Curriculum Vitae

James A. Solomon, MD, PhD

Chief Dermatology, North Las Vegas VA Medical Center
Director, Koch's Postulates Dermatology Consulting
Professor Dermatology, University Central Florida, College of Medicine
Professor University of Nevada Las Vegas School of Medicine
Assistant Clinical Professor, Carle-Illinois, College of Medicine
Editor, Journal Medical Internet Research Dermatology
Assistant Editor, Journal American Academy Dermatology

Diplomate of American Board of Dermatology, December 1983
State of Florida, ME 91126, 8/03/2004 – current
NV Med License 26355 09/05/24-currrent

Clinical and Research Office:

Las Vegas VA Medical Center
6900 North Pecos Road,
North Las Vegas, NV 89086.
Office: V: 702.791.9000,15214
James.solomon@va.gov
james.solomon@ucf.edu
james.solomon@unlv.edu
avibenzv@illinois.edu

Consulting Office (Preferred telephone number and email address)

Koch's Postulates Dermatology Consulting
8317 Aqua Spray Ave
Las Vegas, NV 89128
M: 217 649 5504
F: 702 779 0727
jasolomonmdphd@kochspostulates.com

Education:

Academic

June 1976 - June 1979
SUNY at Buffalo School of Medicine
3435 Main Street, Buffalo, NY
MD

June 1972 - May 1976
SUNY Upstate Medical Center
766 Irving Avenue, Syracuse, NY
PhD – Immunology

September 1968 - May 1972
Johns Hopkins University
3400 N Charles Street, Baltimore, MD
BA: Natural Sciences

Fellowships
September 1978 - June 1979
Massachusetts General Hospital, Boston, MA
Research Fellow in Medicine
January 1977 – August 1978
Roswell Park Memorial Institute
Louis Sklarow Memorial, Buffalo, NY
Dermatology/Oncology Research Fellowship
July 1972 - June 1976
SUNY, Upstate Medical, Syracuse, NY
National Institutes of Health Training Fellowship

Residency
July 1980 - June 1983
University Hospital, Boston, MA
Dermatology

Internship
November 1979 - June 1980
Boston City Hospital, Boston, MA
Pathology
July 1979 - October 1979
New England Medical Center Hospital, Boston, MA
Pediatrics

Work History
April 2023-Present
Las Vegas VA Medical Center
North Las Vegas , NV
Chief Dermatology
July 2021- Present
Koch's Postulates Dermatology Consulting
Ormond Beach, FL
Consultant
January 2009 – March 2023
Leavitt Medical Associates of FL, Inc. d/b/a)
Ameriderm Research, Ormond Beach, FL
Director
January 2006 - 2008
Leavitt Medical Associates of FL, Inc. d/b/a
Research Division, Advanced Dermatology Cosmetic Surgery, Ormond Beach, FL
Director
August 2004 - 2023
Leavitt Medical Associates of FL, Inc. d/b/a
Advanced Dermatology and Cosmetic Surgery, Ormond Beach, FL

Dermatologist
August 2001 - July 2004
Decatur Memorial Hospital
2300 N Edwards, Decatur, IL
Dermatologist
January - July 200
Host, "Perfect Skin", Talk radio Sunday 11am WNNZ 640 am.
March 1990 - July 2001
Aesthetique Skin International/James A Solomon MD PhD PC
171 Dwight Road, Longmeadow, MA 01106
Owner
February, 1986 - February, 1990
Medical West Community Health Plan, Chicopee, MA
Chief Dermatology
February 1985 - January 1986
Medical West Community Health Plan, Chicopee, MA
Staff Dermatologist
July 1983 - February 1985
Boston University School of Medicine
Department of Dermatology, Boston, MA
Assistant Research Professor

Teaching and Staff Appointments:

2023-Present
Professor, University of Nevada Las Vegas, Kirk Kerkorian School of Medicine
2019-present
Assistant Clinical Professor, Dermatology Carle-Illinois College of Medicine
2018-2023
Adjunct Professor Dermatology, Kansas City University of Medicine & Biosciences
2017- 2023
Faculty & Primary Advisor Residents Research Activities. Kansas City University of Medicine & Biosciences- GME Consortium. Advanced Dermatology & Cosmetic Surgery (ADCS) Orlando Dermatology Residency
2017- present
Clinical Professor Dermatology, Florida State University, College of Medicine
2016 - present
Professor of Dermatology, University of Central Florida, College of Medicine, Department of Medical Education
2007-2015
Associate Professor of Dermatology, University of Central Florida, College of Medicine, Department of Medical Education
2001-2019
Assistant Clinical Professor, Dermatology University Illinois Urbana-Champaign, College of Medicine
2004-2008
Teaching and Courtesy Staff, Dermatology Department, Halifax Hospital, Daytona Beach, FL

