



IND SAFETY REPORT LOG

Protocol Title:	Principal Investigator:
	GCO # _____ Protocol #: _____

Adverse Event	Date of Event	Mfr. Report # or Report's Unique Identifier	Is the event <u>serious, unexpected & associated</u> with the research?		Does the research protocol place any subjects or others at a greater risk of harm?		Will the consent form need to be modified?		PI initials
			<u>Yes</u>	<u>No</u>	<u>Yes</u>	<u>No</u>	<u>Yes</u>	<u>No</u>	