

Healthcare Compliance Forms and Tools Sample Request to Conduct Research Form

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Research Review Committee Documentation of Review and Approval

Request to Conduct Research at Facility

Project Title:	IRB No.:
Submitted by:	Affiliation:
Email:	Tele:

Primary Investigation (if different from person submitting):

Project Location(s):

Community Approval:

Management Company Approval:

Funding Source(s):

Grant Title (if applicable and if different from project title):



Research Submission:

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Informed consent, dated	
Privacy notice	
Drug or device brochure(s), dated	
Protocol, include any questionnaire(s), dated	
Summary Safety Guard Statement, dated	
Advertisement (if applicable), dated	
Authorization, dated	
Other (description), dated	
Consents	
IRB approval letter	

Please mail all materials to:

I assure the Research Review Committee (RRC) that all procedures performed under the project will be conducted in strict accordance with those federal regulations and internal policies that govern research involving human subjects. I agree to submit any deviation from the project in the form of an amendment for RRC approval prior to implementation. By signing this form, I am certifying that all co-investigators listed in the study are aware of the research and are agreeing to participate.

NOTE: Applications and any additional material requested by the RRC will not be processed unless neatly typed and legible, properly prepared, and signed personally by the principal investigator.

Date _____ Principal Investigator (Signature)

This protocol and informed consent statement for use of subjects in research has been reviewed and approved by the RRC for a maximum of a one-year period beyond the final approval date unless otherwise indicated as follows: ______.

Authorized RRC Signature

RRC Approval Date

Recorded in the Minutes of:

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