

## **Protocol Deviation Reporting Log**

## **TriHealth Institutional Review Board**

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Please utilize this form to report minor protocol deviations and simple logistical conflicts at the time of continuing review. None of the items on this list should include any violations from currently approved protocol procedures that adversely affect the rights, safety, health or welfare of subjects, nor significantly impact the integrity of research data. Those should be reported as an **Unanticipated Problem**.

Study teams are responsible for updating this log in a timely manner and for maintaining this log in the Regulatory Binder.

4 or more **Minor** deviations of the same type require you to submit this form to the IRB as soon as the fourth deviation is recognized. The IRB Chair will decide if the multiple deviations constitute a **Major** violation.

## **Protocol Deviation Codes:**

- 1=Recruitment Procedure
- 2=Documentation
- 3=Withdrawal Criteria
- 4=Concomitant Medications
- 5=Event Outside Time Frame
- 6=Study Conduct
- 7=Informed Consent
- 8=IRB Reporting
- 9=Administrative Issues
- 10=Drug Accountability
- 11=Missing Lab Results
- 12=Other

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Event Date				
			<u></u>	<del>-</del>
Subject ID/Site/ Initials				
Deviation Code (1-12)				
Description of Deviation (including				
assessment of the net effect on Risk/Benefit)				
Reason for Deviation and Date of Sponsor				
Approval				
Corrective Action				
Taken to Avoid Recurrence				
Reduirence				
certify the above as being an a	ccurate and comple	ete accounting of	the minor deviations	that occurred
luring the reporting period.	comple	oto accounting of	e minor deviations	
make at Dainelia at Lacrastics				
nature of Principal Investiga	tor:			
te:				
Submit this protocol dev				

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