



Proyecto ACTU

ESTUDIOS ABIERTOS A RECLUTAMIENTO	Criterios Inclusión Generales
A5369: HIV-1-Gag Conserved-Element DNA Vaccine (p24CE) as Therapeutic Vaccination in HIV-Infected Persons with Viral Suppression on Antiretroviral Therapy	➤ Diagnóstico VIH+ ➤ entre 18 y 65 años de edad. ➤ Conteo de CD4+ >500 cells/uL ➤ Carga Viral de VIH indetectable ➤ Estable en ART por los pasados 2 años
A5359: Long-Acting Antiretroviral Therapy in Non-adherent HIV-Infected Individuals	➤ Diagnóstico VIH+, ≥18 años de edad ➤ Poco adherencia a la terapia ART ➤ Carga Viral de VIH >200 copias/mL
A5375: An Open-Label, Phase II Pharmacokinetic Study to Evaluate Double-Dose Levonorgestrel Emergency Contraception in Combination with Efavirenz-Based Antiretroviral Therapy or Rifampicin-Containing Anti-Tuberculosis Therapy	➤ Diagnóstico VIH+ ➤ Mujeres entre ≥18 años de edad. ➤ Terapia ART estable que incluya una vez al día DTG 50 mg o EFV 600 mg
Próximos a Comenzar Reclutamiento:	
A5377: A Phase I, First-in-human, Ascending Dose Study of SAR441236, a Tri-specific Broadly Neutralizing Antibody in HIV-infected Participants.	➤ Diagnóstico VIH+, entre 18 y 70 años de edad. ➤ RAMA A: Conteo de CD4+ ≥200 cells/uL, carga Viral de VIH <50 copias/mL y estar en ART por los pasados 12 meses. ➤ RAMA B: ≥350 cells/uL y Carga Viral de VIH >500 y <100,000 copias/mL, No estar en ART
A5357: A Proof-of-Concept Study of Long-Acting Cabotegravir Plus VRC01LS to Maintain Viral Suppression in HIV-1-infected Adults.	➤ Diagnóstico VIH+, ≥18 años de edad. ➤ Conteo de CD4+ ≥350 cells/uL ➤ Carga Viral de VIH <50 copias/mL entre los pasados 2 años
A5368: "Safety, Pharmacokinetics and Immunotherapeutic Activity of an Anti-PD-1 Antibody in HBV Infected Participants on Suppressive Antiviral Therapy: A Phase I/II Ascending Multiple Dose Study"	➤ entre 18 y 70 años de edad. ➤ Mono Infección VHB
A5380: Glecaprevir/pibrentasvir Fixed-dose Combination Treatment for Acute Hepatitis C Virus Infection (PURGE-C)	➤ ≥18 años de edad. ➤ VHC co-infectado con VIH ➤ o VHC mono infectado
A5371: A Single-Arm, Open-Label, Pilot Study of Semaglutide for Non-Alcoholic Fatty Liver Disease (NAFLD), a Metabolic Syndrome with Insulin Resistance, Increased Hepatic Lipids, and Increased Cardiovascular Disease Risk (The SLIM LIVER Study)	➤ Diagnóstico VIH+ ➤ ≥18 años de edad. ➤ Conteo de CD4+ ≥200 cells/uL
A5379: HBV Vaccination in HIV-Infected Persons: Randomized, Controlled Trials of HEPLISAV-B vs. Engerix-B.	➤ Diagnóstico VIH+, Entre ≥18 y ≤70 años de edad, Conteo de CD4+ ≥100 cells/uL y HIV-1 RNA <1000



Futuros Estudios:

A5355: Phase I, Double-Blind, Randomized, Placebo-Controlled Trial to Evaluate the Safety, Tolerability, and Immunogenicity of a Modified Vaccinia Ankara (MVA)-based anti-CMV Vaccine (Triplex®), in Human Immunodeficiency Virus (HIV)-1 and CMV Co-Infected Adults who are on Potent Combination ART with Conserved Immune Function

A5363: A Multicenter, Prospective, Double-Blind, Randomized Placebo-Controlled Study to Evaluate the Effects of Cenicriviroc on Arterial Inflammation in People Living with HIV.

A5364: A Phase 1, Randomized, Open-Label, Study of the Safety, Pharmacokinetics and Ability to Durably Prevent Viral Relapse During a Monitored Analytical Treatment Interruption of a Combination of the Broadly Neutralizing Antibodies 3BNC117-LS and 10-1074-LS

A5371: A Single-Arm, Open-Label, Pilot Study of Semaglutide for Non-Alcoholic Fatty Liver Disease (NAFLD), a Metabolic Syndrome with Insulin Resistance, Increased Hepatic Lipids, and Increased Cardiovascular Disease Risk (The SLIM LIVER Study)

A5376: Pharmacokinetic Interactions of Etonogestrel Subdermal Implants (ENG) with Rilpivirine (RPV) Combined with Tenofovir Alafenamide (TAF) and Emtricitabine (FTC)

A5382: A Phase II Randomized Study Assessing the Safety and Efficacy of 24 Weeks of REP 2139-Mg, Followed by 48 Weeks of Combined Therapy with REP 2139-Mg and Pegylated Interferon Alpha-2a in Participants with HBeAg Negative Chronic HBV Infection Suppressed with Nucleos(t)ides

A5383: Randomized, Placebo-Controlled Trial to Evaluate the Safety of Letemovir (Prevymis) in Human Immunodeficiency Virus (HIV)-1 & CMV Co-Infected Adults who are on Suppressive ART & its Effect on Chronic Inflammation, HIV Persistence & Other Clinical Outcomes

A5386: IL-15 Superagonist (N-803) with and without Combination Broadly Neutralizing Antibodies

A5387: Two bNAbs (PGT121.BIJ414.LS and VRC07-523LS) in Combination with TLR9 agonist (Lefitolimod)

A5388: Combination HIV-Specific Broadly-Neutralizing Monoclonal Antibodies Combined with ART Initiation

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