*For National Haemovigilance Office use only*

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| HV/NM/Sequence/Year | Date received  |  | Signature |  |

 **Attachment 3: Blood Establishment: Notification of a Near Miss / Serious Adverse Event to National Haemovigilance Office**

|  |
| --- |
| **Reporting establishment** |
| **Report identification** |
| **Reporting date (year/month/day)** |
| **Date of serious event (year/month/day)** |
| **Serious adverse event, due to a deviation in:**  | **Specification** |
| **Component defect** | **Equipment failure** | **Human error** | **System Failure**  | **Materials**  | **Other****(specify)** |
| **Processing**  |  |  |  |  |  |  |
| **Component Selection**  |  |  |  |  |  |  |
| **Compatibility testing/Cross-matching**  |  |  |  |  |  |  |
| **Storage** |  |  |  |  |  |  |
| **Issue**  |  |  |  |  |  |  |
| **Distribution** |  |  |  |  |  |  |
| **Others (specify)** |  |  |  |  |  |  |
| **Short Description of event:** |
| **Has this event been reviewed by a Consultant****Haematologist/Pathologist?** | **Yes** | **No** |

**Signed:** \_\_\_\_\_\_\_\_\_\_ **Date:** \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

**Email Address \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**