Non Serious Adverse Drug Reactions Report

Start Date:2016-06-22 End Date:2016-06-23

Study :EMR200136_583	Investigator :NA		Country of Investigator :Romania	SiteNo:005		
Subject No :005-0007	Subject Initials :	DOB :05/06/1966	Sex:Female	Race:Caucasian	Height:165(cm)	Weight:70(kg)
First administration date of batch :			Batch number :			

Study Drug	Start Date	Dose	Change in Dose
Visit 5 (Month12)/Early Termination			
Visit 1/ Baseline (Day 1)	03/05/2015	9	

Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity
local erythema and induration local site injection	04/10/2015			Suspected	Mild

Causality Factors	Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)	Drug withdrawn	Led to study termination	Ongoing		

Event description:

Subject received concomitant medications

Name of medication	Start Date	Ongoing	End Date	Dose	Unit	Frequency

Does the subject have any relevant past or present medical conditions:Yes

Condition	Start Date	Related to study condition	Ongoing
caesarean section			No
uterine fibromas			Yes

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No Data between these 2016-06-22 and 2016-06-23