	Non Se	rious Adv	erse Drug	Reactions	s Report			
Start Date:2016-08-	17 End Date:2016-08-1	8			_			
Study :EMR700623-541	Investigator :Fei Gon	g	Country of Investigator :China	SiteNo:C01				
Subject No :C01-0001	Subject Initials :TTW	DOB :05/13/1988	Sex:Female	Race:Asian	Height:157cm	Weight:3050g		
First administration	date of batch :		Batch number :	Batch number :				
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/16/2015		112.5					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/06/2015	06/15/2015		Related	Moderate			
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:)		Not applicable	Concomitant procedure**	Resolved				
Event Description:N 15-Jun-2015,2900m	lausea, vomiting, yellow nl;	urine with less volum	e, chest pelvic effusion	ascites puncture 10-,	lun-2015,2200ml;ascite	s puncture		
Subject received co	ncomitant medications:							
Does the subject ha	ive any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Bilateral fallopian tube obstruction				04/11/2014	Not on treatment/medicatio n	Ongoing		

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	Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-17 End Date:2016-08-18								
Study :EMR700623-541	Investigator :Fei Gon	g	Country of Investigator :China	SiteNo:C01				
Subject No :C01-0068	Subject Initials :X-S	DOB :04/16/1989	Sex:Female	Race:Asian Height:160cm We		Weight:2500g		
First administration d	late of batch :	•	Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	06/27/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	07/23/2015	08/04/2015		Related	Severe			
Causality Factors Action Taken with Study Treatment			Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:)		Not applicable	Concomitant procedure**	Resolved				
	odominal distension, Na 80Jul2015,2500ml; 04A		n less volume, chest pe	elvic effusion , Ascites	puncture 23Jul2015,1	000ml;		
Subject received con	comitant medications:							
Does the subject have	ve any relevant past or	present medical condit	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
The left Hydrosalpinx and adhesions, less patency. Right fallopian tube partially blocked			03/22/2014	Not on treatment/medicatio n	Ongoing			
right fallopian tube resection because of Ectopic pregnancy			UK-Oct-2011	Not on treatment/medicatio n				

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-17 End Date:2016-08-18								
Study :EMR700623-541	Investigator :Fei Gon	g	Country of Investigator :China	SiteNo:C01				
Subject No :C01-0158	Subject Initials :T-Y	DOB :06/17/1983	Sex:Female	Race:Asian	Weight:3300g			
First administration d	late of batch :	•	Batch number :					
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	n 09/16/2015 112.5							
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	10/12/2015	10/12/2015		Related	Moderate			
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:)		Not applicable	Concomitant procedure**	Resolved				
Event Description:Ab	odominal distension, Na	ausea, yellow urine with	h less volume, chest pe	elvic effusion,ascites p	ouncture 12-Oct-2015,2	500ml		
Subject received con	comitant medications:							
Does the subject hav	ve any relevant past or	present medical condit	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
right fallopian tube is less patency,left fallopian tube is obstruction				06/04/2015	Not on treatment/medicatio n	Ongoing		
Endometrial polyps h		08/24/2015	Not on treatment/medicatio n	Ongoing				

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0002	Subject Initials :LLL	DOB :02/12/1987	Sex:Female	Race:Asian	Race:Asian Height:158cm			
First administration	date of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	en 05/08/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/25/2015	06/03/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.	•		•		
Subject received co	ncomitant medications:							
Does the subject ha	ive any relevant past or	present medical cond	itions:Yes					
Condition			Start Date	Related to study condition	Ongoing			
salpingemphrxis,lala	ace peritoneoscope			Uk-Jul-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	SiteNo:C02			
Subject No :C02-0008	Subject Initials :R-H	DOB :02/06/1993	Sex:Female	Race:Asian	an Height:158cm Weight:51.0			
First administration date of batch :			Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	Pen 05/11/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/27/2015	06/05/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.	•				
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
laparoscope hydrotu	ubation		Uk-Unk-2013	Not on treatment/medicatio n				

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-	17 End Date:2016-08-1								
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02					
Subject No :C02-0010	Subject Initials :Y-H DOB :10/18/1985		Sex:Female	Race:Asian	Height:153cm	Weight:47.0kg			
First administration	date of batch :		Batch number :	•	-				
Study Drug	Start Date		Dose	Change in Dose		-			
Gonal-f New Pen Stimulation Treatment	05/11/2015		150						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	05/29/2015	06/02/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.			•			
Subject received con	ncomitant medications:								
Does the subject ha	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02	SiteNo:C02			
Subject No :C02-0011	Subject Initials :P-X	DOB :12/07/1994	Sex:Female	Race:Asian	Height:160cm	Weight:60.0kg		
First administration	date of batch :		Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	n 05/11/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/27/2015	06/03/2015		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:M	Ioderate OHSS occurs	canceled embryo trans	sfer, OHSS improveme	nt				
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
ectopic pregnancy S	Salping ectomy			Uk-Unk-2013	On treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0023	Subject Initials :JYF	DOB :11/10/1981	Sex:Female	Race:Asian	Height:163cm	Weight:70.0kg		
First administration	date of batch :	-	Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	en 05/12/2015		300					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/30/2015	06/05/2015		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:M	Ioderate OHSS occurs of	canceled embryo trans	sfer, OHSS improveme	nt		<u> </u>		
Subject received con	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Hysteroscopy Salpir	ıgemphraxis			Uk-Mar-2015	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-1	17 End Date:2016-08-18	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0024	Subject Initials :HXG	DOB :10/06/1992	Sex:Female	Race:Asian	Height:150cm	Weight:55.0kg			
First administration date of batch :		Batch number :							
Study Drug	Start Date		Dose	Change in Dose		•			
Gonal-f New Pen Stimulation Treatment	05/12/2015		200						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	05/30/2015	06/05/2015		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:Mo	oderate OHSS patients	, improved canceled a	after embryo transfer.			•			
Subject received con	comitant medications:								
Does the subject hav	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-17 End Date:2016-08-18								
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0026	Subject Initials :HCJ	DOB :07/07/1993	Sex:Female	Race:Asian	Height:161cm	Weight:53.0kg		
First administration d	late of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/12/2015		200					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/30/2015	06/05/2015		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:Mo	oderate OHSS occurs of	canceled embryo trans	fer, OHSS improveme	nt				
Subject received con	ncomitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparotomy with left tubal ectopic pregnancy resection				Uk-Unk-2011	Not on treatment/medicatio n			
Under ectopic pregna	ancy laparoscopic cons	ervative surgery		Uk-Unk-2012	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08	-17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02					
Subject No :C02-0028	Subject Initials :XHL	DOB :04/12/1981	Sex:Female	Race:Asian	Height:158cm	Weight:65.0kg			
First administration	date of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose		,			
Gonal-f New Pen Stimulation Treatment	ulation		225						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	05/28/2015	06/03/2015		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 a	after embryo transfer.			•			
Subject received co	oncomitant medications:								
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
Left abdominal ecto	opic pregnancy salpinged	tomy		Uk-Unk-2002	Not on treatment/medicatio n				
Right next laparoscopic tubal ectopic pregnancy surgery				Uk-Unk-2010	Not on treatment/medicatio n				
Conservative treatment of ectopic pregnancy				Uk-Unk-2004	Not on treatment/medicatio n				
Conservative treatr	ment of ectopic pregnanc	у		Uk-Unk-2005	Not on treatment/medicatio n				

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8			-			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	SiteNo:C02			
Subject No :C02-0029	Subject Initials :XLX	DOB :06/08/1981	Sex:Female	Race:Asian Height:158cm V		Weight:70.0kg		
First administration of	date of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	n 05/12/2015		250					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/01/2015	06/07/2015		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:M	loderate OHSS occurs o	canceled embryo trans	sfer, OHSS improveme	nt	•			
Subject received cor	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Pelvic mass laparos drilling, tubal.	copic pelvic surgery stic	cky points, bilateral tu	bal ostomy, left ovarian	Uk-Unk-2013	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-	17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0034	Subject Initials :L-Z DOB :09/04/1981		Sex:Female	Race:Asian	Height:156cm	Weight:53.0kg			
First administration of	date of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose		-			
Gonal-f New Pen Stimulation Treatment	05/13/2015		150						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	05/30/2015	06/05/2015		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:M	oderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt					
Subject received cor	ncomitant medications:								
Does the subject has	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0043	Subject Initials :F-H	DOB :03/13/1983	Sex:Female	Race:Asian Height:164cm		Weight:54.0kg		
First administration d	late of batch :	•	Batch number :					
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/13/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/30/2015	06/05/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.			•		
Subject received con	ncomitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparoscopic tubal ectopic pregnancy at the right side of the window to take embryo			ow to take embryo	Uk-Unk-2005	Not on treatment/medicatio n			
Under the right fallopian tube ectopic pregnancy laparoscopic surgery				Uk-Unk-2007	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0045	Subject Initials :H-T	DOB :09/08/1983	Sex:Female	Race:Asian	Height:158cm	Weight:57.5kg	
First administration date of batch :			Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment			250				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/31/2015	06/05/2015		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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt	_		
Subject received co	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Hysteroscopic polyp	pectomy			Uk-Aug-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				-	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0046	Subject Initials :Q-H DOB :10/07/1985		Sex:Female	Race:Asian	Height:161cm	Weight:50.0kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		-
Gonal-f New Pen Stimulation Treatment	05/13/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	05/28/2015	06/05/2015		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:M	oderate OHSS patients	, improved canceled a	after embryo transfer.			-
Subject received cor	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:No			
Condition				Start Date	Related to study condition	Ongoing

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0049	Subject Initials :HPT	DOB :09/16/1980	Sex:Female	Race:Asian	Height:155cm Weight:50.0			
First administration	First administration date of batch :				•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment			187.5					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/31/2015	06/03/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.	•		•		
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparoscopic surge	ry appendicitis			Uk-Unk-2012	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8			-			
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0051	Subject Initials :YQX	DOB :01/15/1983	Sex:Female	Race:Asian	Height:154cm	Weight:50.0kg		
First administration date of batch :			Batch number :	•	•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	n 05/14/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/31/2015	06/05/2015		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:M	oderate OHSS occurs o	canceled embryo trans	sfer, OHSS improveme	nt		•		
Subject received con	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition			Start Date	Related to study condition	Ongoing			
salpingemphraxis,la	paroscopic operation			Uk-Unk-2012	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1	8				
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0052	Subject Initials :YLL	DOB :02/07/1992	Sex:Female	Race:Asian	Height:165cm	Weight:54.0kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	05/14/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	05/30/2015	06/05/2015		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:M	loderate OHSS patients	s, improved canceled	after embryo transfer.			
Subject received con	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:No			
Condition				Start Date	Related to study condition	Ongoing

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0053	Subject Initials :YND	DOB :04/13/1992	Sex:Female	Race:Asian	:Asian Height:155cm		
First administration date of batch :			Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	/ Pen 05/14/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/30/2015	06/05/2015		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:M	loderate OHSS occurs of	canceled embryo tran	sfer, OHSS improveme	nt	_	•	
Subject received co	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Laparoscopic tubal	colostomy, pelvic surger	ry sticky points.	Uk-Unk-2013	Not on treatment/medicatio n			

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	Non Se	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-7	18			-	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0059	Subject Initials :J-L DOB :12/23/1984		Sex:Female	Race:Asian	Height:155cm	Weight:50.0kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		-
Gonal-f New Pen Stimulation Treatment	05/14/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	05/31/2015	06/05/2015		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:	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt	•	
Subject received co	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:No			
Condition				Start Date	Related to study condition	Ongoing

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0060	Subject Initials :Y-W	DOB :11/10/1981	Sex:Female	Race:Asian	Height:150cm	Weight:45.5kg	
First administration	date of batch :		Batch number :	•			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	tion		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/30/2015	06/03/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.			•	
Subject received co	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Laparoscopic pelvic	sticking points, ovarian	drilling, tubal surgery	to clear.	Uk-Unk-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0061	Subject Initials :T-D	DOB :02/15/1988	Sex:Female	Race:Asian	Height:161cm	Weight:60.0kg	
First administration	First administration date of batch :						
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	v Pen 05/14/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/31/2015	06/05/2015		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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt	_		
Subject received co	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
salpingemphraxis,la	paroscopic operation			Uk-Unk-2014	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0062	Subject Initials :FQR	DOB :07/25/1983	Sex:Female	Race:Asian	Height:162cm	Weight:46.0kg		
First administration	date of batch :		Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	v Pen 05/14/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/30/2015	06/03/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.		•			
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Hysteroscopic tubal	inspection			Uk-May-2014	Not on treatment/medicatio n			

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	Non S	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	-17 End Date:2016-0		0			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0063	Subject Initials DOB :07/25/1981 :HMC		Sex:Female	Race:Asian	Height:162cm	Weight:50.0kg
First administration	date of batch :		Batch number :	Batch number :		
Study Drug	Start Date		Dose	Change in Dose		4
Gonal-f New Pen Stimulation Treatment	05/15/2015		250			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/01/2015	06/05/2015		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:N	Ioderate OHSS occu	rs canceled embryo tran	sfer, OHSS improveme	nt	1	1
Subject received co	oncomitant medication	าร:				
Does the subject ha	ave any relevant past	or present medical cond	litions:Yes			
Condition				Start Date	Related to study condition	Ongoing
Hysteroscopic surgery through liquid				Uk-Unk-2014	Not on treatment/medicatio n	

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-7	17 End Date:2016-08-1	8			•				
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0070	Subject Initials :ZQC	DOB :02/08/1977	Sex:Female	Race:Asian	Height:164cm	Weight:67.0kg			
First administration of	date of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose		-			
Gonal-f New Pen Stimulation Treatment	05/15/2015		200						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	05/28/2015	06/03/2015		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:M	oderate OHSS patients	, improved canceled a	after embryo transfer.						
Subject received cor	ncomitant medications:								
Does the subject hav	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0071	Subject Initials :LLX	Subject Initials :LLX DOB :07/13/1987		Race:Asian	Height:157cm	Weight:41.5kg	
First administration	First administration date of batch :						
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment			150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/31/2015	06/05/2015		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:	Ioderate OHSS occurs	canceled embryo trans	sfer, OHSS improveme	nt	_		
Subject received co	ncomitant medications:						
Does the subject ha	ive any relevant past or	present medical cond	litions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Laparoscopic pelvic	sub sticky, sticky points	s left fallopian tube su	Uk-Unk-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0076	Subject Initials :F-L	DOB :05/15/1987	Sex:Female	Race:Asian	Weight:51.5kg			
First administration d	late of batch :		Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	05/15/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	05/30/2015	06/03/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.			•		
Subject received con	comitant medications:							
Does the subject hav	e any relevant past or	present medical condit	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Sticky points lower pelvic laparoscopy surgery, tubal surgery.				Uk-Unk-2012	Not on treatment/medicatio n			
Hysteroscopic tubal surgery				Uk-Sep-2014	Not on treatment/medicatio n			

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	Non Se	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				•	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0082	Subject Initials :Y-L DOB :09/04/1986		Sex:Female	Race:Asian	Height:162cm	Weight:59.0kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	05/18/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/02/2015	06/05/2015		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:M	oderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt		
Subject received con	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:No			
Condition				Start Date	Related to study condition	Ongoing

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0083	Subject Initials :Y-L	Subject Initials :Y-L DOB :08/20/1988		Race:Asian	Height:158cm	Weight:54.0kg		
First administration of	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	on		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/03/2015	06/05/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.	•		•		
Subject received con	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparoscopic pelvic	sticky points left salping	gostomy		Uk-Unk-2013	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				-	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0086	Subject Initials :YZZ DOB :07/07/1987		Sex:Female	Race:Asian	Height:150cm	Weight:55.0kg
First administration	date of batch :		Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	05/18/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/04/2015	06/12/2015		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:M	oderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt	•	
Subject received con	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:No			
Condition				Start Date	Related to study condition	Ongoing

