Non Serious Adverse Drug Reactions Report Start Date:2016-07-29 End Date:2016-07-30

| Study :EMR700623-541 | Investigator :NA | | Country of Investigator :China | SiteNo:C02 | iteNo:C02 | | |
|---|--------------------------|--------------------------------------|-----------------------------------|-------------------------|------------------------------------|---------------------------|--|
| Subject No :C02-0201 | Subject Initials :J-L | 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 | Jose | | |
| Gonal-f New Pen Stimulation Treatment | 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:M | 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 occlusion | | | | Uk-Unk-2012 | Not on treatment/medicatio n | | |

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| Non Serious Adverse Drug Reactions Report | | | | | | | |
|---|-------------------------|--------------------------------------|-----------------------------------|-------------------------|------------------------------------|---------------------------|--|
| Start Date:2016-07-2 | 29 End Date:2016-07-3 | C | | | | | |
| 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 | Height:161cm | Weight:60kg | |
| First administration date of batch : | | | Batch number : | | | | |
| Study Drug | Start Date | | Dose | Change in Dose | | | |
| 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 has | 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 abortion | | | | 05/18/2012 | Not on treatment/medicatio n | | |

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| Non Serious Adverse Drug Reactions Report | | | | | | | | |
|---|-------------------------|--------------------------------------|-----------------------------------|-------------------------|----------------------------|---------------------------|--|--|
| Start Date:2016-07-2 | 29 End Date:2016-07-3 | 0 | | | | | | |
| 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 | <u> </u> | - | - | | | | |
| 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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