

Kendra Rininger, NP-C

Sub-Investigator

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EDUCATION

2003-2006	Indiana University – Indianapolis, IN Master of Science in Nursing – Family Nurse Practitioner
1993-1997	Indiana State University – Terre Haute, IN Associates of Science in Nursing & Bachelor of Science in Nursing
1989-1993	Northview High School – Brazil, IN High School Diploma

MEDICAL WORK EXPERIENCE

11/2025-Present	Sub-Investigator The Indiana Clinical Trials Center – Plainfield, IN
12/2024-Present	Nurse Practitioner/Injector/Aesthetics The Dermatology Center of Indiana/Optima Dermatology – Plainfield, IN
01/2024-12/2024	Nurse Practitioner/Injector/Aesthetics Her MD – Carmel, IN
11/2020-01/2024	Nurse Practitioner/Injector/Aesthetics The Dermatology Center of Indiana/Optima Dermatology – Plainfield, IN
10/2019-10/2020	Nurse Practitioner IU Health Physicians – Indianapolis, IN

LICENSURE AND CERTIFICATION

Nurse Practitioner License:

- License Number: 71002428A
- NPI Number: 1528264322

Registered Nursing License:

- License Number: 28131799A

AANP Family Nurse Practitioner Certification

- Date: 09/2026 – Present
- Certification Number: F0906092

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Certifications:

11/2025-Present	Good Clinical Practice Certification (GCP)
10/2025-Present	ICH Good Clinical Practice (GCP) Training R3
11/2025-Present	Protecting Human Research Participants Certification
12/2023-Present	Basic Life Support (BLS)
11/2021-Present	Laser Certification
09/2021-Present	Chemical Peels/PCA
09/2012-Present	Neurotoxin and Dermal Filler

CURRENT TRIALS

2025 – Present	Randomized, Multi-Center, Evaluator-Blind, Vehicle-Controlled Study to Evaluate Efficacy and Safety of Reformulated Levulan Kerastick plus Photodynamic Therapy (PDT) for Field-Directed Treatment in Patients with Actinic Keratosis (AK) of Upper Extremities
2025 – Present	Randomized, Multi-Center, Evaluator-Blind, Vehicle-Controlled Study to Evaluate Efficacy and Safety of Reformulated Levulan Kerastick plus Photodynamic Therapy (PDT) for Field-Directed Treatment in Patients with Actinic Keratosis (AK) of Upper Extremities
2025 – Present	COVE-2: A Phase 3, Double-Blind, Randomized, Vehicle-Controlled Study to Evaluate the Efficacy and Safety of YCANTH (VP-102) in Subjects with Common Warts (Verruca Vulgaris)
2026 – Present	COVE-4: A Phase 3, Open-Label, Long-Term Follow-Up Study to Evaluate the Safety and Efficacy of YCANTH (VP-102/TO-208) in Subjects with Common Warts (Verruca Vulgaris)

I have signed my CV to acknowledge that it is accurate, complete, and current, as of the date of signing.

Signed: Kendra Ruzinger NP-C Date: 02 JUN 2026