

<b>Adverse Event Form</b>	<b>No: ART</b>
Date received at MA department:	Staff initials:

<b>1. Reporter/Veterinarian</b>	
Date report:	
Veterinary practise:	
Name reporter:	Position:
Address	
City:	
Phone number	
<b>2. Animal</b>	
Name patient (animal and last name owner):	
Animal species/breed/gender:	Species: Breed: Gender:
Age:	
Weight:	
Physical condition (neutered, in foal etc:)	
Condition (relevant medical history):	
<b>3. Used product in adverse event</b>	
Name product:	
Concentration (if applicable):	
Batch no:	
Expiry date:	
Doses, frequency:	Dose: <input type="checkbox"/> ml / <input type="checkbox"/> gr / <input type="checkbox"/> tabs / <input type="checkbox"/> caps / <input type="checkbox"/> chew ( <i>mark what is applicable</i> ) Frequency:  Administered by: <input type="checkbox"/> veterinarian / <input type="checkbox"/> owner :  <input type="checkbox"/> up dosing phase OR <input type="checkbox"/> maintenance phase
Date product used for the first time: (first dose in course administered)	Day:                    Month:                    Year:  If unknown please indicate how long the patient is on the product: approx.                    year(s) and                    months
Date & Time last administration:	<b>(dose that caused the onset of the adverse event):</b>  Day:                    Month:                    Year:  Time:                    Dose:                    ml / gr / tabs / caps / chew (mark what is applicable)
Date and time onset adverse event:	Day:                    Month:                    Year:  Time event became visible: (for example: 30 minutes after administration, or 2 days after administration)

Treatment continued after adverse event	<input type="checkbox"/> Yes, as usual <input type="checkbox"/> Yes, with dose reduction as follows: <input type="checkbox"/> No	
<b>4. Other drugs taken during the time of the adverse event (when applicable) and reason</b>		
YES <input type="checkbox"/> / NO <input type="checkbox"/>		
<b>If yes, please specify below:</b>		
Name drug:		
Batch no:		
Doses, frequency:		
Date of administration:		
Reason/indication:		
<b>5. Data concerning side effects</b>		
Adverse event in <b>keywords</b>		
Narrative		
Adverse reaction treated with:	<input type="checkbox"/> None <input type="checkbox"/> Dose reduction to <input type="checkbox"/> Treatment stopped <input type="checkbox"/> Therapeutic intervention:	
Date treated:		
Response to corrective treatment:		
Outcome of adverse event:	<input type="checkbox"/> Recovered: <input type="checkbox"/> Stabilized with rest symptoms Description of rest symptoms:	
Date of outcome:	<input type="checkbox"/> Death <input type="checkbox"/> Other:	
<b>6. Narrative of treatment of adverse event. How and with which results?</b>		
Laboratory data available: <input type="checkbox"/> Yes (please provide) <input type="checkbox"/> No		
<b>7. Do you think that the reaction was caused by Artuvetrin?</b>		
<input type="checkbox"/> probable <input type="checkbox"/> possible <input type="checkbox"/> unlikely <input type="checkbox"/> unassessable		
<b>8. Other relevant history or information</b>		

Signature reporter:

Date: