

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541		Investigator :Fei Gong		Country of Investigator :China		SiteNo :C01	
Subject No :C01-0001		Subject Initials :TTW		DOB :05/13/1988		Sex :Female	
				Race :Asian		Height :157cm	
						Weight :41.5kg	
First administration date of batch :				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	
OHSS		06/06/2015		06/16/2015		Related	
						Causality to study drug	
						Moderate	
Causality Factors		Action Taken with Study Treatment		Other action taken		Outcome	
None(Othervalue:)		Not applicable		Concomitant procedure**		Resolved	
Event Description :							
Subject received concomitant medications:							
Does the subject have any relevant past or present medical conditions :Yes							
Condition				Start Date		Related to study condition	
Bilateral fallopian tube obstruction				04/11/2014		Not on treatment/medication	
						Ongoing	

12-OCT-15

Fei Gong/EMR700623-541/C01-0001

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541	Investigator :Fei Gong		Country of Investigator :China	SiteNo :C01		
Subject No :C01-0068	Subject Initials :X-S	DOB :04/16/1989	Sex :Female	Race :Asian	Height :160cm	Weight :43kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	06/26/2015		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		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :Yes						
Condition				Start Date	Related to study condition	Ongoing
The left Hydrosalpinx and adhesions,less patency.Right fallopian tube partially blocked				03/22/2014	Not on treatment/medication	Ongoing
right fallopian tube resection because of Ectopic pregnancy				UK-Oct-2011	Not on treatment/medication	

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Fei Gong/EMR700623-541/C01-0068

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541	Investigator :NA		Country of Investigator :China	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
Concomitant medication**(Other value:)		Not applicable	Led to study termination	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :No						
Condition				Start Date	Related to study condition	Ongoing

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NA/EMR700623-541/C02-0010

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541	Investigator :NA		Country of Investigator :China	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	05/19/2015		150			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS risk				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			
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :Yes						
Condition				Start Date	Related to study condition	Ongoing
Palace laparoscopy				Uk-Unk-2013	Not on treatment/medication	

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NA/EMR700623-541/C02-0102

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541	Investigator :Ying Zhong		Country of Investigator :China	SiteNo :C05		
Subject No :C05-0001	Subject Initials :XFZ	DOB :01/21/1988	Sex :Female	Race :Asian	Height :159cm	Weight :50kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	08/01/2015		225			
Adverse Event	Start Date	End Date	Time related to study treatment	Causality to study drug	Severity	
OHSS	08/16/2015	08/18/2015		Related	Severe	
Causality Factors		Action Taken with Study Treatment	Other action taken	Outcome	AE Special Interest	AE dose limiting toxicity
None(Othervalue:)		Dose not changed	Concomitant medication **	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :Yes						
Condition				Start Date	Related to study condition	Ongoing
oophorocystectomy				UK-Jul-2013	Not on treatment/medication	
salpingoplasty				UK-Feb-2014	Not on treatment/medication	
salpingitis after previous tubal occlusion				UK-Feb-2014	Not on treatment/medication	Ongoing

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Ying Zhong/EMR700623-541/C05-0001

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

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 :74kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
Gonal-f New Pen Stimulation Treatment	06/02/2015		225			
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	Led to study termination	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :No						
Condition				Start Date	Related to study condition	Ongoing

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NA/EMR700623-541/k01-0059

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

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 :56kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change 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	Led to study termination	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :No						
Condition				Start Date	Related to study condition	Ongoing

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NA/EMR700623-541/k01-035

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

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 :54kg
First administration date 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	Led to study termination	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :No						
Condition				Start Date	Related to study condition	Ongoing

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NA/EMR700623-541/k01-036

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541	Investigator :NA		Country of Investigator :Korea	SiteNo :K01		
Subject No :k01-040	Subject Initials :LBH	DOB :03/05/1985	Sex :Female	Race :Asian	Height :164cm	Weight :61kg
First administration date of batch :			Batch number :			
Study Drug	Start Date		Dose	Change in Dose		
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	Led to study termination	Resolved		
Event Description:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :No						
Condition				Start Date	Related to study condition	Ongoing

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NA/EMR700623-541/k01-040

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

Study :EMR700623-541		Investigator :NA		Country of Investigator :Korea		SiteNo :K01	
Subject No :k01-049	Subject Initials :PGR	DOB :02/20/1983	Sex :Female	Race :Asian	Height :168cm	Weight :62kg	
First administration date of batch :			Batch number :				
Study Drug	Start Date		Dose	Change in Dose			
Gonal-f New Pen Stimulation Treatment	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	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 **,Led to study termination	Resolved			
Event Description:							
Subject received concomitant medications:							
Does the subject have any relevant past or present medical conditions :Yes							
Condition				Start Date	Related to study condition	Ongoing	
laparoscopic ovary cystectomy				UK-UNK-2004	Not on treatment/medication		

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NA/EMR700623-541/k01-049

Non Serious Adverse Drug Reactions Report

Start Date:2015-10-08 End Date:2015-10-12

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 :46kg
First administration date of batch :			Batch number :			
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:						
Subject received concomitant medications:						
Does the subject have any relevant past or present medical conditions :Yes						
Condition				Start Date	Related to study condition	Ongoing
ovary cystectomy				UK-Jan-2014	Not on treatment/medication	

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NA/EMR700623-541/k01-050

