



POSADA

RESEARCH CENTER

Precision in every detail.

A Coral Gables Site Built for *Precision & Partnership*

Multi-therapeutic Phase II–IV clinical research site • Coral Gables, Florida
Embedded in an established medical practice • Bilingual English / Spanish

INDEPENDENT

Direct access to site leadership and investigators.

MULTI-THERAPEUTIC

Eight outpatient therapeutic areas and growing.

PRACTICE-EMBEDDED

Pre-screening from an established adult patient base.

BILINGUAL

English–Spanish consent, visits and follow-up.

AT A GLANCE

- ◇ Board-certified Internal Medicine & Nephrology PI with 18+ years of Phase II–IV experience
- ◇ ACRP-certified site director (CCRC) with extensive academic cancer-center trial experience
- ◇ Team experience with AstraZeneca, Boehringer Ingelheim, Eli Lilly, Roche-Genentech and NCI cooperative-group trials
- ◇ Refrigerated and ambient centrifugation with a dedicated phlebotomy station
- ◇ Locking 2–8 °C pharmacy-grade IP refrigerator with continuous temperature logging
- ◇ Vital signs, standardized blood pressure, column scale with stadiometer
- ◇ Private sponsor monitoring room with workstation and printer
- ◇ Access to Miami-Dade's diverse, largely Hispanic adult population
- ◇ Minutes from Little Havana, Brickell and Downtown Miami

747 Ponce De Leon Boulevard, Suite 409 • Coral Gables, FL 33134

786-837-5463 | posadaresearchcenter.com

Feasibility questionnaires, site qualification requests and CDAs may be sent directly to site leadership.



SCAN TO CONNECT

Posada Research Center is an independent, multi-therapeutic clinical research site in Coral Gables, Florida, conducting outpatient Phase II–IV studies. Because the site operates inside an established medical practice, studies reach patients who are already in care, with bilingual staff, a single accountable operations lead and investigators who stay directly involved from feasibility to close-out.

SITE SNAPSHOT

LEGAL ENTITY	Posada Research Center LLC
ADDRESS	747 Ponce De Leon Boulevard, Suite 409, Coral Gables, FL 33134
MARKET	Miami-Dade County — Coral Gables, Little Havana, Coconut Grove, Westchester, Kendall, Doral
TRIAL PHASES	Phase II – Phase IV, outpatient
FACILITY	Practice exam rooms, research room, investigational product area, sponsor monitoring room
OWNERSHIP	Independent single site
IRB	Central IRB of record designated by the sponsor
LANGUAGES	English and Spanish — staff, consent (when IRB-approved) and participant materials

FACILITY & SPONSOR AMENITIES

- ◆ Private exam rooms for informed consent, physical exams and study visits
- ◆ Research room with lockable cabinets for regulatory binders and source documents
- ◆ Dedicated investigational product area with a locking 2–8 °C refrigerator
- ◆ Specimen collection and processing area with refrigerated and ambient centrifuges
- ◆ Private monitoring room with workstation, display and printer for CRA visits

PARTICIPANT SUPPORT

- ◆ Bilingual English–Spanish communication at every visit
- ◆ Pre-screening from the practice's records under HIPAA, with first contact made by the treating physician
- ◆ Flexible scheduling with visit windows tracked

RESEARCH TECHNOLOGY

- ◆ Sponsor EDC, IRT / IWRS (including Endpoint) and eCOA platforms
- ◆ SAE reporting, including FDA MedWatch 3500 and sponsor safety forms
- ◆ Electronic and paper regulatory binders



PRINCIPAL INVESTIGATOR

**Jorge L. Posada, MD****PRINCIPAL INVESTIGATOR • INTERNAL MEDICINE & NEPHROLOGY**

Board-certified in Internal Medicine and Nephrology, with more than 25 years of clinical, academic and medical-practice experience and more than 18 years as Principal Investigator and Sub-Investigator in Phase II–IV industry-sponsored trials. Fellowship-trained in Nephrology & Hypertension and in Transplant Nephrology.

