SUMMARY OF CLINICAL TRIAL ADVERSE EVENTS

IRB Summary Form
(Form Date 11/2017)

Study Title:

Patient ID:	**Patient Initials**
Enrollment Date:	**Off Study/Completion Date:**
General Summary of subjects progress in the study:	

ADVERSE EVENT DESCRIPTION	ADVERSE EVENT WORST GRADE	RELATIONSHIP TO STUDY MEDICATION/DEVICE (PER INVESTIGATOR)

Study Nurse Signature: _____ **Date:** _____