

# Instruction for Use

## MONITORING SITE VISIT LOG

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<b>Template</b>	
<b>Purpose</b>	Keep a record all monitoring visits, starting with Site Initiation
<b>Use</b>	Study coordinator, principal investigator, monitor
<b>Details</b>	This log should provide a comprehensive list of all monitoring visits
<b>Best Practice Recommendations</b>	<p>Record monitoring visits as they occur, to ensure completeness and accuracy of the data. If a visit occurs over more than one day, each day should be recorded on a separate line.</p> <p>In the upper part of the page identifying the study, delete (Local) if the study is monocentric, and ( ) if the study is multicentric</p> <p>The clinical site monitor should sign each line as a monitoring visit begins</p> <p>The Local Investigator or site staff should sign each line at the end of the visit</p> <p>Number each page and maintain this log in the ISF (a copy will be collected by the CRA at the end of the study for the TMF)</p> <p>Store pages in reverse chronological order, with the newest pages of the log placed at the front of the section.</p>

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