ADVERSE EVENT TRACKING LOG

Principal Investigator: _____ Study #: _____ Sponsor: _____

Study Title: _____

#	Subject ID	Date of Event	Date PI Aware	Description of Event	Serious		Expected		Related to Study Drug			Reported to MHRIRB*	Reported to Sponsor*	Reported to FDA*
					Yes	No	Yes	No	Yes	No	Unk			
1					☐	☐	☐	☐	☐	☐	☐			
2					☐	☐	☐	☐	☐	☐	☐			
3					☐	☐	☐	☐	☐	☐	☐			
4					☐	☐	☐	☐	☐	☐	☐			
5					☐	☐	☐	☐	☐	☐	☐			
6					☐	☐	☐	☐	☐	☐	☐			
7					☐	☐	☐	☐	☐	☐	☐			
8					☐	☐	☐	☐	☐	☐	☐			
9					☐	☐	☐	☐	☐	☐	☐			
10					☐	☐	☐	☐	☐	☐	☐			
11					☐	☐	☐	☐	☐	☐	☐			
12					☐	☐	☐	☐	☐	☐	☐			

**** If applicable***