# **Clinical Trial Agreement**

Date: \_\_\_\_\_

Study Title: \_\_\_\_\_

This Clinical Trial Agreement (the "Agreement") is made and entered into by and between:

Institution: \_\_\_\_\_

Principal Investigator: \_\_\_\_\_

Sponsor: \_\_\_\_\_

## 1. Purpose

The purpose of this Agreement is to set forth the terms and conditions under which the clinical trial, titled "\_\_\_\_\_\_", will be conducted at multiple centers.

# 2. Study Sites

The multi-center study will be conducted at the following sites:

- Site 1: \_\_\_\_\_
- Site 2: \_\_\_\_\_

### 3. Responsibilities

#### Institution Responsibilities:

- Conduct the study in accordance with the protocol.
- Ensure compliance with regulatory requirements.

#### **Sponsor Responsibilities:**

- Provide necessary funding for the trial.
- Manage data collection and analysis.

## 4. Compensation

The Sponsor agrees to compensate the Institution as follows:

\_\_\_\_\_ (detailed terms).

# 5. Confidentiality

Both parties agree to maintain confidentiality of all proprietary information exchanged.

# 6. Term and Termination

This Agreement shall commence on the date first above written and shall continue until the completion of the study unless terminated earlier by either party upon written notice.

# 7. Signatures

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

Authorized Signatory for Institution

Authorized Signatory for Sponsor