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0087	Subject Initials :XPT	DOB :06/15/1988	Sex:Female	Race:Asian	Height:158cm	Weight:57.0kg		
First administration	date of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment								
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/02/2015	06/05/2015		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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.	•		•		
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Hysteroscopy norma	al			Uk-Mar-2015	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	-17 End Date:2016-08-1	8				
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0088	Subject Initials :HLK	DOB :06/14/1986	Sex:Female	Race:Asian	Height:159cm	Weight:59.0kg
First administration date of batch :			Batch number :	•		
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	05/18/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/02/2015	06/05/2015		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:N	Ioderate OHSS occurs of	canceled embryo trans	sfer, OHSS improveme	nt	•	
Subject received co	ncomitant medications:					
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes			
Condition				Start Date	Related to study condition	Ongoing
Palace laparoscopic endometriosis lesior	c pelvic surgery sticky po ns fulguration.	oints, bilateral tubal pl	astic surgery,	Uk-Unk-2012	Not on treatment/medicatio n	

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0089	Subject Initials :XYZ	DOB :06/12/1986	Sex:Female	Race:Asian	Height:152cm	Weight:48.5kg		
First administration	date of batch :	•	Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	ation		187.5					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/07/2015	06/12/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.			•		
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Sticky points pelvic	laparoscopy surgery, tu	bal surgery to clear		Uk-Nov-2012	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-1	17 End Date:2016-08-18	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0095	Subject Initials :MLG	DOB :08/19/1990	Sex:Female	Race:Asian	Height:155cm	Weight:54.0kg			
First administration d	late of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose		•			
Gonal-f New Pen Stimulation Treatment	05/19/2015		150						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	06/03/2015	06/05/2015		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:Mo	oderate OHSS occurs o	anceled embryo trans	sfer, OHSS improveme	nt		•			
Subject received con	comitant medications:								
Does the subject hav	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0102	Subject Initials :HYD	DOB :10/14/1985	Sex:Female	Race:Asian	Height:163cm	Weight:57.0kg		
First administration	date of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	/ Pen 05/19/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/04/2015	06/12/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.	•				
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Palace laparoscopy				Uk-Unk-2013	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0103	Subject Initials :CPZ	DOB :05/12/1989	Sex:Female	Race:Asian	Height:165cm	Weight:63.5kg	
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	en 05/19/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/05/2015	06/12/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.	4			
Subject received co	ncomitant medications:						
Does the subject ha	ive any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
The right side of the	open line on the right s	ide of salpingectomy	Uk-Unk-2011	Not on treatment/medicatio n			

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	Non S	Serious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	-17 End Date:2016-0	8-18			•		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0111	Subject Initials DOB :10/10/1979 :CXW		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	06/05/2015	06/12/2015		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:M	Ioderate OHSS patie	ents, improved canceled	after embryo transfer.		1	1	
Subject received co	oncomitant medication	ns:					
Does the subject ha	ave any relevant past	or present medical cond	litions: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/medicatio n		
Tubal lipiodol angiography				Uk-Unk-2013	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-1	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02			
Subject No :C02-0115	Subject Initials :KJT	DOB :07/02/1988	Sex:Female	Race:Asian	Weight:52.0kg		
First administration d	late of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/21/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/05/2015	06/12/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.				
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical condi	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG examination: bilateral tubal occlusion				Uk-Unk-2012	Not on treatment/medicatio n		
Laparoscopy surgery: pelvic sticky points, bilateral tubal ostomy + right side mesosalpinx cyst removal.			right side	Uk-May-2015	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8			•			
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0118	Subject Initials :MLF	DOB :08/03/1984	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	05/21/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/07/2015	06/12/2015		Related	Moderate			
·····		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	anceled embryo trans	sfer, OHSS improveme	nt				
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tuba	l obstruction			Uk-Unk-2007	Not on treatment/medicatio n			
Laparoscopic pelvio	c sticky points, bilateral o	warian drilling		Uk-Unk-2007	Not on treatment/medicatio n			
Cervical biopsy sho	owing inflammation			Uk-Unk-2015	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0124	Subject Initials :Y-C	DOB :07/17/1982	Sex:Female	Race:Asian	Weight:48.5kg			
First administration d	late of batch :		Batch number :	Batch number :				
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	v Pen 05/21/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/07/2015	06/12/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.		_			
Subject received con	comitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: the left fallopian tube obstruction and hydrocephalus right salpingitis			alpingitis	Uk-Unk-2013	Not on treatment/medicatio n			
Hysteroscopy normal				Uk-Mar-2015	Not on treatment/medicatio n			

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	Non S	erious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	-17 End Date:2016-0	8-18			-		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0126	Subject Initials :RCM			Race:Asian	Height:165cm	Weight:60.0kg	
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/21/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/07/2015	06/12/2015		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:N	Ioderate OHSS patie	nts, improved canceled	after embryo transfer.		1	1	
Subject received co	ncomitant medicatio	ns:					
Does the subject ha	ave any relevant past	or present medical cond	litions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: the right fallopian tube obstruction, poor uterine shape				Uk-Unk-2013	Not on treatment/medicatio n		
Hysteroscopy: uterine spindle-shaped, single horn.				Uk-Unk-2014	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8			•			
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	,				
Subject No :C02-0134	Subject Initials :QYL	DOB :06/17/1986	Sex:Female	Race:Asian	Height:155cm	Weight:47.0kg		
First administration	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	05/22/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/06/2015	06/12/2015		Related	Moderate			
		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 of	canceled embryo trans	sfer, OHSS improveme	nt		•		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparotomy: pelvic abscess incision and drainage, the right accessories cystector				Uk-Unk-2009	Not on treatment/medicatio n			
Hysteroscopy normal uterine shape				Uk-Unk-2013	Not on treatment/medicatio n			
Hysteroscopic cure	ttage		Uk-Unk-2015	Not on treatment/medicatio n				

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0136	Subject Initials :Q-L	DOB :03/28/1984	Sex:Female	Sex:Female Race:Asian Height:154cm		Weight:47.0kg	
First administration	date of batch :		Batch number :	•	•		
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/22/2015		200				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/08/2015	06/12/2015		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:N	loderate OHSS patients	, improved canceled a	after embryo transfer.	•			
Subject received co	ncomitant medications:						
Does the subject ha	ive any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal	obstruction incomplete			Uk-Unk-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8			-		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0137	Subject Initials :YLF	DOB :11/01/1989	Sex:Female	Race:Asian	ice:Asian Height:153cm Weigh		
First administration	date of batch :		Batch number :		•		
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	n 05/22/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/09/2015	06/16/2015		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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt			
Subject received co	ncomitant medications:						
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes				
Condition			Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal	HSG: bilateral tubal obstruction				Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0138	Subject Initials :TTJ	DOB :06/20/1990	Sex:Female	Race:Asian Height:150cm		Weight:41.0kg	
First administration of	date of batch :		Batch number :	•			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/22/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/06/2015	06/12/2015		Related	Moderate		
Causality Factors Action Taken with Study Treatment			Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
None(Othervalue:)		Not applicable	None	Resolved			
Event Description:M	oderate OHSS patients	, improved canceled a	after embryo transfer.			•	
Subject received cor	ncomitant medications:						
Does the subject has	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Ectopic pregnancy laparoscopic surgery: the right fallopian tube embryo pelvic sticking points				Uk-Jan-2014	Not on treatment/medicatio n		
Hysteroscopy: endometrial polyps.				Uk-Feb-2015	Not on treatment/medicatio n		

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	Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-1	7 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0142	Subject Initials :HMT	DOB :02/19/1988	Sex:Female	Race:Asian Height:168cm		Weight:46.0kg		
First administration d	late of batch :		Batch number :	Batch number :				
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	Pen 05/22/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/07/2015	06/12/2015		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:Mo	oderate OHSS occurs o	anceled embryo trans	fer, OHSS improveme	nt	_	•		
Subject received con	comitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal obstruction and hydrocephalus			Uk-Unk-2014	Not on treatment/medicatio n				
Laparoscopy surgery: pelvic stars stick + fulguration of endometriosis foci, tubal plastic surgery			Uk-Unk-2014	Not on treatment/medicatio n				

