



(Annexure 6)

Protocol Violation/Deviation Reporting Form (Reporting by case)

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EC Ref. No. (For office use):

Title of study:

Principal Investigator (Name, Designation and Affiliation):

1. Date of EC approval dd mm yy Date of start of study dd mm yy
2. Participant ID: Date of occurrence dd mm yy
3. Total number of deviations /violations reported till date in the study:
4. Deviation/Violation identified by: Principal Investigator/study team Sponsor/Monitor
SAE Sub Committee/EC
5. Is the deviation related to (Tick the appropriate box):

Consenting	<input type="checkbox"/>	Source documentation	<input type="checkbox"/>
Enrollment	<input type="checkbox"/>	Staff	<input type="checkbox"/>
Laboratory assessment	<input type="checkbox"/>	Participant non-compliance	<input type="checkbox"/>
Investigational Product	<input type="checkbox"/>	Others (specify) <input style="width: 150px; height: 20px; border: 1px solid black;" type="text"/>	<input type="checkbox"/>
Safety Reporting	<input type="checkbox"/>		
6. Provide details of Deviation/Violation:
.....
.....
.....
7. Corrective action taken by PI/Co-I:
.....
.....
.....
8. Impact on (if any): Study participant Quality of data
9. Are any changes to the study/protocol required? Yes No
If yes, give details.....
.....

Signature of PI:

dd mm yy