August 2001 — 2004
Consulting Staff, Decatur Memorial Hospital, Decatur IL
2001— 2004
Assistant Clinical Professor, Dermatology, Southern Illinois University, School of Medicine, Springfield, IL
1987 - 2001
Assistant Clinical Professor of Dermatology, Tufts University School of Medicine, Boston IL
1985 - August 2001
Associate Staff, Dermatology, Baystate Medical Center, Springfield, MA
1985 - 1991
Courtesy Staff, Dermatology, Holyoke Hospital, Holyoke, MA
February 1985 - August 2001
Courtesy Staff, Dermatology, Mercy Hospital, Springfield, MA
February 1985 - 1986
Adjunct Assistant Professor, Department Microbiology, Boston University Medical School, Boston, MA
July, 1983 - February, 1985
Consultant in Dermatology, Lemuel Shattuck Hospital, Boston, MA
July, 1983 - February, 1985
Assistant Visiting Physician in Dermatology, Boston City Hospital, Boston, MA
July, 1983 - February, 1985
Assistant Visiting Physician in Dermatology, University Hospital, Boston, MA
July, 1983 - February, 1985
Assistant Research Professor of Dermatology, Boston University School of Medicine, Boston, MA
July, 1981 - June, 1983
Research Instructor, Boston University School of Medicine, Boston, MA

Certifications

CITI Program course: **Human Research, VA Human Subjects Protection, 2-Refresher Course**
Course under requirements of **Las Vegas [VA] NV-593**
Completion Date 21 June 2023 Expiration Date 21 June 2025
CITI Program course: **Biosecurity, VA ORD Biosecurity Training, 1- Basic Course**
Course under requirements of **Las Vegas [VA] NV-593**
Completion Date 21 June 2023 Expiration Date n/a
CITI Program course: **Financial Disclosure and Conflicts of Interest for PHS Investigators 1 – Stage 1.**
Course under requirements of **University of Illinois at Urbana- Champaign**
Completion Date 06-March-2021 Expiration Date 05-March-2024
CITI Program course: **Core IRB Training 1. Basic Course.**
Course under requirements of **University of Illinois at Urbana- Champaign**
Completion Date 06-March-2021 Expiration Date 05-March-2024
CITI Program course: **Biomedical Responsible Conduct of Research Course 1. RCR**
Course under requirements of **University of Illinois at Urbana- Champaign**
Completion Date 06-March-2021 Expiration Date 05-March-2024
CITI Program course: **Human Subjects Research -Group 1. Biomedical Research Investigators and Key Personnel 1. Basic Course**
Course under requirements of **University of Central Florida**
Completion Date 05 March 2021 Expiration Date 04 March 2024
CITI Program course: **Investigators (Biomedical) 1. Basic Course**
Course under requirements of **Advarra IRB**
Completion Date 05-March-2021 Expiration Date 05-March-2023

CITI Program course: **Additional Groups in the Course for The Protection of Human Subjects** (Curriculum Group) Investigators (Biomedical) 2 - Refresher Course Under requirements of **Advarra IRB**
Completion Date 16-May-2019 Expiration Date 15-May-2023
CITI Program course: **CITI Good Clinical Practice CITI** 1 - GCP Refresher Course Under requirements of **Advarra IRB**
Completion Date 16-May-2019 Expiration Date 15-May-2023
CITI Program course: **Good Clinical Practice Course for Clinical Trials Involving Drugs (ICH focus)** 2 - Refresher Course
Under requirements of **Kansas City University of Medicine and Biosciences**
8/7/18 valid until 8/7/20
CITI Program course: **Good Clinical Practice Course for Clinical Trials Involving Drugs** (ICH focus) (Curriculum Group) **GCP for Clinical Trials with Investigational Drugs and Biologics** (ICH Focus) (Course Learner Group) 1 - **Basic Course**
Under requirements set by: **Kansas City University of Medicine and Biosciences**
Completion Date 21-May-2018 Expiration Date 20-May-2020
CITI Program course**: Human Research** (Curriculum Group) **Biomedical Investigators** (Course Learner Group) 3 - **Refresher Course** (Stage).
Under requirements set by: **Kansas City University of Medicine and Biosciences**
Completion Date 21-May-2018 Expiration Date 20-May-2021
CITI Program course: **Conflict of Interest 1- Stage 1.**
Course under requirements of **University of Central Florida**
Completion Date 21-May-2018 Expiration Date 20-May-2022
CITI Program course**: Human Research** (Curriculum Group) Group 1.**Biomedical Research Investigators and Key Personnel** (Course Learner Group) 4 - **Refresher Cours**e (Stage)
Under requirements set by: **University of Central Florida**
Completion Date: 21-May-2018 • Expiration Date: 20-May-2021
CITI Human Research: • **Curriculum Group: Human Research** • Course Learner Group: Group 1. **Biomedical Research Investigators and Key Personnel** • Stage: Stage 4 - **Refresher Course** • Record ID: 26079449 •
Institution Affiliation: **University of Central Florida** (ID: 405)
Completion Date: 21-May-2018 • Expiration Date: 20-May-2020
CITI Program course**:** • (ID: 443)• Curriculum Group: **Human Research** • Course Learner Group: Group 1. **BIOMEDICAL RESEARCH** • Stage: Stage 3 - Refresher Course • Record ID: 22221517 •
Institution Affiliation: **Western IRB**
Completion Date 21-May-2018 Expiration Date 20-May-2021
CITI Human Research: (ID: 829) • Curriculum Group: CITI Health Information Privacy and Security (HIPS) • Course Learner Group: **CITI Health Information Privacy and Security (HIPS) for Clinicians** • Stage: Stage 1 - **Basic Course** • Record ID: 23461056 •
Institution Affiliation: **Kansas City University of Medicine and Biosciences**
Completion Date 20-Mar-2018 Expiration Date 19-Mar-2020
CITI Program course: **CITI Health Information Privacy and Security (HIPS)** (Curriculum Group) **CITI Health Information Privacy and Security (HIPS) for Clinical Investigators** (Course Learner Group) 1 - **Basic Course** (Stage)
Under requirements set by: **Kansas City University of Medicine and Biosciences**
Completion Date: 07/09/2017 • Expiration Date: 07/28/2021
American AED/CPR Association Certificate Healthcare Provider CPR Course
Advanced Dermatology & Cosmetic Surgery, Moh's Microscopic Controlled Surgery Training
1997
Certificate, Clinical Use of the C02 Laser in Cosmetic Skin Resurfacing, Luxar Corporation

