|  | Non S                                | Serious Adv                          | erse Drug                                 | Reactions               | Report                      |                           |
|--|--------------------------------------|--------------------------------------|---|-------------------------|-----------------------------|---------------------------|
| Start Date:2015-09-18                    | End Date:2015-09-24                  |                                      |   |                         | -                           |                           |
| Study<br>:EMR700623-541                  | Investigator :Fei Go                 | ong                                  | Country of Investigator : China           | SiteNo:C01              |                             |                           |
| Subject No<br>:C01-0001                  | Subject Initials :TTW DOB:05/13/1988 |                                      | <b>DOB</b> :05/13/1988 <b>Sex:</b> Female | Race:Asian              | Height:157cm                | Weight:41.5kg             |
| First administration date of batch :     |                                      |                                      | Batch number :                            |                         |                             |                           |
| Study Drug                               | Study Drug Start Date                |                                      | Dose                                      | Change in Dose          |                             | 1                         |
| Gonal-f New Pen<br>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                           | 06/16/2015                           |   | Related                 | Moderate                    |                           |
| Causality Factors                        | •                                    | Action Taken with<br>Study Treatment | Other action taken                        | Outcome                 | AE Special Interest         | AE dose limiting toxicity |
| None(Othervalue:)                        |                                      | Not applicable                       | Concomitant procedure**                   | Resolved                |                             |                           |
| Event Description:                       |                                      |                                      |   | l                       |                             | L                         |
| Subject received conce                   | omitant medications:                 |                                      |   |                         |                             |                           |
| Does the subject have                    | e any relevant past or               | present medical conditio             | ns:Yes                                    |                         |                             |                           |
| Condition                                |                                      |                                      |   | Start Date              | Related to study condition  | Ongoing                   |
| Bilateral fallopian tube obstruction     |                                      |                                      |   | 04/11/2014              | Not on treatment/medication | Ongoing                   |

Fei Gong/EMR700623-541/C01-0001

|   | Non Se                                | rious Adv              | erse Drug                         | Reactions               | Report                      |                           |  |
|---|---------------------------------------|------------------------|-----------------------------------|-------------------------|-----------------------------|---------------------------|--|
| Start Date:2015-09-18   | 8 End Date:2015-09-24                 |                        |                                   |                         | -                           |                           |  |
| <b>Study</b> :EMR700623-541   | Investigator :Fei Gong                | g                      | Country of<br>Investigator :China | SiteNo:C01              |                             |                           |  |
| Subject No<br>: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          |                             | 1                         |  |
| Gonal-f New Pen<br>Stimulation Treatmen   | 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:  |                                       | l                      | · I                               | l                       |                             | I                         |  |
| Subject received cond   | comitant medications:                 |                        |                                   |                         |                             |                           |  |
| Does the subject hav  | ve any relevant past or pr            | esent medical conditio | ns: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 |                           |  |

Fei Gong/EMR700623-541/C01-0068

|  | Non Se                                | rious Adv                            | erse Drug                       | Reactions               | Report                     |                           |  |
|--|---------------------------------------|--------------------------------------|---------------------------------|-------------------------|----------------------------|---------------------------|--|
| Start Date:2015-09-18                    | End Date:2015-09-24                   |                                      | <u> </u>                        |                         | •                          |                           |  |
| <b>Study</b> :EMR700623-541              | Investigator :NA                      |                                      | Country of Investigator :China  | SiteNo:C02              | SiteNo:C02                 |                           |  |
| Subject No<br>: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                               | Study Drug Start Date                 |                                      | Dose                            | Change in Dose          |                            | -1                        |  |
| Gonal-f New Pen<br>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                   |                           |  |
| <b>Causality Factors</b>                 |                                       | Action Taken with<br>Study Treatment | Other action taken              | Outcome                 | AE Special Interest        | AE dose limiting toxicity |  |
| Concomitant medicati                     | on**(Othervalue:)                     | Not applicable                       | Led to study termination        | Resolved                |                            |                           |  |
| Event Description:                       |                                       |                                      | ·                               |                         | 1                          | 1                         |  |
| Subject received conc                    | omitant medications:                  |                                      |                                 |                         |                            |                           |  |
| Does the subject have                    | e any relevant past or pr             | esent medical conditio               | ns:No                           |                         |                            |                           |  |
| Condition                                |                                       |                                      |                                 | Start Date              | Related to study condition | Ongoing                   |  |

