#

**CABRINI RESEARCH GOVERNANCE**

**SAFETY REPORTING FORM**

*Refer to Cabrini’s Safety Monitoring and Reporting in Research Policy for guidance on reportable events.*

| **Cabrini Project #:** | **Cabrini Project Title:** |
| --- | --- |
| **Participant Study ID:** |  |
| **Report****Type** (e.g. initial, follow-up 1, 2, 3 etc) | **Report Date** | **Incident****Date**(the same for each report) | **Nature of Event** | **Was the event related to the study intervention (drug, device, procedure)?** | **Was****it a death?** | **Investigator’s comments & recommendations e.g.:****Is action required?****Has dosing been suspended?****Does the participant remain on study?****Has the event resolved?** |
| (Delete rows below as required) |
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| **Principal Investigator name**:  |
| **Principal Investigator signature**:  | **Date**:  |