A3.22 NON-SERIOUS ADVERSE EVENTS LOG (PAGE ONE)

Has the participant experienced any Adverse Events since signing the Informed Consent to the trial?	**YES, specify below**	1	**NO**	0

AE no.	**Adverse Event** [Diagnosis or symptom (if known) or signs/symptoms]	**Start Date** (dd/mm/yyyy)	**Stop Date** (dd/mm/yyyy)	**Severity** Mild = 1 Moderate = 2 Severe = 3	**Relationship to Study Drug?** Definite = 1 Probable = 2 Possible = 3 Remote = 4 None = 5	**Action Taken with Study Drug** None (1) Temp. Dose Reduction = 2 Perm. Dose Reduction = 3 Temp. Discontinuation = 4 Perm. Discontinuation = 5	**Staf Initia & Dat**
1				1	1	1	
				2	2	2	
				3	3	3	
					4	4	
					5	5	
2				1	1	1	
				2	2	2	
				3	3	3	
					4	4	
					5	5	
3				1	1	1	
				2	2	2	
				3	3	3	
					4	4	
					5	5	