

Clinical Trials

- Is the trial being conducted under the auspices of an Institutional Review Board; if so confirm the name of the IRB and the affiliated hospital.
- Will the trial include any of the following types of patients: maternity, mentally or developmentally disabled, prisoners, minors, or elderly?
- How many clinical trials annually?
- Is there a written contract with the pharmaceutical company?
- Does the contract contain a contractual hold harmless or indemnification clause?
- Submit a complete copy of the contract and a copy of the consent form.
- Underwriting and risk management advice will follow upon receipt of this information.

Source: ProAssurance 2007

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