

## Non Serious Adverse Drug Reactions Report

Start Date:2015-06-01 End Date:2015-06-03

<b>Study</b> :EMR700623-541	<b>Investigator</b> :NA		<b>Country of Investigator</b> :Korea	<b>SiteNo</b> :K01		
<b>Subject No</b> :k01-049	<b>Subject Initials</b> :PGR	<b>DOB</b> :02/20/1983	<b>Sex</b> :Female	<b>Race</b> :Asian	<b>Height</b> :168cm	<b>Weight</b> :62kg
<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	05/02/2015	300				
<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	05/13/2015	05/29/2015		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:)		Dose not changed	Concomitant medication **,Led to study termination	Resolved		
<b>Event description:</b>						
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>
laparoscopic ovary cystectomy				UK-UKN-2004	Not on treatment/medication	

01-JUN-15

NA/EMR700623-541/k01-049

