

January 2021

Curriculum Vitae
William Allen Bethea, DO
Joined January 2015



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Personal Data: **Advanced Dermatology and Cosmetic Surgery**
725 West Granada Blvd., Suite #44
Ormond Beach, FL 32174

Leavitt Medical Associates of Florida d/b/a Ameriderm Research
725 West Granada Blvd., Suite #44
Ormond Beach, FL 32174
Phone: 386-523-0768
Fax: 386-366-9212
Email: drwbethea@leavittmgt.com

Physician **January 2015-Present** **Ormond Beach, FL**

- Advanced Dermatology and Cosmetic Surgery
- Leavitt Medical Associates of Florida d/b/a Ameriderm Research
- Physician/Sub- Investigator

Residency: **July 2010 - December 2014** **Trenton, MI**

- Oakwood Southshore Hospital/Michigan State University Consortium
- Chief Resident July/2012-2013

Military Service: **July 2006 – June 2010** **MacDill AFB, FL**

- United States Air Force, Flight Surgeon
- Rank: Major Select

Internship: **July 2005 - June 2006** **Largo, FL**

- Sun Coast Hospital
- Traditional Internship

Medical Education: **August 2001 – May 2005** **Davie, FL**

- Nova Southeastern University
- D.O.

Undergraduate Education: **August 1997 – May 2000** **Gainesville, FL**

- University of Florida
- B.S., Entomology

Certificates/Licensure: **Licensed Doctor of Osteopathic Medicine**

- Florida License OS 9887
- ACLS/BLS
- CITI Good Clinical Practice (Expires June 2020)

Affiliations:

- American Osteopathic College of Dermatology
- Michigan Dermatological Society
- American Osteopathic Association

Publications/Presentations:

- Publications
 - Bethea WA. Brooke-Spiegler Syndrome
Journal of the AOCD; 2011
Oct, Vol 21, No 151-2
 - Bethea WA. Case Report: Vaccinia Vaccine Major Reaction.
Medical Corps Examiner
July 2008
- Presentations
 - Drug Hyperpigmentation
American Osteopathic College of Dermatology
Mid-Year Meeting, 2013
 - Michigan Dermatological Society Meeting, 2011
 - Graft Versus Host Disease
Michigan Dermatological Society Meeting, 2012

Accomplishments:

- Health Professions Scholarship Program
- Military
 - Meritorious Service Award
 - National Defense Service Medal
 - Afghanistan Campaign Medal
 - Iraq Campaign Medal
 - Global War on Terrorism Medal
 - Air Force Expeditionary Service Ribbon
 - NATO Medal
- University of Florida Men's Volleyball Team

Volunteer Experience:

- Space Shuttle STS-131 (Discovery) medical support team, 4/2010
- Hurricane Ike disaster relief team, 8/2008
- Director of smoking cessation program 3/2007
- Summer camp medical provider 6/2006

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Research Studies:

Efficacy and Safety of xxx in field Treatment of Actinic Keratosis on Full Face, Balding Scalp or Approximately xxx on the chest

Efficacy and Safety of xxx in field Treatment of Actinic Keratosis on Approximately xxx on Trunk or Extremities

Efficacy and Safety of xxx in Field Treatment of Actinic Keratosis on Balding Scalp including 12-month follow-up

A Multi-Center, Randomized, xxx-Blind, Parallel-Group, xxx Controlled Study To Compare The Efficacy and Safety of xxx Versus xxx In Subjects With Acne Vulgaris

A Phase II, Randomized, Double-Blind, xxx-Controlled Study to Evaluate The Safety and Efficacy of xxx In Patients With Persistent Moderate To Severe Atopic Dermatitis That Is Inadequately Controlled By Topical Corticosteroids

A Multicenter, Randomized, xxx-Blind Study Comparing the Efficacy and Safety of Xxx Dosing Regimens in Patients with Moderate-To-Severe Plaque Psoriasis

A Phase 3 Multi-Center, Randomized, Double-Blinded, xxx-Controlled, Parallel Group Study Comparing the Efficacy, Tolerability and Safety of xxx and xxx Once Daily in the Treatment of Acne Vulgaris

A Phase 2B Randomized, Double-Blind, xxx-Controlled, Parallel, Multicenter, Dose-Ranging, Study to Evaluate the Efficacy and Safety Profile of xxx in Subjects with Moderate to Severe Atopic Dermatitis

Xxx Versus xxx and xxx comparators in a randomized double blind for Maintenance use in Moderate to severe plaque type psoriasis – 2 (xxx)

Multicenter, Randomized, Double-Blind, xxx-Controlled, Phase 2A Study of xxx Tablets in Androgenetic Alopecia in Males with a xxx Arm

A Multicenter, xxx, Double-Blind, xxx-Controlled Study to Assess The Safety And Efficacy of xxx in Treatment-Resistant Puritus Associated with Atopic Dermatitis

Efficacy and Safety of xxx in Field Treatment of Actinic Keratosis on Face or Chest including 12-month follow-up/Phase 3

