Becoming an Investigator

The advancement of medicine relies on dedicated clinical trial partners. If you are interested in working with Pfizer as an investigator, please send your contact details, institution or site name, and therapeutic areas of interest to clinicaltrials@pfizer.com

Investigator Responsibilities

- Protocol compliance
- Participant safety
- Informed consent
- Investigational product (IP) handling/accountability
- Staff training and oversight
- Communication with the IRB/IEC
- Timely completion of CRF documentation
- Communication with other facilities of the study site, such as laboratory and pharmacy
- Confidentiality requirements
- Resourcing

There are many roles involved in clinical trial duties and functions

- Investigator 01
- Sub Investigator 02
- Site coordinator/Study nurse 03
- Laboratory technician 04
- Referring physician(s) 05
- Other specialists 06
- Data coordinator 11
- Information Technology (IT) 10
- Pharmacist 09
- Regulatory 08
- Recruitment 07