My electronic signature indicates acknowledgment of the following terms:

- The Georgia CTSA Clinical Research Centers (GCRC) supports research participants aged 16 years and older, weighing ≥ 40 Kg., at EUH and EUHM. Special approval is required for participants <18 years old at other EHC sites.
- A valid speedtype is required for all studies.
- A monthly invoice will be generated for ongoing studies and the team has 7 days to raise inquiries or contest any charges. **If the GCRC does not receive communication from the PI within the stipulated period, invoice approval will be inferred.**
- Research studies will incur charges based on the current fee schedule listed on the Georgia Clinical and Translational Science Alliance website, which is approved by the Emory University School of Medicine. These fee schedules are subject to adjustments every two years.
- If there is a change in the fee schedule prior to the start of the study, the approved budget will be revised to reflect the updated fee schedule.
- Any modifications to the requested services originally outlined in the initial protocol/budget require a revision of the approved budget.
- Study teams are responsible for procuring necessary supplies pertinent to their studies, with a few exceptions (i.e., personal protective equipment, hand sanitizer, phlebotomy supplies including standard of care blood tubing and saline flushes, needles, syringes, blood glucose supplies, standard small urine collection cups, urine pregnancy tests, IV tubbing, absorbent pads, and biohazard waste bag).
- For space-only visits, should any staff assistance be requested whatsoever—be it from nursing, laboratory, coordinator, or bionutrition—a fee will be applied in accordance with the GCRC fee schedule.
- Prior to study initiation, the study teams must ensure proficiency in EPIC system usage, including order entry, preference list management, and order sets. Proficiency in CR-Assist is also required.