checkbox"/>			
Date : ____/____/____			
DESCRIPTION OF THE EVENT Indicate also if the reaction has been treated, how and with what was the result?			
Causality assesment related to Masivet@ and adverse reaction _____		Probable <input type="checkbox"/> Possible <input type="checkbox"/> Unclassified <input type="checkbox"/> Inconclusive <input type="checkbox"/> Unlikely <input type="checkbox"/>	
Necropsy <input type="checkbox"/> File number _____ Investigations <input type="checkbox"/> : _____			
OTHER RELEVANT DATA (attach further papers if necessary : investigations carried out, copy of necropsy report, copy of medical reports for human beings exposure or ongoing)			
HUMAN EXPOSURE CASE (if the reported case refers to human being, please also complete the details of exposure below)			
<ul style="list-style-type: none"> • Contact with treated animal <input type="checkbox"/> • Oral ingestion <input type="checkbox"/> • Topical exposure <input type="checkbox"/> • Others <input type="checkbox"/> 			
Dose d'exposition :			
Date: _____ Place: _____ Name and signature of sender: _____			
Contact point if different from number given on page 1 _____			