

## Non Serious Adverse Drug Reactions Report

Start Date:2015-06-29 End Date:2015-06-30

<b>Study</b> :EMR700623-541	<b>Investigator</b> :Fei Gong		<b>Country of Investigator</b> :China	<b>SiteNo</b> :C01		
<b>Subject No</b> :C01-0001	<b>Subject Initials</b> :TTW	<b>DOB</b> :05/13/1988	<b>Sex</b> :Female	<b>Race</b> :Asian	<b>Height</b> :157cm	<b>Weight</b> :41.5kg
<b>First administration date of batch</b> :			<b>Batch number</b> :			
<b>Study Drug</b>	<b>Start Date</b>		<b>Dose</b>	<b>Change in Dose</b>		
Gonal-f New Pen Stimulation Treatment	05/16/2015		113			
<b>Adverse Event</b>	<b>Start Date</b>	<b>End Date</b>	<b>Time related to study treatment</b>	<b>Causality to study drug</b>	<b>Severity</b>	
OHSS	06/06/2015	06/16/2015		Related	Moderate	
<b>Causality Factors</b>		<b>Action Taken with Study Treatment</b>	<b>Other action taken</b>	<b>Outcome</b>	<b>AE Special Interest</b>	<b>AE dose limiting toxicity</b>
None(Othervalue:)		Not applicable	Concomitant procedure**	Resolved		
Event Description:						
Subject received concomitant medications:						
<b>Does the subject have any relevant past or present medical conditions</b> :Yes						
<b>Condition</b>				<b>Start Date</b>	<b>Related to study condition</b>	<b>Ongoing</b>
Bilateral fallopian tube obstruction				04/11/2014	Not on treatment/medication	Ongoing

29-JUN-15

Fei Gong/EMR700623-541/C01-0001

