ARISTOTLE

UCL

Serious Adverse Event (SAE) Report

Patient trial number: ARI – ☐☐☐

Concomitant medications (continued)

Drug Name	Brand	Indication	Dose	Units	Frequency	Route	Start date (d d m m m y y)	Ongoing?	End date (d d m m m y y)
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	
								Y N	

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Serious Adverse Event (SAE) Report

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Additional Information *(use for event summary description and relevant medical history/concurrent conditions)*