A Multicenter, xxx label study to assess the safety and efficacy of xxx for maintenance in Moderate to severe plaque type psoriasis/Phase 3

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A Phase 3, Randomized, Double-Blind, xxx-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of xxx versus xxx in Subjects 12 years of age or older with Cutaneous Common Warts

A Randomized, Multicenter, Double-blind, Vehicle-controlled Study to Evaluate the Safety and Efficacy of xxx --% Topical xxx Foam Compared to Vehicle in the Treatment of Facial Papulopustular Rosacea

A Randomized, Double-Blind, Vehicle-Controlled Study to Evaluate the Efficacy and Safety of Topical Administration of xxx for 12 Weeks in the Treatment of Moderate-To-Severe Acne Vulgaris/Phase 3

A Randomized, Double-Blind Dose-Ranging Study of xxx xxx Cream in Subjects With Vitiligo

A Phase 2, Randomized, Dose-Ranging, Vehicle-Controlled and xxx xx% Cream-Controlled Study to Evaluate the Safety and Efficacy of xxx Cream Applied Topically to Adults With Atopic Dermatitis

An Open-Label Study to Evaluate the Long-Term Safety of Topical Administration of xxx for 40 weeks in the Treatment of Moderate to Severe Facial Papulopustular Rosacea.

A Phase 3, Double-Blind, xxx-Controlled, Randomized, Parallel Group, Multicenter, Efficacy and Safety Study of xxx Ointment 1% in Adult Subjects with Actinic Keratosis on the Face of Scalp

A Phase 3, Multicenter, Randomized, xxx-Blind, xxx-And Active Comparator-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety xxx in Adult Subjects With Moderate to Severe Chronic Plaque Psoriasis

A Phase 3, Multicenter, xxx-label Study to Assess the Long-Term Safety, Tolerability, and Efficacy of xxx in Adult Subjects with Moderate to Severe Chronic Plaque Psoriasis

A Multicenter, Randomized, xxx-Blind, xxx-Controlled, Parallel-Group Study to Evaluate the Efficacy and safety of xxx in Adult Subjects with Moderate To Severe Chronic Plaque Psoriasis

A Phase 3 Randomized, XXX-Controlled, XXX-Blind Study to Evaluate XXX in Adolescent and Adult Subjects with Moderate to Severe Atopic Dermatitis

A Multicenter, Randomized, Double-blind, XXX-controlled, Phase 3 Efficacy and Safety Study of XXX, x%, for the Reduction of Disease Burden of Persistently Developing Basal Cell Carcinomas (BCCs) in Subjects with Basal Cell Nevus Syndrome

Phase 2B, Randomized, Double-blind, xxx Controlled, Parallel Group, xxx Ranging Study to Assess Efficacy, Safety, Tolerability and Pharmacokinetics of xxx Applied Once or Twice Daily for 6 Weeks in Participants with Mild or Moderate Atopic Dermatitis

Phase 2B, Randomized, Double-blind, xxx Controlled Parallel Group, xxx Ranging Study to Assess Efficacy, Safety, Tolerability and Pharmacokinetics of xxx Applied Once or Twice Daily for 12 Weeks in Participants with Mild to Moderate Chronic Plaque Psoriasis

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Phase 3, Randomized, Double-Blind, xxx-Controlled, Multicenter Study Investigating the Efficacy and Safety of xxx Co-Administered with Background Medicated xxx Therapy in Adolescent Participants 12 to <18 Years of Age with Moderate to Severe Atopic Dermatitis.

Phase 2, Randomized, Double-blind, xxx-Controlled Study to Evaluate the Safety and Efficacy of xxx in Participants with Moderate to Severe Plaque Psoriasis.

A multicenter, randomized, xxx-blind, xxx-controlled, 24-week study, with a 28-week open-label extension, to assess the safety and efficacy of xxx in subjects with moderate-to-severe alopecia areata

A Phase 3, Randomized, xxx-Blind, xxx-Controlled, Multicenter Study Evaluating The Efficacy and Safety of xxx in Study Participants with Moderate to Severe Hidradenitis Suppurativa

A Phase II, Randomized, xxx-Blind, xxx-Controlled Dose-Ranging Study To Evaluate The Efficacy And Safety of xxx In Subjects With Non-Segmental Vitiligo

A Randomized, xxx-Blind, xxx-Controlled Study to Assess the Efficacy and Safety of xxx (xxx) in Subjects with Moderate-to-Severe Atopic Dermatitis

A Randomized, xxx-Blind, xxx-Controlled Study to Assess the Efficacy and Safety of xxx (xxx) in Subjects with Prurigo Nodularis

A Prospective, Multicenter, xxx-Term Study to Assess the Safety and Efficacy of xxx (xxx) in Subjects with Prurigo Nodularis (PN)

A Phase 3, Multicenter, Open-Label Extension Study of xxx Topical Gel, xxx in Subjects with Gorlin Syndrome (Basal Cell Nevus Syndrome).

A xxx-Blind, xxx-Controlled, Randomized Withdrawal and Treatment Extension Study to Assess the Long-Term Efficacy and Safety of xxx Cream in Participants With Vitiligo