

## Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541		Investigator :NA		Country of Investigator :China		SiteNo:C02	
Subject No :C02-0210		Subject Initials :GLW	DOB :05/08/1983	Sex:Female	Race:Asian	Height:160cm	Weight:55.0kg
First administration date of batch :				Batch number :			
Study Drug		Start Date		Dose		Change in Dose	
Gonal-f New Pen Stimulation Treatment		08/28/2015		225			
Adverse Event		Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS		09/15/2015	09/20/2015		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:							
Subject received concomitant medications:							
Does the subject have any relevant past or present medical conditions:Yes							
Condition				Start Date		Related to study condition	Ongoing
HSG: the right fallopian tube obstruction				Uk-Unk-2014		Not on treatment/medication	

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NA/EMR700623-541/C02-0210

# Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541	Investigator :NA	Country of Investigator :China	SiteNo:C02			
Subject No :C02-0215	Subject Initials :L-X	DOB :08/05/1989	Sex:Female	Race:Asian	Height:165cm	Weight:56.0kg
First administration date of batch :			Batch number :			
Study Drug	Start Date	Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/28/2015	150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/14/2015	09/18/2015		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:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions:Yes						
Condition				Start Date	Related to study condition	Ongoing
HSG: bilateral tubal occlusion				Uk-Unk-2014	Not on treatment/medication	
Under laparoscopy surgery to clear the fallopian tubes, pelvic adhesions dissection				Uk-Unk-2014	Not on treatment/medication	

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NA/EMR700623-541/C02-0215

# Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541	Investigator :NA	Country of Investigator :China	SiteNo:C02			
Subject No :C02-0217	Subject Initials :YQX	DOB :02/10/1987	Sex:Female	Race:Asian	Height:157cm	Weight:60.0kg
First administration date of batch :			Batch number :			
Study Drug	Start Date	Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/31/2015	187.5				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/15/2015	09/20/2015		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 occurs canceled embryo transfer, OHSS improvement						
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 therapy chemotherapy to kill embryos				Uk-Unk-2011	Not on treatment/medication	
HSG: bilateral tubal obstruction.				Uk-Unk-2010	Not on treatment/medication	
Laparoscopy surgery: pelvic adhesions dissection, bilateral tubal surgery to clear				Uk-Unk-2010	Not on treatment/medication	

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NA/EMR700623-541/C02-0217

# Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541	Investigator :NA	Country of Investigator :China	SiteNo:C02			
Subject No :C02-0219	Subject Initials :YLZ	DOB :06/17/1986	Sex:Female	Race:Asian	Height:160cm	Weight:46.0kg
First administration date of batch :			Batch number :			
Study Drug	Start Date	Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/31/2015	150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/15/2015	09/20/2015		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 occurs canceled embryo transfer, OHSS improvement						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions:Yes						
Condition				Start Date	Related to study condition	Ongoing
HSG: bilateral tubal obstruction.				Uk-Unk-2011	Not on treatment/medication	
HSG: bilateral tubal patency, pelvic adhesions.				Uk-Unk-2014	Not on treatment/medication	

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NA/EMR700623-541/C02-0219

# Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541	Investigator :NA	Country of Investigator :China	SiteNo:C02			
Subject No :C02-0220	Subject Initials :CCT	DOB :05/15/1985	Sex:Female	Race:Asian	Height:152cm	Weight:51.0kg
First administration date of batch :			Batch number :			
Study Drug	Start Date	Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/31/2015	150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/15/2015	09/20/2015		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
HSG: bilateral tubal patency.				Uk-Unk-2014	Not on treatment/medication	

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NA/EMR700623-541/C02-0220

# Non Serious Adverse Drug Reactions Report

Start Date:2016-04-11 End Date:2016-04-12

Study :EMR700623-541	Investigator :NA	Country of Investigator :China	SiteNo:C02			
Subject No :C02-0222	Subject Initials :YMD	DOB :04/19/1983	Sex:Female	Race:Asian	Height:158cm Weight:42.0kg	
First administration date of batch :			Batch number :			
Study Drug	Start Date	Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/31/2015	225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/16/2015	09/20/2015		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 occurs canceled embryo transfer, OHSS improvement						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions:Yes						
Condition			Start Date	Related to study condition	Ongoing	
HSG: the side of tubal passable.			Uk-Unk-2012	Not on treatment/medication		
Laparoscopic surgery: bilateral tubal surgery, right fallopian tube patency.			Uk-Unk-2012	Not on treatment/medication		

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NA/EMR700623-541/C02-0222