Medical Director and Principal Investigator at Horizon Research Group (2009–2023), investigator at Veritas Research (2012–2020) and Principal Investigator at Professional Research Center since 2021. Personally oversees eligibility, informed consent, safety assessments, AE/SAE evaluation and protocol-required procedures, with hands-on EDC, IWRS and SAE-reporting experience and long working relationships with sponsors, CROs and monitors.

His Coral Gables internal medicine and nephrology practice houses the site, so pre-screening draws on an adult patient base he already treats.

RESEARCH FOCUS

Chronic kidney disease stages 1–5 · diabetic kidney disease · IgA nephropathy · FSGS · cystic kidney disease · dialysis and ESRD · anemia of CKD · hyperkalemia · hyperphosphatemia · uremic pruritus · type 2 diabetes · hypertension · ASCVD and hypercholesterolemia · heart failure · migraine · neuropathic pain

BOARD CERTIFICATION

Internal Medicine (2005)
Nephrology (2010)

FELLOWSHIPS

Transplant Nephrology — University of South Florida / LifeLink Foundation
Nephrology & Hypertension — University of Puerto Rico

RESIDENCY & DEGREES

Internal Medicine, University of Puerto Rico
(Chief Resident 2005–06)
MD — Universidad Autónoma de Centro América
BS Chemistry — University of Florida

APPOINTMENTS

Attending physician, Coral Gables Hospital (since 2015)
Former Assistant Professor of Internal Medicine, UPR School of Medicine

HOSPITAL AFFILIATIONS

Baptist Hospital of Miami · Jackson Health System · HCA Florida Kendall · South Miami Hospital · University of Miami Hospital · Larkin Community Hospital South

LICENSURE & TRAINING

Active Florida medical license · ECFMG certified · GCP (NIDA) · Human Subjects Protection (Advarra) · CITI HIPAA, AE reporting, informed consent

SOCIETIES • LANGUAGES

American Society of Nephrology · American Society of Transplantation
English · Spanish

SITE LEADERSHIP

**Christian Hernández, CCRC, CPT****DIRECTOR OF SITE OPERATIONS • CERTIFIED CLINICAL RESEARCH COORDINATOR (ACRP)**

Clinical Research Coordinator II in the Lymphoma and Hematology-Oncology program at Sylvester Comprehensive Cancer Center, University of Miami Miller School of Medicine, coordinating Phase 1–3 industry-sponsored, investigator-initiated and NCI cooperative-group trials. Took over a full caseload of more than 60 participants within his first three months and ran it on his own.

Earlier roles include industry-sponsored trials at Clinical Research of South Florida and research data and coordination at the University of Miami. Has handled every stage of a study: pre-screening, consent and re-consent, eligibility packets, visit windows, specimen processing and regulated shipping, EDC queries, deviations and CAPA, SAE reporting, IP accountability, monitoring visits, audits and close-out. Writes SOPs and templates and trains coordinators on GCP and protocol procedures. Serves as the site's single accountable contact from feasibility through close-out.

CERTIFICATIONS

Certified Clinical Research Coordinator (ACRP)
Certified Phlebotomy Technician (NHA)

EDUCATION

MS Data Science, Bioinformatics — University of Miami
BS Biological Science, Biotechnology — Miami Dade College

TRAINING

CITI GCP for Drugs & Biologics · CRC Foundations · Informed Consent · HIPAA · Shipping of Regulated Biological Materials · OSHA Bloodborne Pathogens

SYSTEMS

Velos · REDCap · sponsor EDC (Medidata, Medrio) · IRT / IWRS

LANGUAGES

English · Spanish

Additional board-certified investigators and sub-investigators are added by therapeutic area as protocols require. CVs, licenses, GCP certificates and financial disclosures are in the site qualification packet.

