Office of Human Research Protection
Institutional Review Board
701 Grove Rd
ESC158
Greenville, SC 29605

STATEMENT OF COMPLIANCE

Prisma Health has three operating Institutional Review Boards. All are structured to operate in compliance with United States regulatory requirements related to the protection of human research participants.

Prisma Health IORG and FWA #'s

FWA: 00001380

IORG: 0000483

Registered Committees:

Committee A: IRB# 00000805

Committee C: IRB# 00000875

Midlands Committee: IRB# 00013931

The Prisma Health Institutional Review Boards comply with HHS Regulations 45 CFR 46, and FDA regulations 21 CFR 50, 21 CFR 56, 21 CFR 312, 21 CFR 812. The Prisma Health Institutional Review Board operates in compliance with components of the Health Insurance of Portability Act of 1996 (HIPAA Privacy Rule) applicable human subject research, as defined in 45 CFR Parts 160 and 164. Guidelines of the International Conference on Harmonization (ICH) and Good Clinical Practice (GCP) are considered and applied when relevant.

If you have any questions or concerns, please feel free to contact the Prisma Health Office of Human Research Protection via e-mail at IRB@PrismaHealth.org.