Mercer University Clinical Research Department is proud to offer a large number of clinical trials for patients and potential patients. Our researchers start new trials throughout the year, so the list is updated frequently. For questions regarding a specific clinical trial, please contact the Clinical Research Manager at 478-301-5846, and she will direct you to the appropriate study coordinator.

ENROLLING STUDIES:

- **DLS GCS 2018 – Suspected or Known Pneumonia**: The collection of biological specimen with companion data from subjects that will be catalogued in a biorespository for research.

- **FOURIER LEGACY**: Long-term Study of LDL-c Lowering with Evolocumab: Observational Follow-up after the FOURIER OUTCOMES Trial

- **GS-US-200-4334**: A Phase 2 Randomized, Open Label, Active Controlled Study Evaluating the Safety and Efficacy of Long-acting Capsid InhibitorGS-6207 in Combination with Tenofovir Alafenamide, AfterInducing Virologic Suppression with Emtricitabine/TenofovirAlafenamide in Antiretroviral Treatment-Naïve Adults with HIV-1Infection

- **HEART FID (1VIT15043)**: A Randomized, Double Blind, Placebo Controlled Study to Investigate the Efficacy and Safety of Injectafer® (Ferric Carboxymaltose) as Treatment for Heart Failure with Iron Deficiency

- **JANSSEN – DEFINE**: A Phase 4, Randomized, Active-Controlled, Open-label Study to Evaluate the Safety and Tolerability of Switching to Once-Daily Darunavir/Cobicistat/Emtricitabine/TenofovirAlafenamed (D/C/F/TAF) Fixed-dose Combination (FDC) Regimen in Virologically suppressed Human Innumodeficiency Virus Type 1 (HIV-1) Infected Participants Experiencing Rapid Weight Gain with an INI + TAF/FTC ARV Regimen

- **MK-8591A – 018**: A Phase 3 Randomized, Active-Controlled, Double-Blind Clinical Study to Evaluate a Switch to Doravirine/Islatravir (DOR/ISL) Once-Daily in Participants with HIV-1 Virologically Suppressed on Bictegravir/Entrivitabine/Tenofovir Alafenamide (BIC/FTC/TAF**)**

- **STEP/YES**: Youth Engagement & Sustained Care: Seek, Test, Treat and Retain for Youth and Young Adults Living with or at High Risk for Acquiring HIV

STUDIES IN FOLLOW UP:

- **CHAMP-HF**: Observational Registry of Treatment Patterns in US Heart Failure Patients with Reduced Ejection Fraction

- **GS-US-380-1490**: A Phase 3, Randomized, Double-Blind Study to Evaluate the Safety and Efficacy of GS- 9883/Emtricitabine/Tenofovir Alafenamide Versus Dolutegravir + Emtricitabine/Tenofovir Alafenamide in HIV-1 Infected, Antiretroviral Treatment-Naïve Adults

- **GS-US-380-4030**: A Phase 3, Randomized, Double-Blind Study to Evaluate the Safety and Efficacy of Switching from a Regimen of Dolutegravir and Either Emtricitabine/Tenofovir Alafenamide or Emtricitabine/Tenofovir Disoproxil Fumarate to a Fixed Dose Combination of Bictegravir/ Emtricitabine/Tenofovir Alafenamide in HIV-1 Infected Subjects who are Virologically Suppressed

- **GS-US-380-4580**: A Phase 3b, Multicenter, Open-Label Study to Evaluate Switching From a Regimen of Two Nucleos(t)ide Reverse Transcriptase Inhibitors (NRTI) plus a Third Agent to a Fixed Dose Combination (FDC) of Bictegravir/Emtricitabine/Tenofovir Alafenamide (B/F/TAF), in Virologically-Suppressed, HIV-1 Infected African American Participants

- **GS-US-412-2055**: A Phase 3, Randomized, Double-blind Study to Evaluate the Safety and Efficacy of Emtricitabine and Tenofovir Alafenamide (F/TAF) Fixed-Dose Combination Once Daily for Pre-Exposure Prophylaxis in Men and Transgender Women Who Have Sex with Men and Are At Risk of HIV-1 Infection

- **GSK-Atlas 2M (207966)**: A study to compare two drugs (called Cabotegravir and Rilpivirine, as oral tablets followed by long-acting injections) to current regimens containing 3 HIV drugs.

- **GSK-FLAIR (201584)**: A Phase III, Randomized, Multicenter, Parallel-group, Open-Label Study Evaluating the Efficacy, Safety, and Tolerability of Long-Acting Intramuscular Cabotegravir and Rilpivirine for Maintenance of Virologic Suppression Following Switch from an Integrase Inhibitor Single Tablet Regimen in HIV-1 Infected Antiretroviral Therapy Naive Adult Participants

- **GSK-POLAR (209035)**: A Phase IIb, Multicenter, Open-label, Rollover Study Evaluating the Efficacy, Safety and Tolerability of Long-acting Cabotegravir Plus Long-acting Rilpivirine Administered Every Two Months in HIV-1 infected Adults who areVirologically Suppressed and Participated in Study LAI116482

- **GSK-Tango (204862)**: A Phase III, randomized, multicentre, parallel-group, non-inferiority study evaluating the efficacy, safety, and tolerability of switching to dolutegravir plus lamivudine in HIV-1 infected adults who are virologically suppressed