SUB-INVESTIGATOR



Yamilet Rodriguez, APRN, FNP

SUB-INVESTIGATOR • FAMILY NURSE PRACTITIONER — NEPHROLOGY

Family Nurse Practitioner and Registered Nurse with more than 25 years in healthcare, trained in medicine in Cuba and in nursing in the United States. Provides nephrology care in Dr. Posada's Coral Gables practice, which houses the site, including weekly follow-up of dialysis patients.

Performs delegated clinical assessments under Principal Investigator oversight: medical history review, physical examination, vital signs, phlebotomy and specimen collection, and documentation of adverse events and concomitant medications. Background in women's health (Pap smears, colposcopy), IV therapy, perioperative care, diagnostic sonography, EKG, Holter and pulmonary function testing. Bilingual English-Spanish.

LICENSURE

Florida APRN (Family Nurse Practitioner)
Florida Registered Nurse

EDUCATION

Master's, Family Nurse Practitioner — Miami Regional University (2025)
MD — Facultad de Ciencias Médicas "Dr. Enrique Cabrera," Havana (1997)

CLINICAL FOCUS

Nephrology and dialysis · cardiometabolic care · women's health

LANGUAGES

Spanish · English

SELECTED TRIAL EXPERIENCE — DR. POSADA

AREA	STUDIES
Renal · CKD · Dialysis	FLOW semaglutide vs placebo on kidney outcomes in T2D with CKD · NEUTRALIZE (AstraZeneca) Phase IIIb sodium zirconium cyclosilicate in hyperkalemia with metabolic acidosis · Boehringer Ingelheim 1378-0005 BI 690517 ± empagliflozin for UACR reduction · RAINIER (NCT06564142) Phase 3 povetacicept in IgA nephropathy · PROACT / REACT (NCT05099770) Phase 3 renal autologous cell therapy in T2D with CKD · Vertex VX19-NEN-801 APOL1 in proteinuric kidney disease · Vadadustat Phase 2-3 programs in anemia of CKD, including INNO2VATE · Tenapanor Phase 3 in hyperphosphatemia · CR845 pruritus in hemodialysis · Kibow Biotech Phase 2 in stage IV CKD · FSGS proteinuria studies · Phase III IV-iron comparisons in iron-deficiency anemia
Diabetes · Cardiovascular · Metabolic	CAROLINA Phase III cardiovascular safety in T2D · FIGARO / FIDELIO cardiovascular and renal outcomes in diabetic kidney disease · Cardiovascular Inflammation Reduction Trial low-dose methotrexate in CAD with T2D or metabolic syndrome · injectable lipid-lowering therapy in ASCVD and hypercholesterolemia · chronic heart failure mortality outcomes · pediatric T2D fixed-dose combinations · basal insulin initiation in insulin-naïve T2D
Neurology · Pain	SPARTAN and GLADIATOR acute treatment of migraine · diabetic peripheral neuropathic pain · pseudobulbar affect · relapsing multiple sclerosis · chronic low back pain and opioid-induced constipation · pediatric pain PK
Other	Uncomplicated urinary tract infection · hereditary cancer gene-panel validation · transdermal irritation and sensitization

TRIAL PORTFOLIO — SITE DIRECTOR

AREA	STUDIES COORDINATED OR SUPPORTED AT SYLVESTER COMPREHENSIVE CANCER CENTER
Lymphoma · Hematology	LOTIS-7 (ADC Therapeutics) loncastuximab tesirine combinations · Corvus Phase 3 soquelitinib in relapsed T-cell lymphoma · Genentech GO43878 Phase III glofitamab in mantle cell lymphoma · Nurix NX-5948 Phase 1 BTK degrader · AstraZeneca PRIMAVERA Phase I/II AZD3470 · BeiGene zanubrutinib combinations · Ono Phase II tirabrutinib in PCNSL · Genmab epcoritamab in marginal zone lymphoma · Eli Lilly J2N-MC-JZNX
Cooperative group · IIT	Alliance A051901 · SWOG S2308 · RADAR Phase III Hodgkin lymphoma · Dana-Farber DLBCL 22-606 · MSK 18-160 and 21-176 · NCI ETCTN · investigator-initiated studies across Phase 1-3

