| | Non Se | rious Adv | erse Drug | Reactions | s Report | | |
|---|--------------------------|---|---------------------------------|------------------------------|----------------------------|---------------------------|--|
| Start Date:2016-04 | -07 End Date:2016-04-0 | 8 | | | • | | |
| Study :EMR700623-541 | Investigator :NA | Investigator :NA Country of Investigator :China | | 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 : | Batch number : | | | |
| Study Drug | Start Date | | Dose | Change in Dose | | | |
| Gonal-f New Pen Stimulation Treatment | 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: | | | -! | | <u>'</u> | | |
| 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 obst | ruction | | Uk-Apr-2014 | Not on treatment/medicatio n | | | |

07-APR-16 NA/EMR700623-541/C02-0200