ALTUS CLINICAL RESEARCH SOUTH FLORIDA

Women's health. Broader expertise.

Independent Phase II-IV research site | Palm Beach County, South Florida

Altus Clinical Research combines an established foundation in women's health with study experience in metabolic conditions, weight management, dermatology and general medicine. Our research model connects clinical trial operations with community physician relationships.

INVESTIGATOR LEADERSHIP

Samuel N. Lederman, MD, FACOG
Medical Director & Principal Investigator
Obstetrics & Gynecology | Clinical research leadership dating to 1996
Medical Director of Altus since 2006

CORE RESEARCH FOCUS	ADDITIONAL EXPERIENCE & INTEREST
Women's health	Obesity and weight management
Menopause and hot flashes	Diabetes and metabolic conditions
PCOS and reproductive health	General medicine
Contraception	Dermatology and psoriasis
Urinary incontinence	Tinea pedis and onychomycosis
HPV and vaginal health	

AN INTEGRATED APPROACH TO STUDY DELIVERY

Experienced clinical research coordinators, regulatory and recruitment support, and community-based physician relationships support protocol assessment and study execution. Patient safety, protocol requirements and data quality guide our approach.

LEADERSHIP	SITE OPERATIONS
Jessica Ravelo, MS, CCRP President 561-248-0001 jessica@altusresearch.com	Yanelys Perez Site Director 561-257-3277 yperez@altusresearch.com
Samuel Lederman, MD Medical Director & Principal Investigator sleder@altusresearch.com	

4671 S. Congress Avenue, Suite 100-B, Lake Worth, FL 33461
altusresearch.com

Investigator experience by indication

Samuel N. Lederman, MD, FACOG | Clinical research experience

STUDY LISTINGS IN THE PRINCIPAL INVESTIGATOR CV

THERAPEUTIC CATEGORY	CV LISTINGS	THERAPEUTIC CATEGORY	CV LISTINGS
Contraception	31	Menopause symptoms	23
Atrophic vaginitis	8	Osteoporosis	16
Overactive bladder / incontinence / urinary disorders	19	Genital infections	14
Uterine fibroids / endometriosis / menstrual disorders	18	Pap smear studies	9
Weight loss / obesity	7	Psoriasis	14
Dermatology	9	Tinea pedis / onychomycosis	7

ADDITIONAL INVESTIGATOR EXPERIENCE

The CV also documents vaccine research, anemia, cardiovascular and lipid disorders, hypertension, gastrointestinal conditions, osteoarthritis, respiratory disorders and diagnostic sample-collection studies. This experience supports assessment of protocols beyond the core women's health portfolio.

SCIENTIFIC EXPERIENCE

The supplied CV lists publications in menopause, osteoporosis and vulvovaginal candidiasis, including co-authored research published in The Lancet and The New England Journal of Medicine. Investigator credentials and indication-specific experience are available for feasibility review.

ADDITIONAL PIPELINE INDICATIONS

PCOS; hot flashes; vaginal atrophy; urinary incontinence associated with obesity; HPV / Pap testing; menopause with weight management; and yeast infection research.

Current studies and follow-up

Enrollment and follow-up by indication

SPONSOR	INDICATION	GOAL	Screened	RANDOMIZED / ENROLLED	GOAL REACHED	STATUS
A*****	Weight loss	10	3	NR	NR	Enrolling
M*******	HPV	50	47	47	94.0%	Enrolling
E********	Obesity / weight management	14	46	36	257.1%	Enrolling
E********	Urinary incontinence	28	14	11	39.3%	Enrolling
M******	Oral contraception	10	13	4	40.0%	Follow-up
M****	Birth control	20	18	11	55.0%	Follow-up
R****	TV/CV testing - study 1	200	270	270	135.0%	Follow-up
R****	TV/CV testing - study 2	150	260	260	173.3%	Follow-up
R**	OB sample collection	50	59	59	118.0%	Follow-up
R****	Type 2 diabetes mellitus (T2DM)	50	56	42	84.0%	Follow-up
G******	Healthy volunteers	100	220	220	220.0%	Follow-up
R****	HPV	100	285	285	285.0%	Follow-up
R**** / C*********	STI	100	192	192	192.0%	On pause
C*****	Obesity with T2DM and metabolic disease	15	11	9	60.0%	Follow-up

NR = not reported

A practical partner for your protocol

Study-specific feasibility, operational assessment and direct access to site leadership

SITE RESOURCES

CLINICAL & PARTICIPANT RESOURCES	STUDY OPERATIONS
Clinical practice relationships	Clinical trial management and EDC experience
Patient examination rooms	Regulatory and recruitment support
Laboratory and phlebotomy capabilities	Restricted-access medication storage
Ultrasound, DEXA, colposcopy and EKG	Temperature-controlled storage
Uroflow capabilities	Monitoring space and written SOPs

START WITH A FEASIBILITY DISCUSSION

Please share the study synopsis or feasibility questionnaire, including key eligibility criteria, enrollment expectations, recruitment window and required assessments. Our team can evaluate investigator fit, relevant study experience, recruitment approach and operational requirements.

CONTACT OUR TEAM

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