

# Christina Lynne Race, PA-C

Dawes Fretzin Clinical Research Group, LLC  
7910 North Shadeland Avenue  
Indianapolis, Indiana 46256  
Phone (317) 516-5030  
Fax (317) 516-5148  
E-mail: christinadfcrg@gmail.com

Dawes Fretzin Dermatology Group, LLC  
7910 North Shadeland Avenue  
Indianapolis, Indiana 46256  
Phone (317) 516-5000  
Fax (317) 516-5146

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## Education

Butler University (Indianapolis, IN)  
Bachelor of Health Science/Physician Assistant, 2000

- Dean's List
- Graduated with Honors

## Professional Experience

**Dawes Fretzin Clinical Research Group, LLC** (Indianapolis, IN)  
*Sub-Investigator, 2004-Present*

**Dawes Fretzin Dermatology Group, LLC** (Indianapolis, IN)  
*Physician Assistant, 2000-Present*

**Indiana Blood Center/Tissue Bank** (Indianapolis, IN)  
*Donor Procurement, 1999-00*

**Methodist Hospital Emergency Department** (Indianapolis, IN)  
*Emergency Room Technician, 1997-98*

## License/Certifications

Indiana State Physician Assistant License 10000471A  
NCCAA 1045335  
CPR/AED Certified  
ACLS Certified  
TB Certified  
Adjunct Faculty Butler University  
CITI Certification, March 2010, May 2012, February 2014, February 2015, February 2016, February 2017, February 2018, January 2021

## Professional Societies

Membership to the American Academy of Physician Assistants  
Membership to the Society of Dermatology Physician Assistants  
Membership to the Indiana Academy of Physician Assistants  
House of Delegates Student Representative for Physician Assistant organization (Butler University)  
Fundraising chair for Butler University Methodist Physician Assistant organization (Butler University)

*C. Race PA C  
01/22/22*

# **Christina Lynne Race, PA-C**

## **Psoriatic Arthritis**

Sponsor: Janssen Research and Development

A Phase 2a, Multicenter, Randomized, Double-blind, Placebo-controlled Study Evaluating the Efficacy and Safety of Guselkumab in the Treatment of Subjects With Active Psoriatic Arthritis April 2015 to Present

Sponsor: Janssen Research and Development

A Multicenter, Randomized, Double-blind, Placebo-controlled Trial of Golimumab, an Anti-TNF $\alpha$  Monoclonal Antibody, Administered Intravenously, in Subjects with Active Psoriatic Arthritis August 2014 to September 2015

Sponsor: Novartis Pharmaceuticals

A phase III, randomized, double-blind, placebo-controlled multicenter study of subcutaneous secukinumab in autoinjectors, to demonstrate efficacy at 24 weeks and to assess the long term safety, tolerability and efficacy up to 3 years in subjects with active Psoriatic Arthritis June 2014 to Present

## **Atopic Dermatitis**

Sponsor: Pfizer

A Phase 2B Randomized, Double-Blind, Placebo-Controlled, Parallel, Multicenter, Dose-Ranging, Study to Evaluate the Efficacy and Safety Profile of PF-04965842 in subjects with Moderate to Severe Atopic Dermatitis. April 2016- Present

Sponsor: GlaxoSmithKline

A Randomized, Blinded, Vehicle-Controlled, Dose-Finding Study of GSK2894512 Cream for the Treatment of Atopic Dermatitis - February 2016 to Present

Sponsor: MedImmune, Limited

A Phase 2a, Randomized, Double-blinded, Placebocontrolled Study to Evaluate the Efficacy and Safety of MEDI9929 in Adult Subjects with Moderate-to-Severe Atopic Dermatitis August 2015 to September 2016

Sponsor: Regeneron Pharmaceuticals

A Phase 3, Randomized, Double-Blind, Placebo-Controlled Study Investigating the Efficacy and Safety of Multiple Dupilumab Dose Regimens Administered as Monotherapy for Maintaining Treatment Response in Patients with Atopic Dermatitis July 2015 to February 2017

Sponsor: Regeneron Pharmaceuticals

An Open-Label Study of Dupilumab In Patients with Atopic Dermatitis Who Participated In Previous Dupilumab Clinical Trials - June 2015 to Present

Sponsor: Regeneron Pharmaceuticals

A phase 3 Confirmatory Study Investigating the Efficacy and Safety of Dupilumab Monotherapy Administered to Adult Patients with Moderate-to-Severe Atopic Dermatitis February 2015 to June 2016

Sponsor: Regeneron Pharmaceuticals

A Randomized, Double-Blind, Placebo-Controlled, Study Investigating Vaccine Responses in Adults with Moderate to Severe Atopic Dermatitis Treated with Dupilumab September 2014 to December 2015

# **Christina Lynne Race, PA-C**

## **Actinic Keratosis**

Sponsor: Therapeutics, Inc

A Multicenter, Randomized, Double-Blind, Vehicle-Controlled, Parallel Group Comparison Study to Determine the Therapeutic Equivalence of a Generic Ingenol Mebutate Gel, 0.015% and Picato Gel, 0.015% in Subjects with Actinic Keratosis on the Face or Scalp July 2016 to Present

Sponsor: LEO Pharma A/S

Efficacy and Safety of Ingenol Mebutate Gel in Field Treatment of Actinic Keratosis on Full Face, Balding Scalp or Approximately 250 cm<sup>2</sup> on the Chest – *May 2015 to Present*

Sponsor: Perrigo UK Finco Limited Par

A Multi-Center, Double-Blind, Randomized, Vehicle-Controlled, Parallel-Group Study to Compare Perrigo UK Finco Ingenol Mebutate Topical Gel 0.015% to Leo Pharma Inc. Picato® Topical Gel 0.015% (Ingenol Mebutate Topical Gel 0.015%), and Both Active Treatments to a Vehicle Control in the Treatment of Actinic Keratosis on the Head Region (Face or Scalp) March 2015 to December 2015

Sponsor: Leo Pharma A/S

Safety of LEO 43204 0.018%, 0.037% and 0.1% for actinic keratosis applied once daily for three consecutive days on face/chest, scalp and trunk/extremities, respectively – *February 2015 to September 2015*

## **Common Warts**

Sponsor: RXi Pharmaceuticals

A Prospective, Phase 2a Study to Evaluate the Effectiveness and Safety of DPCP Ointment (Samcyprone) on the Clearance of Verruca Vulgaris (Common Warts) in Subject Ages 18-65 Years. July 2016 – Current