|   | Non S                          | Serious Adv                          | erse Drug                       | Reactions               | Report                      |                           |
|---|--------------------------------|--------------------------------------|---------------------------------|-------------------------|-----------------------------|---------------------------|
| Start Date:2015-09-1                    | 8 End Date:2015-09-24          |                                      | <u> </u>                        |                         | *                           |                           |
| <b>Study</b> :EMR700623-541             | Investigator :NA               | Investigator :NA                     |                                 | SiteNo:C02              |                             |                           |
| Subject No<br>:C02-0102                 | Subject Initials<br>:HYD       | <b>DOB</b> :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<br>Stimulation Treatmer | inulation 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                    |                           |
| <b>Causality Factors</b>                | •                              | Action Taken with<br>Study Treatment | Other action taken              | Outcome                 | AE Special Interest         | AE dose limiting toxicity |
| None(Othervalue:)                       |                                | Not applicable                       | None                            |                         |                             |                           |
| Event description:                      |                                | L                                    | L                               |                         |                             |                           |
| Subject received cond                   | comitant medications:          |                                      |                                 |                         |                             |                           |
| Does the subject hav                    | ve any relevant past or        | present medical condition            | ons:Yes                         |                         |                             |                           |
| Condition                               |                                |                                      |                                 | Start Date              | Related to study condition  | Ongoing                   |
| Palace laparoscopy                      |                                |                                      |                                 | Uk-Unk-2013             | Not on treatment/medication |                           |
|   |                                |                                      |                                 | l                       |                             | 24-                       |

| us Adv   | erse Drug                       | Reactions                | Report                                |   |
|--|---------------------------------|--------------------------|---------------------------------------|---|
|  |                                 |                          | •                                     |   |
|  | Country of Investigator : China | SiteNo:C02               |                                       |   |
| <b>3</b> :05/12/1989   | Sex:Female                      | Race: Asian              | Height:165cm                          | Weight:63.5kg   |
|  | Batch number :                  | !                        |                                       |   |
| Study Drug Start Date  |                                 | Change in Dose           |                                       | ·I  |
| onal-f New Pen 05/19/2015 mulation Treatment                                   |                                 |                          |                                       |   |
| Date   | Time related to study treatment | Causality to study drug  | Severity                              |   |
|  |                                 | Related                  | Moderate                              |   |
| on Taken with<br>y Treatment   | Other action taken              | Outcome                  | AE Special Interest                   | AE dose limiting toxicity                               |
| applicable   | None                            | Resolved                 |                                       |   |
|  | !                               | !                        |                                       | ·!  |
|  |                                 |                          |                                       |   |
| medical condition  | ns:Yes                          |                          |                                       |   |
| Condition  |                                 |                          | Related to study condition            | Ongoing   |
| The right side of the open line on the right side of salpingectomy tubal pregr |                                 |                          | Not on treatment/medication           |   |
| r  | ngectomy tubal p                | ngectomy tubal pregnancy | ngectomy tubal pregnancy  Uk-Unk-2011 | condition ngectomy tubal pregnancy  Uk-Unk-2011  Not on |

|   | Non Se                  | rious Adv              | erse Drug                       | Reactions               | Report                      |             |  |  |
|---|-------------------------|------------------------|---------------------------------|-------------------------|-----------------------------|-------------|--|--|
| Start Date:2015-09-18                               | End Date:2015-09-24     |                        |                                 |                         |                             |             |  |  |
| <b>Study</b> :EMR700623-541                         | Investigator : Ying Zho | •                      | Country of Investigator : China | SiteNo:C05              |                             |             |  |  |
| Subject No<br>:C05-0001                             | Subject Initials :XFZ   | <b>DOB</b> :01/21/1988 | Sex:Female                      | Race: Asian             | Height:159cm                | Weight:50kg |  |  |
| First administration date of batch :                |                         |                        | Batch number :                  | <u> </u>                |                             |             |  |  |
| Study Drug Start Date                               |                         |                        | Dose                            | Change in Dose          |                             |             |  |  |
| Gonal-f New Pen<br>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:                                  |                         |                        | •                               |                         | -1                          | •           |  |  |
| Subject received conco                              | mitant medications:     |                        |                                 |                         |                             |             |  |  |
| Does the subject have                               | any relevant past or pr | esent medical conditio | ns: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 previou                           | us tubal occlusion      |                        |                                 | UK-Feb-2014             | Not on treatment/medication | Ongoing     |  |  |

