Non Serious Adverse Drug Reactions Report

Start Date:2016-05-27 End Date:2016-05-30

Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05		
Subject No :C05-0001	Subject Initials :XFZ	DOB :01/21/1988	Sex:Female	Race:Asian	Height:159cm	Weight:50kg
First administration date of batch :			Batch number :			

Study Drug	Start Date	Dose	Change in Dose
Gonal-f New Pen Stimulation Treatment	08/01/2015	225	

Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	08/16/2015	08/18/2015		Related	Severe	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)		Not applicable	None	Resolved		

Event Description:Hydroxyethyl Starch 130/0.4 and Sodium Chloride Injection 500ml，ivgtt bid calcium gluconate injection 10ml+dextrose injection 500ml ivgtt qd

Subject received concomitant medications:

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

Condition	Start Date	Related to study condition	Ongoing
oophorocystectomy	UK-Jul-2013	Not on treatment/medication	
salpingoplasty	UK-Feb-2014	Not on treatment/medication	
salpingitis after previous tubal occlusion	UK-Feb-2014	Not on treatment/medication	Ongoing

27-MAY-16

Ying Zhong/EMR700623-541/C05-0001

Non Serious Adverse Drug Reactions Report

Start Date:2016-05-27 End Date:2016-05-30

Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05		
Subject No :C05-0021	Subject Initials :FYZ	DOB :08/28/1990	Sex:Female	Race:Asian	Height:150cm	Weight:45kg
First administration date of batch :			Batch number :			

Study Drug	Start Date	Dose	Change in Dose
Gonal-f New Pen Stimulation Treatment	08/04/2015	225	

Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity
OHSS	08/19/2015	08/21/2015		Related	Mild

Causality Factors	Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
Protocol procedure**,Concomitant medication**(Othervalue:)	Not applicable	None	Resolved		

Event Description:abdominal distention;Drug : Hydroxyethyl Starch 130/0.4 and Sodium Chloride Injection 500ml , ivgtt bid calcium gluconate injection 10ml+dextrose injection 500ml ivgtt qd 19-Aug-2015 / 21-Aug-2015

Subject received concomitant medications:

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

Condition	Start Date	Related to study condition	Ongoing
salpingocatheterism	UK-Unk-2012	Not on treatment/medication	

29-MAY-16

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