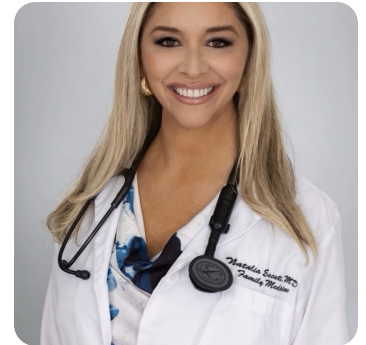


Natalia Escuti, M.D.

Sub-Investigator | Integrated Health Research LLC

Clinical Research Operations and Family Medicine

Miami-Dade County, Florida | 818-481-2248 | docnjescutti@gmail.com



PROFESSIONAL PROFILE

Dr. Escuti brings family medicine practice and clinical research operations experience to Integrated Health Research. Her CV documents study management roles with AbbVie, ICON Clinical Research and Precision for Medicine; work as a physician investigator and medical monitor; and residency leadership in inpatient and outpatient care. She reports proficiency in English, Spanish and French.

CLINICAL AND PHYSICIAN INVESTIGATOR EXPERIENCE

Larkin Community Hospital, Palm Springs Campus, Hialeah | Chief Physician, Family Medicine Residency Program - Inpatient | July 2022-June 2024

Clinical experience across inpatient services, ICU, cardiology, emergency medicine, pediatrics, obstetrics and gynecology, and other rotations. Participated in grand rounds and weekly didactics.

Isla Morada Medical Center, Tavernier | Chief Physician, Family Medicine Residency Program - Outpatient | July 2022-June 2024

Operated a family medicine clinic providing primary and urgent care, preventive care, and outpatient procedures. CV reports approximately 15-20 patients per day, including pediatrics and obstetrics and gynecology.

Plasma Med Research, LLC, Boca Raton | Physician Investigator and Medical Monitor (Consultant) | August 2015-March 2018

Served as substitute Principal Investigator for consent and physical examinations and followed participants during studies. Performed medical monitoring focused on participant safety, protocol adherence and data.

EDUCATION

M.D. | Ross University School of Medicine | June 2014

Medical degree.

B.A., Psychology | California State University | June 2006

Undergraduate degree.

A.S., Biology | Citrus College | June 2004

Associate degree.

CLINICAL RESEARCH OPERATIONS

AbbVie Inc., Chicago | Study Manager Associate II (Remote) | May 2020-May 2022

Oversaw planning and execution of clinical studies across phases and geographies, including site selection and startup, recruitment, monitoring, data retrieval, database lock and closeout. Contributed to risk mitigation and cross-functional plans.

ICON Clinical Research, North Wales | Clinical Trial Manager (Consultant, Remote) | March 2018-May 2022

Managed studies from startup through closeout. Reviewed protocol synopses, informed consent forms, case report forms and study plans. Oversaw clinical operations teams, vendor interactions, safety planning, site performance, budgets and quality standards.

Precision for Medicine, Bethesda | Clinical Trial Manager (Remote) | June 2014-February 2018

Managed timelines, clinical research associates and site issues. Contributed to protocol and case report form drafting, clinical validation studies across several regions, database lock, and quality assurance initiatives.

University of Miami, Neurology Research Department | Clinical Research Coordinator / Medical Writing Intern | July 2014-June 2015

Supported protocol review and submissions, case reports, manuscript review and a poster presented at the 2015 American Headache Society conference. Assisted with protocol design for an intracranial hypertension study.

THERAPEUTIC AREAS DOCUMENTED IN THE SOURCE CV

Women's health, dermatology, endocrine conditions, psychiatry, respiratory and cardiovascular conditions, vaccinations, rheumatology and neurology.

SELECTED STUDY EXPERIENCE

- Phase 3 upadacitinib induction and maintenance study in Crohn's disease (2020-2022).
- Phase 3 ozanimod induction, maintenance and open-label extension studies in Crohn's disease (2018-2022).
- Clinical evaluation of a preeclampsia diagnostic test using ACOG 2013 reference criteria (2017-2018).
- NGAL test for risk assessment of acute kidney injury in an intensive care population (2017).

SELECTED STUDY EXPERIENCE, CONTINUED

- Randomized pilot study of digital interventions for cognitive dysfunction in adults with major depressive disorder (2016-2017).
- Phase 3 study of an anticholinergic agent added to an antispasmodic in irritable bowel syndrome (2015-2017).

TRAINING AND SKILLS

Good Clinical Practice, NIDA Clinical Trials Network - last update July 6, 2024.

Languages: English, Spanish and French.

Office software: Microsoft Word, PowerPoint and Excel.

CREDENTIALS IN THE SOURCE CV

The 2024 source CV lists a medical doctor license with expiration January 2026 and ACLS, BLS and PALS certificates with expiration April 2026. DEA registration was listed as processing. Current status has not been independently verified and should be updated before sponsor submission.

Experience and study descriptions above are based on the physician-provided 2024 CV. Her IHR role is shown for the requested site profile; protocol delegation, licensure, GCP training and institutional requirements should be confirmed for each study.