CLINICAL TRIALS

WE UNDERSTAND THE COMPLEXITIES OF TODAY'S CLINICAL TRIALS ENVIRONMENT!

Eligible Risks Include:

- ✓ **Independent Research Sites (IRS's)**
- ✓ **Contract Research Organizations (CRO's)**
- ✓ **Site Management Organizations (SMO's)**
- ✓ **Institutional Review Boards (IRB's)**
- ✓ **Data Management Safety Boards (DSMB's)**
- ✓ **NFP Research Institutes**
- ✓ **Trial Sponsors/ Biotech Firms (Less than $50M budgets)**

Our broad definition of "Clinical Research Professional Services" includes medical professional services, manufacturing, designing, handling or distribution of a drug or device in a clinical trial, pre-clinical services, quality assurance, protocol design, participant recruiting, consulting, market evaluation, data management, medical writing, audit and regulatory advisory services.

- Up to $10M Total Capacity
- Primary & Excess Placements
- Follow Form Available (above select lead markets)
- Defense Outside the Limit
- Individuals Named on FDA 1572 Form Covered
- Crisis Response/ Regulatory Coverage Included
- Worldwide Coverage Included
- HNOA Available
- Sexual Abuse Liability Available

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