Good Clinical Practice Guidelines (Dummy)

Section 1: Responsibilities

1.1 The Investigator should be qualified by education, training, and experience.

1.2 The Investigator should maintain a list of appropriately qualified persons to whom the in

Section 2: Safety Reporting

2.1 All serious adverse events (SAEs) should be reported immediately to the sponsor.

2.2 The immediate reports should be followed promptly by detailed, written reports.

2.3 Adverse events and/or laboratory abnormalities identified in the protocol as critical to sa

Section 3: Protocol Compliance

3.1 The investigator should conduct the trial in compliance with the protocol agreed to by th

3.2 The investigator should not implement any deviation from, or changes of the protocol w