





**SUMMARIZE THE RESULTS YOU HAVE ACHIEVED THUS FAR IN YOUR STUDY.**

Include a description of any adverse events or unanticipated problems involving risks to participants or others, withdrawal of participants from research, or complaints about the research; a summary of any recent literature, findings, or other relevant information, especially information about risks associated with the research; and a copy of the current informed consent document(s).

**REVIEW OF ANNUAL PROGRESS REPORT SUBMISSION**

Renewal is **APPROVED**. New expiration date    /    /

Renewal is **APPROVED** contingent upon modifications as described below. [Submit to UWW ORSP for approval.]  
 See attachment for details

New expiration date    /    /

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Requires **FULL BOARD REVIEW** and will be included on the next IRB agenda    /    /

**RESUBMIT** with modifications/additional information as described below. [Submit to UWW ORSP for approval.]  
 See attachment for details

**REVIEW OF PROTOCOL MODIFICATIONS (AND/OR CONDITION FULFILLMENTS)**

I have reviewed the modifications to the protocol renewal and determined that the modified protocol is

is **APPROVED**.       must be **RESUBMITTED** with modifications as described below. [Submit to UWW ORSP for approval.]  
 See attachment for details

New expiration date    /    /

**REVIEW AUTHORIZATION**

<b>TYPED/PRINTED NAME IRB CHAIR/DESIGNEE</b>	<b>SIGNATURE</b>	<b>DATE</b>
<i>If you should make any changes in the protocol involving 1) method, 2) participants, 3) informed consent, and/or 4) subject identification, you must resubmit the protocol. The case number assigned to this protocol is listed on the Protocol Coversheet; please reference this number in all future correspondence. You are responsible for submission of an Annual Progress Report (if necessary) and for maintaining all records related to this project for at least three years after completion of the research project.</i>		