	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-05-	-26 End Date:2016-05-2	7			•		
Study :EMR700623-541	Investigator :Fei Gon	g	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 Treatment	09/16/2015		112.5				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	10/11/2015	10/12/2015		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:	Abdominal distension, Na	ausea, yellow urine wi	th less volume, chest p	elvic effusion,ascites	puncture 12-Oct-2015,2	500ml	
Subject received co	oncomitant medications:						
Does the subject ha	ave any relevant past or	present medical cond	itions: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/medicatio n	Ongoing	
Endometrial polyps hyperplasia				08/24/2015	Not on treatment/medicatio n	Ongoing	

26-MAY-16

Fei Gong/EMR700623-541/C01-0158