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	Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02	SiteNo:C02			
Subject No :C02-0143	Subject Initials :F-L	DOB :11/06/1982	Sex:Female	Race:Asian Height:155cm		Weight:51.0kg		
First administration d	late of batch :		Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	en 05/22/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/08/2015	06/14/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.			•		
Subject received con	comitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: the left fallopian tube obstruction, incomplete right fallopian tube obstruction			tube obstruction	Uk-Unk-2012	Not on treatment/medicatio n			
Laparoscopy surgery: pelvic stars stick + tubal surgery, endometriosis lesions fulguration			Uk-Unk-2012	Not on treatment/medicatio n				

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				•	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0153	Subject Initials :H-H DOB :07/12/1979		Sex:Female	Race:Asian	Height:153cm	Weight:43.5kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	05/25/2015		250			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/10/2015	06/16/2015		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:M	oderate OHSS patients	, improved canceled a	after embryo transfer.		•	
Subject received cor	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	itions:No			
Condition				Start Date	Related to study condition	Ongoing

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	,				
Subject No :C02-0157	Subject Initials :Y-Z	DOB :07/22/1984	Sex:Female	Race:Asian	Height:156cm	Weight:55.0kg		
First administration	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	05/25/2015		200					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/10/2015	06/16/2015		Related	Moderate			
		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 trans	sfer, OHSS improveme	nt		•		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal	lobstruction			Uk-Unk-2006	Not on treatment/medicatio n			
Laparoscopic Surge	ery: Tubal clear.			Uk-Unk-2007	Not on treatment/medicatio n			
Ectopic pregnancy	laparoscopic surgery: tu	bal embryo window.		Uk-Jan-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8			•			
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	China SiteNo:C02				
Subject No :C02-0161	Subject Initials :H-Y	tials :H-Y DOB :12/12/1985 Sex:Female Race:Asi			Height:166cm	Weight:57.0kg		
First administration	date of batch :		Batch number :		•			
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	05/26/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/10/2015	06/16/2015		Related	Moderate			
		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:	Noderate OHSS patients	, improved canceled a	after embryo transfer.	-		•		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal obstruction				Uk-Unk-2007	Not on treatment/medicatio n			
Laparoscopic tubal	surgery sticking points.			Uk-Unk-2007	Not on treatment/medicatio n			
Open left fallopian t	ube ectopic pregnancy	surgery		Uk-Unk-2012	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-1	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02			
Subject No :C02-0164	Subject Initials :AHC	DOB :12/12/1982	Sex:Female	Race:Asian	Weight:49.0kg		
First administration d	late of batch :		Batch number :	Batch number :			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/26/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/12/2015	06/20/2015		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:Mo	oderate OHSS occurs o	anceled embryo trans	fer, OHSS improveme	nt			
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical condi	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal occlusion side				Uk-Unk-2012	Not on treatment/medicatio n		
Laparoscopy Surgery: salpingostomy, pelvic adhesions dissection.				Uk-Unk-2012	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				-	
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0169	Subject Initials :L-W	DOB :06/08/1985	Sex:Female	Race:Asian Height:166cm We		Weight:50.0kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change in Dose		-
Gonal-f New Pen Stimulation Treatment	05/27/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/12/2015	06/20/2015		Related	Moderate	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)		Not applicable	None	Resolved		
Event Description:M	loderate OHSS occurs of	canceled embryo tran	sfer, OHSS improveme	nt		•
Subject received co	ncomitant medications:					
Does the subject ha	ive any relevant past or	present medical cond	litions:Yes			
Condition				Start Date	Related to study condition	Ongoing
HSG: bilateral tubal	occlusion insufficiency			Uk-Jun-2014	Not on treatment/medicatio n	

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China					
Subject No :C02-0174	Subject Initials :YXC	Subject Initials :YXC DOB :06/04/1982		Race:Asian	Height:155cm	Weight:54.0kg		
First administration	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/27/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/12/2015	06/16/2015		Related	Moderate			
		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:	Ioderate OHSS occurs of	anceled embryo trans	sfer, OHSS improveme	nt				
Subject received co	ncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Laparoscopic surgery: left ovarian endometriosis cystectomy				Uk-Unk-2010	Not on treatment/medicatio n			
HSG: the left fallopi	an tube obstruction, righ	mation.	Uk-Unk-2012	Not on treatment/medicatio n				
Laparoscopic surge	ry: pelvic adhesions dis	section, left fallopian to	ube plastic surgery	Uk-Unk-2012	Not on treatment/medicatio n			

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	Non Serious Adverse Drug Reactions Report							
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0175	Subject Initials :L-T	DOB :02/15/1989	Sex:Female	Race:Asian	Weight:40.0kg			
First administration d	late of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	05/27/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/11/2015	06/16/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.					
Subject received con	comitant medications:							
Does the subject hav	e any relevant past or	present medical condit	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal obstruction.			Uk-Aug-2014	Not on treatment/medicatio n				
Laparoscopy surgery: pelvic adhesions dissection, tubal plastic surgery to repair the left corpus luteum cyst cystectomy,				Uk-Unk-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0176	Subject Initials :KBF	DOB :05/19/1986	Sex:Female	Race:Asian	Height:160cm	Weight:57.0kg		
First administration date of batch :			Batch number :	4				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment			200					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/11/2015	06/17/2015		Related	Moderate			
Causality Factors Action Taken with Study Treatment			Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:) Not applicable			None	Resolved				
Event Description:	OHSS occurs not put em	bryos				-		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	litions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Tubal examination:	passable			Uk-Unk-2008	Not on treatment/medicatio n			
HSG: bilateral tuba	l occlusion		Uk-Unk-2009	Not on treatment/medicatio n				
Laparoscopy surgery: bilateral tubal repair plastic surgery, pelvic adhesions dissection				Uk-Unk-2011	Not on treatment/medicatio n			
Ectopic pregnancy:	open surgery			Uk-Unk-2012	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8			•			
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	· ·				
Subject No :C02-0178	Subject Initials :XSC	Subject Initials :XSC DOB :10/11/1988		Race:Asian	Height:155cm	Weight:54.0kg		
First administration	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/27/2015		200					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/10/2015	06/16/2015		Related	Moderate			
		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 a	after embryo transfer.			•		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tuba	l resistance.			Uk-Jan-2013	Not on treatment/medicatio n			
Laparoscopy surge surgery, normal ute	ry: pelvic adhesions diss rine shape.	ection, bilateral salpir	igo-repair plastic	Uk-Jan-2013	Not on treatment/medicatio n			
Hysteroscopy norm	al, water surgery: incom	plete right fallopian tu	be obstruction	Uk-Feb-2015	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0179	Subject Initials :HYZ	DOB :05/14/1991	Sex:Female	Race:Asian	Height:158cm	Weight:53.0kg		
First administration	date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		-		
Gonal-f New Pen Stimulation Treatment	05/27/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/11/2015	06/16/2015		Related	Moderate			
		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 of	canceled embryo trans	sfer, OHSS improveme	nt		•		
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Ectopic pregnancy	laparotomy: the left fallo	pian tube removal.		Uk-Unk-2009	Not on treatment/medicatio n			
Abdominal ectopic	pregnancy conservative	surgery.		Uk-Unk-2010	Not on treatment/medicatio n			
Hysteroscopy: endo	ometrial polyp excision.			Uk-Jan-2015	Not on treatment/medicatio n			

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	Non S	erious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	-17 End Date:2016-0	8-18			-		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0180	Subject Initials :YPW			Race:Asian	Height:156cm	Weight:57.0kg	
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/27/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/13/2015	06/16/2015		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:	Ioderate OHSS patie	nts, improved canceled a	after embryo transfer.		1	1	
Subject received co	ncomitant medicatio	ns:					
Does the subject ha	ave any relevant past	or present medical cond	litions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Ectopic pregnancy: treatment of chemical drugs to kill embryos				Uk-Unk-2012	Not on treatment/medicatio n		
HSG: bilateral tubal	occlusion			Uk-Unk-2008	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8			-		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	SiteNo:C02		
Subject No :C02-0185	Subject Initials :YMX	DOB :09/07/1975	Sex:Female	Race:Asian	ace:Asian Height:159cm		
First administration	date of batch :	•	Batch number :	•			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	n 05/28/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/15/2015	06/20/2015		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:N	Ioderate OHSS occurs of	canceled embryo tran	sfer, OHSS improveme	nt			
Subject received co	ncomitant medications:						
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal	occlusion			Uk-Unk-2013	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1				•	
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0190	Subject Initials :F-H	DOB :03/09/1984	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	05/28/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	06/12/2015	06/16/2015		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:M	oderate OHSS patients	, improved canceled a	after embryo transfer.		_	
Subject received con	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	itions:No			
Condition				Start Date	Related to study condition	Ongoing

