Non Serious Adverse Drug Reactions Report

Start Date:2015-10-14 End Date:2015-10-21

Study :EMR700623-541	**Investigator :**NA		**Country of Investigator :China**	**SiteNo:**C02		
Subject No :C02-0111	**Subject Initials :**CXW	**DOB :**10/10/1979	**Sex:**Female	**Race:**Asian	**Height:**162cm	**Weight:**68.0kg
First administration date of batch :			**Batch number :**			

Study Drug	**Start Date**	**Dose**	**Change in Dose**
Gonal-f New Pen Stimulation Treatment	05/20/2015	200	

Adverse Event	**Start Date**	**End Date**	**Time related to study treatment**	**Causality to study drug**	**Severity**
OHSS				Related	Moderate

Causality Factors	**Action Taken with Study Treatment**	**Other action taken**	**Outcome**	**AE Special Interest**	**AE dose limiting toxicity**
None(Othervalue:)	Not applicable	Led to study termination	Resolved		

Event description: Moderate OHSS patients, improved canceled after embryo transfer.

Subject received concomitant medications:

Does the subject have any relevant past or present medical conditions: Yes

Condition	**Start Date**	**Related to study condition**	**Ongoing**
Ectopic pregnancy laparoscopic surgery, the left fallopian tube embryo window.	Uk-Unk-2012	Not on treatment/medication	
Tubal lipiodol angiography	Uk-Unk-2013	Not on treatment/medication	