

Participant ID							ADVERSE EVENTS LOG			

Description of adverse event (provide additional information on notes pages if required)	Date of onset DD/MM/YYYY	Date reported to Investigator /team DD/MM/YYYY	Severity 1. Mild 2. Moderate 3. Severe	Causality 1. Unrelated 2. Possible 3. Probable 4. Definite	Action taken – please list all that apply 1. None 2. Hospitalisation 3. Intervention stopped 4. Intervention reduced 5. Intervention interrupted 6. Con Meds commenced * 7. Other (specify)	Outcome 1. Recovered 2. Ongoing 3. Disability or incapacity 4. Death 5. Unknown	Is this a Serious AE? YES** or NO	Date resolved (Enter date resolved or tick if ongoing at end of study) DD/MM/YYYY	PI Signature and Date DD/MM/YYYY
	//___	_/_/___						_/_/___ or <input type="checkbox"/>	_/_/___
	//___	_/_/___						_/_/___ or <input type="checkbox"/>	_/_/___
	//___	_/_/___						_/_/___ or <input type="checkbox"/>	_/_/___
	//___	_/_/___						_/_/___ or <input type="checkbox"/>	_/_/___
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	//___	_/_/___						_/_/___ or <input type="checkbox"/>	_/_/___

* Record on Con Meds Log

** If adverse event meets criteria for a serious adverse event (SAE), please submit an online SAE report within 24 hours of becoming aware of the event