

List on this form all of the IND Safety Reports which the sponsor requires you to submit to the MWMC IRBs. Attach a copy of the individual reports to this form. Receipt of these reports will be confirmed by the MWMC IRB but the reports will <u>not</u> be reviewed.

*If a change in the protocol or consent form is required by the IND Safety Report submit a Change in Research Form with the IND Safety Report attached.

Please no more than 10 per submission

| A. Study Information | | | | | | |
|----------------------|---------------------------------|------------------|--|--|--|--|
| Date: | | | | | | |
| 1. | MWMC Study Number: | Protocol Number: | | | | |
| 2. | Name of Study: | | | | | |
| 3. | Name of Principal Investigator: | | | | | |

| IND # | Event Name | Study Drug | Relatedness | Changes to ICF |
|-------|------------|------------|-------------|----------------|
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Dear Dr. Desai, Enclosed are copies of the IND safety reports generated by the sponsor. I have personally reviewed each report.

Principal Investigator

Date

| Name of Person Completing Form (contact person for questions): | | | | |
|--|--|--|--|--|
| Name: | | | | |
| Title: | | | | |
| Phone Number: | | | | |
| Email Address: | | | | |
| Signature: | | | | |

PLEASE EMAIL COMPLETED FORMS AND REQUIRED DOCUMENTS TO <u>MWMCIRB@mwmc.com</u> you will receive an acknowledgement once all required information has been submitted.

| CHECKLIST |
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| Please submit all supporting documents for IND's |