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-1	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02	SiteNo:C02			
Subject No :C02-0191	Subject Initials :SYZ	DOB :02/16/1983	Sex:Female	Race:Asian	Weight:58.0kg			
First administration d	late of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		•		
Gonal-f New Pen Stimulation Treatment	05/28/2015		200					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	06/14/2015	06/18/2015		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:Mo	oderate OHSS occurs of	canceled embryo trans	fer, OHSS improveme	nt		•		
Subject received con	ncomitant medications:							
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal occlusion with effusion				Uk-Unk-2008	Not on treatment/medicatio n			
Tubal treatment: Tips patency				Uk-Unk-2008	Not on treatment/medicatio n			

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	Non Se	erious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-7	18					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0195	Subject Initials :J-L	DOB :08/15/1984	Sex:Female	Race:Asian	Weight:48.0kg		
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	Pen 05/28/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/14/2015	06/18/2015		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:	Ioderate OHSS patients	s, improved canceled a	after embryo transfer.				
Subject received co	ncomitant medications:						
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes				
Condition			Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal	patency, pelvic adhesi	ons.		Uk-Unk-2012	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8			-		
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0200	Subject Initials :L-Z	DOB :12/17/1988	Sex:Female	Race:Asian	Height:155cm	Weight:70.0kg	
First administration	date of batch :	-	Batch number :		•		
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	en 05/28/2015		187.5				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	06/13/2015	06/18/2015		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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt			
Subject received co	ncomitant medications:						
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes				
Condition			Start Date	Related to study condition	Ongoing		
HSG: bilateral obstruction				Uk-Apr-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	SiteNo:C02		
Subject No :C02-0201	Subject Initials :J-L	DOB :12/15/1990	DOB :12/15/1990 Sex:Female Race:Asian Height:168cm			Weight:65kg	
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	en 09/07/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/26/2015	10/02/2015		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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt	_		
Subject received co	ncomitant medications:						
Does the subject ha	ive any relevant past or	present medical cond	litions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal	HSG: bilateral tubal occlusion				Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	SiteNo:C02			
Subject No :C02-0202	Subject Initials :F-Q	DOB :06/09/1985	Sex:Female	Race:Asian	Weight:65.0kg			
First administration	date of batch :		Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose		,		
Gonal-f New Pen Stimulation Treatment	08/27/2015		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	09/11/2015	09/16/2015		Related	Moderate			
		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 a	after embryo transfer.	•				
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal patency				Uk-Unk-2013	Not on treatment/medicatio n			
Laparoscopic surge	ery: pelvic adhesions dis	section, bilateral meso	osalpinx cystectomy	Uk-Unk-2014	Not on treatment/medicatio n			
Tubal examination:	smooth			Uk-Unk-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08	-17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0206	Subject Initials :YPZ	DOB :10/20/1980	Sex:Female	Race:Asian	Height:150cm	Weight:47.0kg			
First administration	date of batch :		Batch number :		•				
Study Drug	Start Date		Dose	Change in Dose					
Gonal-f New Pen Stimulation Treatment	nulation		187.5						
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:	Moderate OHSS occurs of	anceled embryo trans	sfer, OHSS improveme	nt	•				
Subject received co	oncomitant medications:								
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
HSG: bilateral tuba	l occlusion			Uk-Unk-2011	Not on treatment/medicatio n				
Laparoscopy surgery: pelvic adhesions dissection, bilateral tubal surgery to clear.				Uk-Unk-2011	Not on treatment/medicatio n				
HSG: bilateral tubal occlusion				Uk-Unk-2013	Not on treatment/medicatio n				
	Laparoscopy surgery: pelvic adhesions dissection, uterine fibroids dug surgery, tubal surgery to clear, coherent liquid skill.				Not on treatment/medicatio n				

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8			•			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0207	Subject Initials :JHZ	DOB :09/03/1984	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		150					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	09/13/2015	09/18/2015		Related	Moderate			
		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 a	after embryo transfer.	•				
Subject received co	oncomitant medications:							
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Ectopic pregnancy	laparoscopic surgery: Le	ft salpingectomy		Uk-Unk-2011	Not on treatment/medicatio n			
Ectopic pregnancy	laparoscopic surgery: Ri	ght salpingectomy		Uk-Unk-2013	Not on treatment/medicatio n			
HSG: bilateral tuba	l occlusion			Uk-Unk-2014	Not on treatment/medicatio n			

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	Non S	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-0	8-18	Ŭ			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0208	Subject Initials :MQW	DOB :09/24/1988	Sex:Female	Race:Asian	Height:156cm	Weight:54.0kg
First administration	date of batch :		Batch number :		•	
Study Drug	Start Date	Start Date		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:N	loderate OHSS occu	rs canceled embryo tran	sfer, OHSS improveme	nt	•	
Subject received co	ncomitant medication	าร:				
Does the subject ha	ive any relevant past	or present medical cond	litions:Yes			
Condition				Start Date	Related to study condition	Ongoing
Hysteroscopic resection of endometrial polyps				Uk-Unk-2015	Not on treatment/medicatio n	

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	Non S	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	-17 End Date:2016-0		0			
Study :EMR700623-541	Investigator :NA	Investigator :NA		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	Start Date		Change in Dose		4
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:	Ioderate OHSS patie	nts, improved canceled	after embryo transfer.	•		4
Subject received co	ncomitant medication	าร:				
Does the subject ha	ave any relevant past	or present medical cond	litions:Yes			
Condition				Start Date	Related to study condition	Ongoing
HSG: the right fallo	pian tube obstruction			Uk-Unk-2014	Not on treatment/medicatio n	

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-17 End Date:2016-08-18								
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	Weight:56.0kg			
First administration d	late 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:Mo	oderate OHSS occurs of	canceled embryo trans	fer, OHSS improveme	nt		•		
Subject received con	comitant medications:							
Does the subject hav	ve any relevant past or	present medical condit	tions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal occlusion				Uk-Unk-2014	Not on treatment/medicatio n			
Under laparoscopy surgery to clear the fallopian tubes, pelvic adhesions			esions dissection	Uk-Unk-2014	Not on treatment/medicatio n			

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8			-			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02	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			
		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:M	loderate OHSS occurs of	anceled embryo trans	sfer, OHSS improvemer	nt		•		
Subject received con	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
Ectopic pregnancy therapy chemotherapy to kill embryos				Uk-Unk-2011	Not on treatment/medicatio n			
HSG: bilateral tubal obstruction.				Uk-Unk-2010	Not on treatment/medicatio n			
Laparoscopy surger	y: pelvic adhesions diss	ection, bilateral tubal	surgery to clear	Uk-Unk-2010	Not on treatment/medicatio n			

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	Non Se	rious Adve	erse Drug	Reactions	s Report		
Start Date:2016-08-1	17 End Date:2016-08-1	8					
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:Mo	oderate OHSS occurs o	anceled embryo trans	fer, OHSS improveme	nt		•	
Subject received con	comitant medications:						
Does the subject hav	ve any relevant past or	present medical condit	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal o	obstruction.			Uk-Unk-2011	Not on treatment/medicatio n		
HSG: bilateral tubal p	patency, pelvic adhesic	ns.		Uk-Unk-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report			
Start Date:2016-08-	17 End Date:2016-08-1	8						
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	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:M	loderate OHSS patients	, improved canceled a	after embryo transfer.		_	•		
Subject received co	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: bilateral tubal	patency.			Uk-Unk-2014	Not on treatment/medicatio n			

