# Clinical Trial Agreement

Date: _____

Parties:

1. [Pharmaceutical Company Name], hereinafter referred to as "Sponsor".

2. [Clinical Site/Institution Name], hereinafter referred to as "Site".

## 1. Purpose

The purpose of this Agreement is to define the terms and conditions under which the Site will conduct the clinical trial entitled "[Trial Title]".

## 2. Study Details

Study Protocol Number: _____

Study Start Date: _____

Study End Date: _____

## 3. Responsibilities

3.1. Sponsor's Responsibilities:

- Provide the investigational drug.
- Ensure compliance with regulatory requirements.

3.2. Site's Responsibilities:

- Conduct the study per the approved protocol.
- Recruit and enroll subjects.

## 4. Compensation

The Sponsor agrees to compensate the Site in accordance with the schedule outlined in Exhibit A attached hereto.

## 5. Confidentiality

Both parties agree to keep all proprietary information confidential as defined in Exhibit B.

## 6. Termination

Either party may terminate this Agreement upon written notice to the other party.

## Signatures

IN WITNESS WHEREOF, the parties hereto have executed this Clinical Trial Agreement as of the date first above written.

_____
[Name, Title]
[Pharmaceutical Company Name]

_____
[Name, Title]
[Clinical Site/Institution Name]