



FINAL REPORT (FORM 3.4)

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|---------------------------------------------------------------------|----------------------|-------------------|----------------------|
| IRB Protocol No. | <input type="text"/> | Approval Date | <input type="text"/> |
| Protocol Title | <input type="text"/> | | |
| Principal Investigator. | <input type="text"/> | | |
| Phone number: | <input type="text"/> | E-mail address : | <input type="text"/> |
| Sponsor's Name | <input type="text"/> | | |
| Address: | <input type="text"/> | | |
| Phone number: | <input type="text"/> | E-mail address : | <input type="text"/> |
| Study site(s): | <input type="text"/> | | |
| Total Number of study participants : | <input type="text"/> | No. of Study Arms | <input type="text"/> |
| Number of participants who received the test articles: | <input type="text"/> | | |
| Study materials: | <input type="text"/> | | |
| Treatment form: | <input type="text"/> | | |
| Study dose(s): | <input type="text"/> | | |
| Duration of the study | <input type="text"/> | | |
| Objectives: | <input type="text"/> | | |
| Results: <i>(Use extra blank paper, if more space is required.)</i> | <input type="text"/> | | |
| Signature of P.I. | <input type="text"/> | | |