24-SEP-15 Ying Zhong/EMR700623-541/C05-0001

|  | Non Se                                | rious Adv                            | erse Drug                       | Reactions               | Report                     |                           |
|--|---------------------------------------|--------------------------------------|---------------------------------|-------------------------|----------------------------|---------------------------|
| Start Date:2015-09-18 I                  | End Date:2015-09-24                   |                                      |                                 |                         | -                          |                           |
| <b>Study</b> :EMR700623-541              | Investigator :NA                      | Investigator :NA                     |                                 | 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 :                  | ı                       | I.                         |                           |
| Study Drug                               | Start Date                            |                                      | Dose                            | Change in Dose          |                            |                           |
| Gonal-f New Pen<br>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                       |                           |
| <b>Causality Factors</b>                 |                                       | Action Taken with<br>Study Treatment | Other action taken              | Outcome                 | AE Special Interest        | AE dose limiting toxicity |
| None(Othervalue:)                        | None(Othervalue:)  Dose not changed   |                                      | Led to study termination        | Resolved                |                            |                           |
| Event Description:                       |                                       |                                      |                                 |                         | •                          | •                         |
| Subject received concor                  | mitant medications:                   |                                      |                                 |                         |                            |                           |
| Does the subject have                    | any relevant past or pro              | esent medical conditio               | ns:No                           |                         |                            |                           |
| Condition                                |                                       |                                      |                                 | Start Date              | Related to study condition | Ongoing                   |

|  | Non Se                                | erious Adv                           | erse Drug                         | Reactions               | Report                     |                           |
|--|---------------------------------------|--------------------------------------|-----------------------------------|-------------------------|----------------------------|---------------------------|
| Start Date:2015-09-18                    | End Date:2015-09-24                   |                                      |                                   |                         | -                          |                           |
| <b>Study</b> :EMR700623-541              | Investigator :NA                      |                                      | Country of<br>Investigator :Korea | l "                     |                            |                           |
| 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 :                    |                         | <u>.</u>                   |                           |
| Study Drug                               | Start Date                            |                                      | Dose                              | Change in Dose          |                            | •                         |
| Gonal-f New Pen<br>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<br>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 conco                   | mitant medications:                   |                                      |                                   |                         |                            |                           |
| Does the subject have                    | any relevant past or pr               | esent medical conditio               | ns:No                             |                         |                            |                           |
| Condition                                |                                       |                                      |                                   | Start Date              | Related to study condition | Ongoing                   |

|  | Non Se                                | erious Adv                           | erse Drug                         | Reactions               | Report                     |                           |
|--|---------------------------------------|--------------------------------------|-----------------------------------|-------------------------|----------------------------|---------------------------|
| Start Date:2015-09-18                    | End Date:2015-09-24                   |                                      |                                   |                         | -                          |                           |
| <b>Study</b> :EMR700623-541              | Investigator :NA                      |                                      | Country of<br>Investigator :Korea | 1 · ·                   |                            |                           |
| 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<br>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                   |                           |
| <b>Causality Factors</b>                 |                                       | Action Taken with<br>Study Treatment | Other action taken                | Outcome                 | AE Special Interest        | AE dose limiting toxicity |
| None(Othervalue:)  Dose not changed      |                                       | Dose not changed                     | Led to study termination          | Resolved                |                            |                           |
| Event Description:                       |                                       | •                                    |                                   |                         | •                          | •                         |
| Subject received conco                   | mitant medications:                   |                                      |                                   |                         |                            |                           |
| Does the subject have                    | any relevant past or pr               | esent medical conditio               | ns:No                             |                         |                            |                           |
| Condition                                |                                       |                                      |                                   | Start Date              | Related to study condition | Ongoing                   |

