CARDAMON Concomitant Medications				Patient Trial Nu	mber: CAR-	Initials	
<b>Type of report:</b> (delete as appropriate)	SAE Report	Urgent Event (TMA) Form	Lactatio	onal Exposure	Report		
Concomitant N	Y Only include non-IMP drugs given	within the 30 days prior to event onset, (exclu	ding treatment for event,	if applicable)			
Drug Name	Indication	Total Daily Dose Prior to this SAE (include units)	Frequency	Route	Date of <u>First</u> Administration of Drug AND Date of <u>Last</u> Administration of Drug Prior to this SAE dd-mm-yyyy		
					First	Ongoing	
					Last		
					First		
					Last		
					First		
					Last		
					First		
					Last		
					First		
					Last		
					First		
					Last		
					First	Ongoing	
					Last		
					First	Ongoing	
					Last		
					First	Ongoing	
					Last		
					First	Ongoing	
					Last		
					First	Ongoing	
					Last		
					First	Ongoing	
					Last		
For UCL CTC use only: Date form r	received:	_ Date form checked:	Date f	orm entered:	Initia	als:	