

# Investigator Responsibilities

A clinical investigator is responsible for ensuring that an investigation is conducted according to the signed investigator statement, the investigational plan, and applicable regulations; for protecting the rights, safety, and welfare of subjects under the investigator's care; and for the control of drugs under investigation. The Clinical Investigator must also meet requirements set forth by the FDA, EMA or other regulatory body. The qualifications must be outlined in a current resume and readily available for auditors.



Protocol  
Compliance



Informed  
Consent



Good  
Documentation



GCP  
Compliance

Safety  
Reporting



Communication  
with  
IRB/IEC



IP  
Accountability



Adequate  
Resources