|  | Non Se                                | rious Adv                            | erse Drug                         | Reactions               | Report                     |                           |
|--|---------------------------------------|--------------------------------------|-----------------------------------|-------------------------|----------------------------|---------------------------|
| Start Date:2015-09-18                    |                                       |                                      | <u> </u>                          |                         |                            |                           |
| <b>Study</b> :EMR700623-541              | Investigator :NA                      |                                      | Country of<br>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 :                    |                         | <u>.</u>                   |                           |
| Study Drug                               | udy Drug Start Date                   |                                      | Dose                              | Change in Dose          |                            |                           |
| Gonal-f New Pen<br>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<br>Study Treatment | Other action taken                | Outcome                 | AE Special Interest        | AE dose limiting toxicity |
| None(Othervalue:)                        | None(Othervalue:)  Dose not changed   |                                      | Led to study termination          | Resolved                |                            |                           |
| Event Description:                       |                                       |                                      |                                   |                         | •                          | •                         |
| Subject received conco                   | mitant medications:                   |                                      |                                   |                         |                            |                           |
| Does the subject have                    | any relevant past or pr               | esent medical conditio               | ns:No                             |                         |                            |                           |
| Condition                                |                                       |                                      |                                   | Start Date              | Related to study condition | Ongoing                   |

|   | Non Se                  | rious Adv              | erse Drug  | Reactions               | Report                      |                           |  |
|---|-------------------------|------------------------|--|-------------------------|-----------------------------|---------------------------|--|
| Start Date:2015-09-18                               | End Date:2015-09-24     |                        |  |                         | -                           |                           |  |
| Study<br>:EMR700623-541                             | Investigator :NA        |                        | Country of SiteNo:K01 Investigator : Korea               |                         |                             |                           |  |
| Subject No :k01-049                                 | Subject Initials :PGR   | <b>DOB</b> :02/20/1983 | Sex:Female   | Race:Asian              | Height:168cm                | Weight:62kg               |  |
| First administration date of batch :                |                         | Batch number :         |  | <u>.</u>                |                             |                           |  |
| Study Drug  | Study Drug Start Date   |                        | Dose   | Change in Dose          |                             | 1                         |  |
| Gonal-f New Pen<br>Stimulation Treatment            |                         |                        | 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<br>medication **,Led to<br>study termination | Resolved                |                             |                           |  |
| Event Description:                                  |                         | I                      | · I  |                         |                             | I                         |  |
| Subject received conco                              | omitant medications:    |                        |  |                         |                             |                           |  |
| Does the subject have                               | any relevant past or pr | esent medical conditio | ns:Yes   |                         |                             |                           |  |
| Condition   |                         |                        |  | Start Date              | Related to study condition  | Ongoing                   |  |
| laparoscopic ovary cystectomy                       |                         |                        |  | UK-UNK-2004             | Not on treatment/medication |                           |  |

|  | Non S                                  | Serious Adv                          | erse Drug                             | Reactions               | Report                      |                           |  |
|--|--|--------------------------------------|---------------------------------------|-------------------------|-----------------------------|---------------------------|--|
| Start Date:2015-09-18                    | End Date:2015-09-24                    |                                      |                                       |                         | -                           |                           |  |
| Study<br>:EMR700623-541                  | Investigator :NA                       |                                      | Country of Si<br>Investigator : Korea | SiteNo:K01              |                             |                           |  |
| Subject No :k01-050                      | Subject Initials: BOB: 04/22/1984: KHG |                                      | 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<br>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<br>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:                       |  | l                                    | L                                     | l                       | I                           |                           |  |
| Subject received conce                   | omitant medications:                   |                                      |                                       |                         |                             |                           |  |
| Does the subject have                    | e any relevant past or                 | present medical conditio             | ns:Yes                                |                         |                             |                           |  |
| Condition                                |  |                                      |                                       | Start Date              | Related to study condition  | Ongoing                   |  |
| ovary cystectomy                         |  |                                      |                                       | UK-Jan-2014             | Not on treatment/medication |                           |  |