AE report date:			D	IVI	IVI	IVI	I	I	I	I			
AE start date:		Γ	D	Μ	Μ	Μ	Y	Y	Y	Y]		
AE end date:	□ Ongoing	Γ	D	Μ	Μ	Μ	Y	Y	Y	Y			
Adverse Event Information													
Participant or infant?	Participa	nnt	□ Infant										
Condition/Diagnosis:													
AE Term (MedDRA Coding):													
Event Description: Include a history of the event chron outcomes and any other relevant in tests/data, treatment/procedures, m	nformation not captu edical history, treat	ured	elsew	here									
Action taken with study medicat No change Other medication(s) started fo 	or AE:			/ mec	licati	on p	erma				tinued ntinued		
Clinical outcome: Recovered/resolved Not yet recovered Unknown 			Reco Study						ied				

Site No.

Subject No.

This section to be completed by the Investigator only

Severity/Intensity

Image: Severe

*If the AE meets the definition of an SAE, complete the SAE form instead

Reporting Centre										
Delegate's Name:										
Signature:										
Investigator's Name:										
Signature:										
Date:	D	D	Μ	Μ	Μ	Y	Y	Y	Y	