Adverse Event Reporting Form

A. Patient Detail (Animal data)

Species:			Breed/production type:		
Sex		Physiological status:			
□ female	□ male	□ pregnant	□ neutered	□ lactating	□ other
Weight (kilogram):			Age:		
State of health at time of treatment:					
□ good	□ fair	□ poor	□ critical	□ unknown	□ other

Reason(s) for treatment (prevention of what disease(s) or initial diagnosis) :

B. Adverse Reaction Details

Adverse Reaction Term (s):	Onset Date: Date & Duration of Reaction:

Description of adverse events: (including all clinical signs, site of reaction, severity, with specific diagnosis, treatment and action taken):

Outcome of the Event: ○ **Recovered** ○ **Not Recovered** ○ **Recovered with sequelae** ○ **Fatal** ○ **Unknown**

Lab test details (Report, if any):

C. Drug Details

Name of the drug: **Strength:**

Indication:

Batch No:		Expiry date:	
Dose and frequency:		Route and site of administration:	
Start date of treatment:	**Stop date:**	Who administered the product: □ veterinarian □ owner □ other	
Use according to label:	□ yes	□ unknown	□ no, explain:
Action taken after reaction:	□ drug withdrawn	□ dose reduced	□ other, explain:

Additional suspect drug (if any), details as above:

Concomitant medications (provide with details):

D. Veterinarian (If not the reporter)	
Name:	
Address: _ **Pin code:** **Tel no.:**	
E. Reporter Details	
Name:	**Occupation:** [ ] Animal Owner [ ] Pharmacist [ ] Other, specify:……
Address:	**Also reported to:** [ ] Regulatory authority [ ] Distributor [ ] None
Tel No:	**Date:** **Signature:**
Email: **Send this report to**	**To be filled by Manufacturer**
Regulatory & Pharmacovigilance Deptt, Zenex Animal Health India Private Ltd, 9th Floor, N.G. Tower, Satellite Cross Roads Ahmedabad-380015, Gujarat, India Fax: 02717-6663727 or 079-26868687 Email: enquiry@zenexah.com	Date of received: Name & Sign of receiver Safety report ID Report type: [ ] Initial [ ] Follow up, number: …….

Confidentiality: Customers identity will be held confidential and shall remain protected. Submission of a report does not constitute an admission that veterinary personnel or manufacturer or the product caused or contributed to the event.