## Non Serious Adverse Drug Reactions Report Start Date:2016-05-20 End Date:2016-05-23

| Otart Date.2010 00 2                        | 0 End Date.2010-03-2   | 0                                    |                                   |                         |                            |                           |  |
|---|------------------------|--------------------------------------|-----------------------------------|-------------------------|----------------------------|---------------------------|--|
| Study<br>:EMR700623-541                     | Investigator :NA       |                                      | Country of<br>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 date of batch :        |                        |                                      | Batch number :                    |                         |                            |                           |  |
| Study Drug                                  | Start Date             |                                      | Dose                              | Change in Dose          |                            |                           |  |
| Gonal-f New Pen<br>Stimulation<br>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<br>Study Treatment | Other action taken                | Outcome                 | AE Special Interest        | AE dose limiting toxicity |  |
| None(Othervalue:)                           |                        | Dose not changed                     | None                              | Resolved                |                            |                           |  |
| Event Description:                          |                        |                                      |                                   |                         |                            |                           |  |
| Subject received con-                       | comitant medications:  |                                      |                                   |                         |                            |                           |  |
| Does the subject hav                        | e any relevant past or | present medical cond                 | tions:No                          |                         |                            |                           |  |
| Condition                                   |                        |                                      |                                   | Start Date              | Related to study condition | Ongoing                   |  |

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