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	Non S	erious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	-17 End Date:2016-0	8-18					
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:M	/loderate OHSS occu	rs canceled embryo tran	sfer, OHSS improveme	ent		1	
Subject received co	oncomitant medicatio	ns:					
Does the subject ha	ave any relevant past	or present medical cond	litions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: the side of tubal passable.				Uk-Unk-2012	Not on treatment/medicatio n		
Laparoscopic surge	ery: bilateral tubal sur	gery, right fallopian tube	patency.	Uk-Unk-2012	Not on treatment/medicatio n		

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	Non Se	rious Adve	erse Drug	Reactions	s Report		
Start Date:2016-08-1	7 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02			
Subject No :C02-0224	Subject Initials :JXL	DOB :11/11/1985	Sex:Female	Race:Asian	Height:158cm	Weight:52.0kg	
First administration date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	09/01/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/19/2015	09/25/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.			•	
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical condit	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal o	occlusion in the left effu	usion		Uk-Unk-2014	Not on treatment/medicatio n		
	adhesions dissection a ion of endometrial poly		plastic ostomy,	Uk-Unk-2014	Not on treatment/medicatio n		

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	Non Se	rious Adve	erse Drug	Reactions	s Report		
Start Date:2016-08-1	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02			
Subject No :C02-0226	Subject Initials :BQX	DOB :07/10/1987	Sex:Female	Race:Asian	Height:160cm	Weight:53.0kg	
First administration date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose		,	
Gonal-f New Pen Stimulation Treatment	09/01/2015		187.5				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/17/2015	09/22/2015		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:Mo	oderate OHSS occurs o	anceled embryo trans	fer, OHSS improveme	nt		•	
Subject received con	comitant medications:						
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Laparoscopic surgery classification pelvic adhesions, tubal fluid p		rosthetics	Uk-Unk-2010	Not on treatment/medicatio n			
Laparoscopic surger	y classification pelvic a	dhesions, tubal fluid pr	rosthetics	Uk-Unk-2011	Not on treatment/medicatio n		

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	Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0227	Subject Initials :YXZ	DOB :12/09/1985	Sex:Female	Race:Asian	Height:164cm	Weight:56.0kg			
First administration	First administration date of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose		,			
Gonal-f New Pen Stimulation Treatment	Stimulation		187.5						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	09/17/2015	09/22/2015		Related	Moderate				
		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:N	Moderate OHSS patients	, improved canceled a	after embryo transfer.	_		•			
Subject received co	oncomitant medications:								
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
Abdominal ectopic pregnancy conservative surgery.				Uk-Unk-2012	Not on treatment/medicatio n				
Ectopic pregnancy laparoscopic conservative surgery.				Uk-Unk-2013	Not on treatment/medicatio n				
HSG: incomplete rig	ght fallopian tube obstru	ction, left fallopian tub	e fluid.	Uk-Unk-2014	Not on treatment/medicatio n				

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	Non Se	rious Adv	erse Drug	Reactions	s Report			
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02				
Subject No :C02-0228	Subject Initials :J-X	Subject Initials :J-X DOB :09/12/1981		Race:Asian	Height:160cm	Weight:51.0kg		
First administration date of batch :		Batch number :						
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	09/02/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	09/20/2015	09/25/2015		Related	Moderate	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:N	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt				
Subject received co	ncomitant medications:							
Does the subject ha	ive any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: right fallopian	tube obstruction.			Uk-Unk-2013	Not on treatment/medicatio n			

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	Non Serious Adverse Drug Reactions Report								
Start Date:2016-08	-17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0229	Subject Initials :HYT	DOB :11/13/1992	Sex:Female	Race:Asian	Height:153cm	Weight:45.5kg			
First administration	First administration date of batch :		Batch number :		-				
Study Drug	Start Date		Dose	Change in Dose		•			
Gonal-f New Pen 09/02/2015 Stimulation Treatment		150							
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	09/18/2015	09/24/2015		Related	Moderate				
		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:	Noderate OHSS patients	, improved canceled a	after embryo transfer.	4	-	•			
Subject received co	oncomitant medications:								
Does the subject ha	ave any relevant past or	present medical cond	litions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
HSG: bilateral tubal occlusion				Uk-Unk-2013	Not on treatment/medicatio n				
Laparoscopic surgery: pelvic adhesions dissection, tubal plasty, u surgery, uterine suspension surgery.			uterine fibroids dug	Uk-Unk-2013	Not on treatment/medicatio n				
Hysteroscopy norm	nal			Uk-Unk-2015	Not on treatment/medicatio n				

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	Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	-17 End Date:2016-08-1	8			•				
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02					
Subject No :C02-0230	Subject Initials :F-X	DOB :07/08/1987	Sex:Female	Race:Asian	Height:156cm	Weight:51.0kg			
First administration date of batch :		Batch number :							
Study Drug	Start Date		Dose	Change in Dose		,			
Gonal-f New Pen 09/02/2015 Stimulation Treatment			150						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	09/18/2015	09/24/2015		Related	Moderate				
		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:M	Ioderate OHSS occurs	canceled embryo trans	sfer, OHSS improveme	nt		•			
Subject received co	ncomitant medications:								
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
Caesarean section				Uk-Unk-2009	Not on treatment/medicatio n				
HSG: bilateral tubal	occlusion			Uk-Mar-2015	Not on treatment/medicatio n				
Hysteroscopy norm	al			Uk-Unk-2015	Not on treatment/medicatio n				

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	Non Se	rious Adve	erse Drug	Reactions	s Report		
Start Date:2016-08-1	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02			
Subject No :C02-0231	Subject Initials :LJS	DOB :08/28/1981	Sex:Female	Race:Asian	Height:160cm	Weight:69.0kg	
First administration date of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	imulation		300				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/17/2015	09/24/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.				
Subject received con	ncomitant medications:						
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: the left fallopia	in tube passable			Uk-Unk-2013	Not on treatment/medicatio n		
Palace laparoscopy				Uk-Unk-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1	8				
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C02		
Subject No :C02-0239	Subject Initials :JJL	Subject Initials :JJL DOB :01/25/1984		Race:Asian	Height:158cm	Weight:49.0kg
First administration date of batch :		Batch number :				
Study Drug Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	09/06/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/23/2015	09/27/2015		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:M	Ioderate OHSS occurs	canceled embryo tran	sfer, OHSS improveme	nt		•
Subject received co	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	litions:Yes			
Condition				Start Date	Related to study condition	Ongoing
HSG: incomplete rig	ht fallopian tube obstru	ction, left fallopian tub	e obstruction	Uk-Mar-2015	Not on treatment/medicatio n	

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	Non S	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08					
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0240	Subject Initials :DMX			Race:Asian	Height:155cm	Weight:47.0kg
First administration of	date of batch :		Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	09/06/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/21/2015	09/26/2015		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:M	oderate OHSS patie	nts, improved canceled a	after embryo transfer.	•		
Subject received cor	ncomitant medicatior	IS:				
Does the subject ha	ve any relevant past	or present medical cond	litions:Yes			
Condition			Start Date	Related to study condition	Ongoing	
Laparoscopy surger fallopian tubes, uteri		eparation surgery, plasti	ic surgery to repair the	Uk-Unk-2013	Not on treatment/medicatio n	

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	Non S	erious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-0		0			
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02		
Subject No :C02-0244	Subject Initials DOB :09/13/1986 Set :DDW		Sex:Female	Race:Asian	Height:160cm	Weight:53.0kg
First administration	date of batch :		Batch number :	Batch number :		
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	09/06/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/22/2015	09/27/2015		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:N	loderate OHSS occu	rs canceled embryo tran	sfer, OHSS improveme	nt		
Subject received co	ncomitant medication	ns:				
Does the subject ha	ive any relevant past	or present medical conc	litions:Yes			
Condition			Start Date	Related to study condition	Ongoing	
HSG: bilateral tubal patency				Uk-Unk-2008	Not on treatment/medicatio n	

