

## Non Serious Adverse Drug Reactions Report

Start Date:2016-06-21 End Date:2016-06-22

|  |                    |                                   |                                  |                         |                            |                           |
|--|--------------------|-----------------------------------|----------------------------------|-------------------------|----------------------------|---------------------------|
| Study :EMR200136_583   | Investigator :NA   |                                   | Country of Investigator :Romania | SiteNo:005              |                            |                           |
| Subject No :005-0001   | Subject Initials : | DOB :08/27/1996                   | Sex:Female                       | Race:Caucasian          | Height:169(cm)             | Weight:65(kg)             |
| First administration date of batch :                                     |                    |                                   | Batch number :                   |                         |                            |                           |
| Study Drug   | Start Date         |                                   | Dose                             | Change in Dose          |                            |                           |
| Visit 5 (Month12)/Early Termination                                      |                    |                                   |                                  |                         |                            |                           |
| Visit 1/ Baseline (Day 1)  | 02/12/2015         |                                   | 5                                |                         |                            |                           |
| Adverse Event  | Start Date         | End Date                          | Time related to study treatment  | Causality to study drug | Severity                   |                           |
| local erythema and induration local site injection                       | 04/28/2015         |                                   |                                  | Suspected               | Mild                       |                           |
| Causality Factors  |                    | Action Taken with Study Treatment | Other action taken               | Outcome                 | AE Special Interest        | AE dose limiting toxicity |
| None(Othervalue:)  |                    | Drug withdrawn                    | Led to study termination         | Ongoing                 |                            |                           |
| Event description:   |                    |                                   |                                  |                         |                            |                           |
| Subject received concomitant medications                                 |                    |                                   |                                  |                         |                            |                           |
| Name of medication   | Start Date         | Ongoing                           | End Date                         | Dose                    | Unit                       | Frequency                 |
| Does the subject have any relevant past or present medical conditions:No |                    |                                   |                                  |                         |                            |                           |
| Condition  |                    |                                   |                                  | Start Date              | Related to study condition | Ongoing                   |

22-JUN-16

NA/EMR200136\_583/005-0001

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Start Date:2016-06-21 End Date:2016-06-22

|  |                                      |  |                                    |                            |                                 |
|--|--------------------------------------|--|------------------------------------|----------------------------|---------------------------------|
| Study<br>:EMR200136_583  | Investigator :NA                     | Country of<br>Investigator<br>:Romania | SiteNo:005                         |                            |                                 |
| Subject No<br>:005-0009  | Subject Initials :                   | DOB :04/22/1992                        | Sex:Female                         | Race:Caucasian             | Height:173(cm)<br>Weight:63(kg) |
| First administration date of batch :                                     |                                      |  | Batch number :                     |                            |                                 |
| Study Drug   | Start Date                           | Dose                                   | Change in Dose                     |                            |                                 |
| Visit 5<br>(Month12)/Early<br>Termination                                |                                      |  |                                    |                            |                                 |
| Visit 1/ Baseline<br>(Day 1)   | 03/26/2015                           | 9                                      |                                    |                            |                                 |
| Adverse Event  | Start Date                           | End Date                               | Time related to<br>study treatment | Causality to study<br>drug | Severity                        |
| local erythema,<br>pain, ecchymosis                                      | 06/09/2015                           |  |                                    | Suspected                  | Mild                            |
| Causality Factors  | Action Taken with<br>Study Treatment | Other action taken                     | Outcome                            | AE Special Interest        | AE dose limiting<br>toxicity    |
| None(Othervalue:)  | Drug withdrawn                       | Led to study<br>termination            | Ongoing                            |                            |                                 |
| Event description:   |                                      |  |                                    |                            |                                 |
| Subject received concomitant medications                                 |                                      |  |                                    |                            |                                 |
| Name of medication   | Start Date                           | Ongoing                                | End Date                           | Dose                       | Unit<br>Frequency               |
| Does the subject have any relevant past or present medical conditions:No |                                      |  |                                    |                            |                                 |
| Condition  | Start Date                           | Related to study<br>condition          | Ongoing                            |                            |                                 |

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