SPONSOR EXPERIENCE OF OUR TEAM

AstraZeneca	Boehringer Ingelheim	Eli Lilly	Roche · Genentech	Vertex	Genmab
BeOne Medicines	ADC Therapeutics	Corvus	Nurix	Ono Pharmaceutical	NCI · Alliance · SWOG

Trials conducted by our investigator and site director at prior sites and institutions. Listing does not imply a contract between these sponsors and Posada Research Center.

04 What Sets Us Apart

Access to patients, accountable leadership and data you can trust.



- ◆ **Patients already in care**
Recruitment starts from an established practice population, not cold advertising, which shortens time to first patient in.
- ◆ **Diverse, bilingual enrollment**
Spanish-speaking staff and IRB-approved Spanish materials help sponsors meet FDA diversity-plan goals in Miami-Dade.
- ◆ **One accountable contact**
An ACRP-certified site director owns feasibility, start-up, conduct and close-out with no layers of hand-offs.
- ◆ **Honest feasibility**
We estimate enrollment from our real patient population and say so when a protocol is not a fit.
- ◆ **Multi-therapeutic by design**
Cardiometabolic, respiratory, vaccine, liver, kidney and gout programs run side by side under one quality system.
- ◆ **Active investigator oversight**
Physicians are directly involved in eligibility, safety and sponsor communication, not delegated away.
- ◆ **Academic-grade coordination**
Coordination experience from an NCI-designated cancer center: complex eligibility, amendments, SAEs and IP accountability.
- ◆ **Sponsor-ready quality systems**
ALCOA+ source documentation, ICH E6 (R3) GCP conduct, temperature and delegation logs maintained on site.

START-UP PATHWAY

01 Feasibility Protocol review with an honest fit and enrollment assessment.	02 Site qualification On-site or remote SQV, staff and facility walkthrough.	03 Contract & IRB Regulatory documents, budget review, CTA negotiation and central IRB submission in parallel.	04 Activation SIV, training, delegation log and regulatory binder complete.	05 First patient in Pre-screening starts at qualification, not at activation.
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Our service commitments to sponsors and CROs

Turnaround targets are confirmed per study and written into the clinical trial agreement.

FEASIBILITY

Prompt, data-based response

START-UP PACKET

Single coordinated submission

DATA ENTRY

Timely after each visit

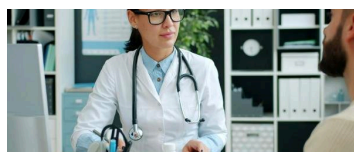
QUERIES

Resolved by the site director

THERAPEUTIC AREAS

Cardiovascular Hypertension · Lp(a) · HF risk	Diabetes & Endocrinology Type 2 diabetes · prediabetes	Obesity Weight management	Vaccines & ID RSV · influenza · COVID-19
Respiratory Asthma · COPD · chronic cough	Liver & Metabolic MASH · fatty liver	Kidney Disease CKD · DKD · IgA · ADPKD	Gout & MSK Gout · osteoarthritis pain

Cardiometabolic overlap. Many adults in the practice live with two or more of these conditions (diabetes, hypertension, excess weight, early kidney disease), so one pre-screened population can support several programs at once.



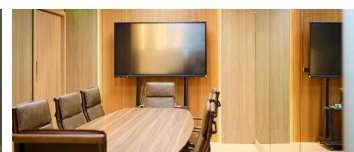
Study visits



Specimen handling



Clinical assessments



Sponsor visits

Representative images.

