	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-05	5-19 End Date:2016-05-2	0			•	
Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05		
Subject No :C05-0001	Subject Initials :XFZ	DOB :01/21/1988	Sex:Female	Race:Asian	Height:159cm	Weight:50kg
First administration date of batch :			Batch number :			
Study Drug	udy Drug Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	08/01/2015		225			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	08/16/2015	08/18/2015		Related	Severe	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)		Dose not changed	Concomitant medication **	Resolved		
Event Description: ivgtt qd	Hydroxyethyl Starch 130/	0.4 and Sodium Chlor	ride Injection 500ml,iv	gtt bid calcium glucor	nate injection 10ml+dext	trose injection 500ml
Subject received c	concomitant medications:					
Does the subject h	nave any relevant past or	present medical cond	itions:Yes			
Condition				Start Date	Related to study condition	Ongoing
oophorocystectom	у			UK-Jul-2013	Not on treatment/medicatio n	
salpingoplasty				UK-Feb-2014	Not on treatment/medicatio n	
salpingitis after pre	evious tubal occlusion			UK-Feb-2014	Not on treatment/medicatio n	Ongoing

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