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	Non Se	rious Adve	erse Drug	Reactions	s Report		
Start Date:2016-08-1	17 End Date:2016-08-18	8	-				
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02			
Subject No :C02-0247	Subject Initials :BSH	DOB :08/12/1987	Sex:Female	Race:Asian Height:155cm		Weight:40.0kg	
First administration of	date of batch :		Batch number :	Batch number :			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	09/07/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/23/2015	09/27/2015		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:Mo	oderate OHSS patients	, improved canceled a	fter embryo transfer.		_	•	
Subject received cor	ncomitant medications:						
Does the subject have	ve any relevant past or	present medical condi	tions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
HSG: the right fallopian tube obstruction, incomplete left fallopian tube obstruction			Uk-Mar-2014	Not on treatment/medicatio n			
Laparoscopy surgery was surgery.	y: dissection of pelvic a	dhesions, tubal repair	plastic surgery, and it	Uk-Apr-2014	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report			
Start Date:2016-08-	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :NA	Investigator :NA		SiteNo:C02				
Subject No :C02-0249	Subject Initials : ZPY	DOB :09/13/1982	Sex:Female	ex:Female Race:Asian Height:162cm		Weight:55.0kg		
First administration date of batch :			Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	09/07/2015		187.5					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	09/22/2015	09/28/2015		Related	Moderate	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:M	oderate OHSS patients	, improved canceled a	after embryo transfer.	4		•		
Subject received con	ncomitant medications:							
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes					
Condition				Start Date	Related to study condition	Ongoing		
HSG: incomplete rig	ht fallopian tube obstru	ction, left fallopian tub	Uk-Unk-2015	Not on treatment/medicatio n				

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-7	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C04			
Subject No :C04-0087	Subject Initials :HZD	DOB :08/04/1983	Sex:Female	Race:Asian	Weight:60kg		
First administration of	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		<u>.</u>	
Gonal-f New Pen Stimulation Treatment	06/26/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	07/07/2015	07/21/2015		Related	Mild		
Causality Factors Action Taken with Study Treatment		Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:)		Dose reduced	None	Resolved			
Event Description:no	one					•	
Subject received cor	ncomitant medications:						
Does the subject hav	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
spontaneous abortion				01/05/2010	Not on treatment/medicatio n		
spontaneous abortio	on			05/18/2012	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08-	17 End Date:2016-08-1	8				
Study :EMR700623-541	Investigator :NA		Country of Investigator :China	SiteNo:C04		
Subject No :C04-0171	Subject Initials :SHL	DOB :08/17/1987	Sex:Female	Race:Asian	Height:154cm	Weight:40kg
First administration date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		-
Gonal-f New Pen Stimulation Treatment	09/29/2015		300			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	09/30/2015	10/09/2015		Related	Mild	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)		Dose reduced	None	Resolved		
Event Description:no	one	•	-		_	•
Subject received cor	ncomitant medications:					
Does the subject ha	ve any relevant past or	present medical cond	itions:No			
Condition				Start Date	Related to study condition	Ongoing

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	Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-	17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :Ying Zho	ong	Country of Investigator :China	SiteNo:C05					
Subject No :C05-0001	Subject Initials :XFZ	DOB :01/21/1988	Sex:Female	Race:Asian	h 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		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity			
None(Othervalue:)		Not applicable	None	Resolved					
Event Description:H ivgtt qd	lydroxyethyl Starch 130/	0.4 and Sodium Chlor	ide Injection 500ml,iv	gtt bid calcium glucon	ate injection 10ml+dext	rose injection 500ml			
Subject received co	ncomitant medications:								
Does the subject ha	ive any relevant past or	present medical condi	itions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
oophorocystectomy				UK-Jul-2013	Not on treatment/medicatio n				
salpingoplasty				UK-Feb-2014	Not on treatment/medicatio n				
salpingitis after prev	vious tubal occlusion			UK-Feb-2014	Not on treatment/medicatio n	Ongoing			

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	Non Se	rious Adv	erse Drug	Reactions	Report		
Start Date:2016-08-7	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator : Ying Zhong		Country of Investigator :China	SiteNo:C05	SiteNo:C05		
Subject No :C05-0010	Subject Initials :Q-Z	DOB :01/31/1990	Sex:Female	Race:Asian	Height:159cm	Weight:45kg	
First administration date of batch :		Batch number :	Batch number :				
Study Drug	Start Date		Dose	Change in Dose		-	
Gonal-f New Pen Stimulation Treatment	08/02/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	08/17/2015	08/24/2015		Related	Moderate		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
Disease under study procedure**,Concom medication**(Otherva	nitant	Dose not changed	Led to study termination	Resolved			
	odominal distention;nau ose injection 500ml ivg			Sodium Chloride Injec	tion 500ml,ivgtt bid c	alcium gluconate	
Subject received cor	ncomitant medications:						
Does the subject have	ve any relevant past or	present medical cond	itions:No				
Condition				Start Date	Related to study condition	Ongoing	

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	Non Se	rious Adv	erse Drug	Reactions	s Report	
Start Date:2016-08	-17 End Date:2016-08-1					
Study :EMR700623-541	Investigator :Ying Zh	ong	Country of Investigator :China	SiteNo:C05		
Subject No :C05-0013	Subject Initials :L-L	DOB :06/12/1986	Sex:Female	Race:Asian	Height:159cm	Weight:54.5kg
First administration	date of batch :		Batch number :			
Study Drug	Start Date		Dose	Change in Dose		-
Gonal-f New Pen Stimulation Treatment	08/02/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	SS 08/18/2015 08/20/2015			Related	Moderate	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
Disease under study**,Protocol procedure**,Concomitant medication**(Othervalue:)		Dose not changed	Led to study termination	Resolved		
	abdominal distention;nau 10ml+dextrose injection			0.4 and Sodium Chlo	ride Injection 500ml , iv	gtt bid calcium
Subject received co	oncomitant medications:					
Does the subject ha	ave any relevant past or	present medical cond	itions:Yes			
Condition				Start Date	Related to study condition	Ongoing
ampullary pregnand	су			UK-Unk-2010	Not on treatment/medicatio n	
ampullary pregnand	су			UK-Unk-2012	Not on treatment/medicatio n	
ampullary pregnand	су			UK-Unk-2013	Not on treatment/medicatio n	
salpingocatheterisn	n			UK-Unk-2013	Not on treatment/medicatio n	

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	Non Se	rious Adv	erse Drug	Reactions	Report		
Start Date:2016-08-1	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator : Ying Zho	ong	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 d	late 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** medication**(Otherva		Not applicable	None	Resolved			
	dominal distention;Dru ion 500ml ivgtt qd 19-A			Chloride Injection 500	ml,ivgtt bid calcium g	luconate injection	
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical condi	itions:Yes				
Condition			Start Date	Related to study condition	Ongoing		
salpingocatheterism				UK-Unk-2012	Not on treatment/medicatio n		

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	Non Se	rious Adv	erse Drug	Reactions	Report	
Start Date:2016-08-7	17 End Date:2016-08-1	8				
Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05		
Subject No :C05-0041	Subject Initials :W-Y	DOB :07/03/1981	Sex:Female	Race:Asian	Height:153cm	Weight:55kg
First administration of	date of batch :		Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	08/08/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	08/23/2015	08/25/2015		Related	Mild	
Causality Factors	•	Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
Protocol procedure** medication**(Otherv	,	Dose not changed	Led to study termination	Resolved		
	odominal distention;Dru tion 500ml ivgtt qd 21-A			Chloride Injection 500	ml, ivgtt bid calcium g	luconate injection
Subject received cor	ncomitant medications:					
Does the subject have	ve any relevant past or	present medical condi	tions:No			
Condition				Start Date	Related to study condition	Ongoing

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05			
Subject No :C05-0056	Subject Initials :X-P	DOB :06/14/1983	Sex:Female	ale Race:Asian Height:158cm		Weight:46kg	
First administration	date of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose		_	
Gonal-f New Pen Stimulation Treatment	08/13/2015		150				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	08/27/2015	08/29/2015		Related	Mild		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
Protocol procedure* medication**(Otherv	,	Not applicable	Led to study termination	Resolved			
	odominal distention;nau I0ml+dextrose injection	, , 0	, , ,	/0.4 and Sodium Chlo	ride Injection 500ml , iv	gtt bid calcium	
Subject received con	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:No				
Condition				Start Date	Related to study condition	Ongoing	

