

## Non Serious Adverse Drug Reactions Report

Start Date:2015-05-28 End Date:2015-05-29

<b>Study</b> :EMR700623-541		<b>Investigator :NA</b>		<b>Country of Investigator :Korea</b>		<b>SiteNo:K01</b>	
<b>Subject No</b> :k01-040	<b>Subject Initials</b> :LBH	<b>DOB</b> :03/05/1985	<b>Sex</b> :Female	<b>Race</b> :Asian	<b>Height</b> :164cm	<b>Weight</b> :61kg	
<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	04/06/2015		225				
<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	04/15/2015	04/30/2015		Related	Mild		
<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	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> No							
<b>Condition</b>				<b>Start Date</b>	<b>Related to study condition</b>	<b>Ongoing</b>	

29-MAY-15

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