

**Frequently Asked Questions and
Guidance on Clinical Research from
the Office for Human Research
Protections (OHRP)**

(April 1, 2019 - March 31, 2020)

**Frequently Asked Questions on
Clinical Research**

**Reviewing and Reporting Unanticipated
Problems Involving Risks to Subjects or
Others and Adverse Events**

**Guidance on IRB Continuing
Review of Research**

Guidance on Written IRB Procedures

**Children Involved as Subjects in Research:
Guidance on the HHS 45 CFR 46.407 “407”
Review Process**

Reporting Incidents to OHRP

**Engagement of Institutions in Human
Subjects Research**

**CFR Title 45, Part 46
Protection of Human Subjects
As of April 1, 2019**