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	Non Se	rious Adv	erse Drug	Reactions	Report		
Start Date:2016-08-7	17 End Date:2016-08-1	8	-				
Study :EMR700623-541	Investigator : Ying Zho	ong	Country of Investigator :China	SiteNo:C05			
Subject No :C05-0068	Subject Initials :YPL	DOB :12/07/1985	Sex:Female	Race:Asian	Height:160cm Weight:50kg		
First administration of	date of batch :	•	Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/15/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	08/27/2015	08/29/2015		Related	Moderate		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
Protocol procedure** medication**(Otherv		Dose not changed	Led to study termination	Resolved			
	odominal distention;nau 0ml+dextrose injection			0.4 and Sodium Chlo	ide Injection 500ml,iv	gtt bid calcium	
Subject received cor	ncomitant medications:						
Does the subject hav	ve any relevant past or	present medical condi	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Fallopian tube repair	r anaplasty			UK-Unk-2010	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-7	17 End Date:2016-08-1	8						
Study :EMR700623-541	Investigator :Ying Zh	Investigator :Ying Zhong		SiteNo:C05	SiteNo:C05			
Subject No :C05-0117	Subject Initials :Y-L	DOB :04/16/1982	Sex:Female	Race:Asian	Weight:48kg			
First administration of	date of batch :	-	Batch number :	•				
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	08/23/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	09/08/2015	09/10/2015		Related	Mild			
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
Protocol procedure** medication**(Otherv	,	Dose reduced	Led to study termination	Resolved				
	odominal distention;asc ose injection 500ml ivg		-	Sodium Chloride Injec	tion 500ml,ivgtt bid ca	alcium gluconate		
Subject received cor	ncomitant medications:							
Does the subject have	Does the subject have any relevant past or present medical conditions:No							
Condition				Start Date	Related to study condition	Ongoing		

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-	17 End Date:2016-08-1	8					
Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05	SiteNo:C05		
Subject No :C05-0132	Subject Initials :HQL	DOB :07/16/1980	Sex:Female	Race:Asian Height:166cm Weight:57kg			
First administration	First administration date of batch :			•			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	08/27/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	09/08/2015	09/16/2015		Related	Mild		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
Disease under study	y**(Othervalue:)	Dose not changed	None	Resolved			
Event Description:			-			•	
Subject received co	ncomitant medications:						
Does the subject ha	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
Fallopian tube repair anaplasty				UK-Unk-2011	Not on treatment/medicatio n		

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-17 End Date:2016-08-18									
Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo:C05					
Subject No :C05-0141	Subject Initials :XHZ	DOB :04/14/1987	Sex:Female	Race:Asian	Weight:48kg				
First administration of	date of batch :		Batch number :	•					
Study Drug	Start Date		Dose	Change in Dose					
Gonal-f New Pen Stimulation Treatment	08/29/2015		150						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	09/12/2015	09/14/2015		Related	Moderate				
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity			
Protocol procedure** medication**(Otherva	,	Dose not changed	Led to study termination	Resolved					
	odominal distention;nau 0ml+dextrose injection			0.4 and Sodium Chlor	ide Injection 500ml,iv	gtt bid calcium			
Subject received cor	ncomitant medications:								
Does the subject hav	ve any relevant past or	present medical condi	tions:Yes						
Condition				Start Date	Related to study condition	Ongoing			
Fallopian tube repair	r anaplasty			UK-Unk-2009	Not on treatment/medicatio n				

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-	17 End Date:2016-08-1	8							
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01					
Subject No :k01-0059	Subject Initials :LKH DOB :08/05/1981		Sex:Female	Race:Asian	Height:153cm	Weight:74g			
First administration	date of batch :		Batch number :	•					
Study Drug	Start Date	Start Date		Change in Dose					
Gonal-f New Pen Stimulation Treatment	06/02/2015	06/02/2015							
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	06/13/2015	07/02/2015		Related	Mild				
Causality Factors	•	Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity			
None(Othervalue:)		Dose not changed	None	Resolved					
Event Description:	Event Description:								
Subject received con	ncomitant medications:								
Does the subject ha	ve any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-1	7 End Date:2016-08-18	8					
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01			
Subject No :k01-035	Subject Initials :PSS DOB :01/19/1983		Sex:Female	Race:Asian	Height:161cm	Weight:56g	
First administration da	ate of batch :		Batch number :				
Study Drug	Start Date		Dose	Change in Dose	ge in Dose		
Gonal-f New Pen Stimulation Treatment	03/11/2015		225				
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	03/23/2015	04/21/2015		Related	Mild		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
None(Othervalue:)		Dose not changed	None	Resolved			
Event Description: no	addtional data			1	•	•	
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical cond	itions:No				
Condition				Start Date	Related to study condition	Ongoing	

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Non Serious Adverse Drug Reactions Report								
Start Date:2016-08-1	7 End Date:2016-08-1							
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01				
Subject No :k01-036	Subject Initials :JSY DOB :07/28/1981		Sex:Female	Race:Asian	Height:162cm	Weight:54g		
First administration da	ate of batch :		Batch number :					
Study Drug	Start Date		Dose	Change in Dose				
Gonal-f New Pen Stimulation Treatment	03/13/2015		225					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity			
OHSS	03/24/2015	04/07/2015		Related	Moderate			
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity		
None(Othervalue:)		Dose not changed	Concomitant medication **	Resolved				
Event Description:								
Subject received con	comitant medications:							
Does the subject hav	e any relevant past or	present medical cond	itions:No					
Condition				Start Date	Related to study condition	Ongoing		

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Non Serious Adverse Drug Reactions Report									
Start Date:2016-08-1	7 End Date:2016-08-1		0						
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01	No:K01				
Subject No :k01-040	Subject Initials :LBH DOB :03/05/1985		Sex:Female	Race:Asian	Height:164cm	Weight:61g			
First administration d	First administration date of batch :			4	1				
Study Drug	Start Date		Dose	Change in Dose		1			
Gonal-f New Pen Stimulation Treatment	04/06/2015		225						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity				
OHSS	04/15/2015	04/30/2015		Related	Mild				
Causality Factors	•	Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity			
None(Othervalue:)		Dose not changed	None	Resolved					
Event Description:				ł					
Subject received con	comitant medications:								
Does the subject hav	e any relevant past or	present medical cond	itions:No						
Condition				Start Date	Related to study condition	Ongoing			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-1	17 End Date:2016-08-18				•		
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01	SiteNo:K01		
Subject No :k01-049	Subject Initials :PGR	DOB :02/20/1983	Sex:Female	Race:Asian	Height:168cm	Weight:62g	
First administration d	ate of batch :		Batch number :	4			
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/02/2015	05/02/2015 300					
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/13/2015	05/29/2015		Related	Mild		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
None(Othervalue:)		Not applicable	Concomitant medication **	Resolved			
Event Description:alk	oumin treatment and re	sloved	_1	4			
Subject received con	comitant medications:						
Does the subject hav	ve any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
laparoscopic ovary c	ystectomy		UK-UNK-2004	Not on treatment/medicatio n			

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	Non Se	rious Adv	erse Drug	Reactions	s Report		
Start Date:2016-08-1	7 End Date:2016-08-18				•		
Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo:K01			
Subject No :k01-050	Subject Initials :KHG	DOB :04/22/1984	Sex:Female	Race:Asian	Height:150cm	Weight:46g	
First administration d	ate of batch :		Batch number :	4	-		
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	05/03/2015 300						
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity		
OHSS	05/14/2015	05/29/2015		Related	Mild		
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity	
None(Othervalue:)		Dose not changed	Led to study termination	Resolved			
Event Description:				<u>•</u>		•	
Subject received con	comitant medications:						
Does the subject hav	e any relevant past or	present medical cond	itions:Yes				
Condition				Start Date	Related to study condition	Ongoing	
ovary cystectomy			UK-Jan-2014	Not on treatment/medicatio n			

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