

## Non Serious Adverse Drug Reactions Report

Start Date:2015-07-02 End Date:2015-07-06

<b>Study</b> :EMR700623-541	<b>Investigator</b> :NA		<b>Country of Investigator</b> :Korea	<b>SiteNo</b> :K01		
<b>Subject No</b> :k01-0059	<b>Subject Initials</b> :LKH	<b>DOB</b> :08/05/1981	<b>Sex</b> :Female	<b>Race</b> :Asian	<b>Height</b> :153cm	<b>Weight</b> :74kg
<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	06/02/2015		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	06/13/2015	07/02/2015		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>

06-JUL-15

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