

## Question 1

### Human Subjects Research

Please choose one of the groups below based on your subjects activities you will conduct. You will be assigned to a group.

- Biomedical Research:** This Basic Course is appropriate for research with human subjects.
- Social & Behavioral Research Investigators:** Choose this group to satisfy CITI training requirements for investigators and staff involved primarily in social and behavioral research with human subjects.
- IRB Members:** This Basic Course is appropriate for IRB or Ethics Committee members.

## Question 2

### Good Clinical Practice (GCP)

Good Clinical Practice (GCP) is an international ethical and scientific quality standard for designing, conducting, monitoring, auditing, reporting results, and disseminating the results of clinical trials involving human subjects.

It is a set of principles that are followed by:

• Investigators, sponsors, clinicians, regulatory agencies, and institutional review boards (IRBs) for clinical trials with investigational drugs and medical devices.

It includes:

- **Quality Management:** Ensuring that the trial is conducted in accordance with the protocol and regulatory requirements.
- **Documentation:** Maintaining accurate and complete records of all trial activities.
- **Monitoring:** Regularly checking the trial progress and data to ensure compliance with GCP.
- **Reporting:** Timely and accurate reporting of trial results to regulatory agencies.

## Question 3

### Responsible Conduct of Research

Please make a selection below to receive the course of Research.

Biomedical Research

Would you like to take the Responsible Conduct of Research course?

Yes

No

Conflicts of Interest (COI)

Would you like to take the Conflicts of Interest course?

- Yes
- No

## Question 5

### Public Health Research

Please make a selection below to receive the course of Public Health Research.

Would you like to take the Public Health Research course?

Yes

No