	Non Se	rious Adv	erse Drug	Reactions	Report		
Start Date:2016-01-1	1 End Date:2016-01-15				-		
Study :EMR700623-541	Investigator : Fei Gong	;	Country of Investigator :China	SiteNo:C01			
Subject No :C01-0158	Subject Initials :T-Y	DOB :06/17/1983	Sex:Female	Race:Asian	Height:163cm	Weight:47kg	
First administration date of batch :		Batch number :	:				
Study Drug	Start Date		Dose	Change in Dose	'		
Gonal-f New Pen Stimulation Treatmen	09/16/2015		112.5				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	10/12/2015	10/15/2015		Related	Moderate		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
None(Othervalue:)		Not applicable	Concomitant procedure**	Resolved			
Event Description:		I.	· I			1	
Subject received cond	comitant medications:						
Does the subject hav	ve any relevant past or pr	esent medical conditio	ns:Yes				
Condition				Start Date	Related to study condition	Ongoing	
right fallopian tube is less patency,left fallopian tube is obstruction				06/04/2015	Not on treatment/medication	Ongoing	
Endometrial polyps h	yperplasia			08/24/2015	Not on treatment/medication	Ongoing	

11-JAN-16 Fei Gong/EMR700623-541/C01-0158