Clinical environment

- ◇ Private exam rooms within the medical practice
- ◇ Mobile vital signs monitor (BP, pulse, temperature)
- ◇ Wall-mounted automated BP monitor plus aneroid backup
- ◇ Column scale with stadiometer and digital floor scales
- ◇ Private space for informed-consent discussions

Laboratory & specimen operations

- ◇ Dedicated phlebotomy station
- ◇ Refrigerated centrifuge and ambient centrifuges
- ◇ Bench-top laboratory incubator
- ◇ Specimen preparation, aliquoting, labeling and packaging
- ◇ Central-lab kit receipt, inventory and expiry tracking

Pharmacy & investigational product

- ◇ Dedicated investigational product area
- ◇ Pharmacy-grade 2–8 °C refrigerator with locking door
- ◇ Secondary medical refrigerator for kits and supplies
- ◇ Accountability, dispensing and reconciliation records, including IRT

Temperature monitoring

- ◇ Wireless data logger with buffered probe in IP storage
- ◇ Traceable min/max thermometer (Fridge-tag 2L)
- ◇ Daily review and documented temperature logs
- ◇ Written excursion process: quarantine, sponsor notification, resolution

EQUIPMENT INVENTORY

CATEGORY	EQUIPMENT	NOTES
Vitals	Welch Allyn vital signs monitor on mobile stand	On site
Blood pressure	Welch Allyn wall-mounted automated monitor; aneroid sphygmomanometer	On site
Anthropometric	Seca column scale with stadiometer; digital floor scales	On site
Specimen collection	Phlebotomy chair with adjustable arms	On site
Lab processing	Eppendorf refrigerated centrifuge	On site
Lab processing	LW Scientific E8 clinical centrifuge	On site
Laboratory	Boekel bench-top incubator	On site
IP storage	Kool-It Ikon pharmacy refrigerator, 2–8 °C, locking door, digital display	On site · continuous monitoring
Cold storage	Accucold medical refrigerator	On site
Monitoring	Wireless temperature data logger, buffered probe, Fridge-tag 2L	On site · daily review
Sponsor visits	Monitoring room with workstation, display and printer	On site

Protocol-specific equipment (for example frozen storage or 12-lead ECG) can be sourced and qualified during study start-up. Calibration and maintenance records are available for review at the site qualification visit.

Study operations

- ◇ Feasibility, site qualification and start-up
- ◇ Regulatory binder and essential documents
- ◇ Contract and budget coordination
- ◇ Visit scheduling with protocol windows tracked
- ◇ Source documentation and timely EDC entry
- ◇ Query resolution and monitoring-visit support
- ◇ Close-out and record retention

Quality, regulatory & data

- ◇ ICH E6 (R3) GCP conduct
- ◇ 21 CFR Parts 50, 56 and 312 compliance
- ◇ ALCOA+ source documentation
- ◇ Delegation and training logs
- ◇ Deviation, CAPA and SAE processes
- ◇ HIPAA-protected participant records
- ◇ Central IRB submissions and reporting

Recruitment & retention

- ◇ Pre-screening from the practice's patient records
- ◇ Treating-physician first contact under HIPAA
- ◇ IRB-approved bilingual materials
- ◇ Community outreach in Miami-Dade
- ◇ Reminder calls and flexible scheduling
- ◇ Interest list for upcoming studies

DOCUMENTS AVAILABLE ON REQUEST

- ◇ Investigator CVs, medical licenses and GCP certificates
- ◇ Site director CV and ACRP certification
- ◇ Signed CDA and site feasibility questionnaire
- ◇ Equipment list and calibration records
- ◇ Temperature monitoring logs
- ◇ Facility photographs and floor information

Let's discuss your next study.

Send a protocol synopsis or feasibility questionnaire and we will reply with an honest assessment of fit and enrollment potential for our population.

PRIMARY CONTACT

Christian Hernández, CCRC – Director of Site Operations
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747 Ponce De Leon Blvd, Suite 409, Coral Gables, FL 33134

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