1980
Diplomat, National Board of Medical Examiners

Honors, Honorary Societies:

2019
University of Central Florida College of Medicine Fire Research Mentor Award
2011
American Academy of Dermatology, AAD Leadership Circle
For Volunteerism, Gold Level
Jan 2007
American Academy of Dermatology, AAD Leadership Circle
For Volunteerism, Bronze Level
May 2007
President's Volunteer Service Award, USA Freedom Corps
1990 — 2004
Physician Recognition Award, American Medical Association
1972
With Honors, Johns Hopkins University
1971
Alpha Epsilon Delta, Johns Hopkins University,

Professional Societies:

Pediatric Dermatology Research Alliance (PeDRA) -2021-present
American Society Clinical Oncology 2020-2022
American Hair Research Society– 2017 present
Acne Core Outcome Network (ACORN) 2015- present
Group for Research & Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA), Member, 2013- present
International Dermatology Outcome Measures (IDEOM) Member–2013-present
International Dermatoepidemiology Association (IDEA), member 2010-present
American DermatoEpidemiology Network (ADEN), Member 2009- present.
Central Florida Society of Dermatology 2009-present
American Federation for Medical Research, Member, 2006- present
American Federation for Clinical Research, Member, 1982 — 1985
Society for Investigative Dermatology, Member, 2006- present
Society for Investigative Dermatology, Member, 1983 – 2000
Society for Investigative Dermatology, Resident— Fellow Member, 1982 -1983
Association of Clinical Research Professionals, Member, 2006 – present
American Society for Laser Medicine and Surgery, Fellow 2002- present
National Psoriasis Foundation, Member, 1988 - present
American Academy of Dermatology, Fellow, 1983 - Present
Massachusetts Medical Society, Member, 1983-Present
Massachusetts Medical Society, Resident Member, 1981 — 1983
Royal Society of Medicine (United Kingdom), Member 2011- 2013
Society of Clinical Research Associates, Member, 2006-2008
American Society of Cosmetic Dermatology and Aesthetic Surgery-member 2002- 2010
Illinois State Medical Society; Member 2001-2004
Macon County Medical Society; Member 2001 —2004
International Society for Hair Replacement Surgery, Member, 1998- 2001
Royal Society of Medicine (United Kingdom), Member-1998 — 2001
Society for Pediatric Dermatology, 1991 - 2001
American Contact Dermatitis Society, 1990 - 2001

Member, Quinsigamond Dermatologic Society (University of Massachusetts Medical School, Department Dermatology), 1990-2001
American Dermatologic Society for Allergy and Immunology, Member,
1985 - 2004
American Medical Association, Member, 1983 –2001
American Medical Association, Resident Member, 1979—1983
American Medical Association, Student Member, 1977—1979
American Association for the Advancement of Science, 1983 — 1985
Massachusetts Academy of Dermatology, 1983 - 1985
Hubert H. Humphrey Cancer Center, Member, 1981 —1985

National, State, and Local
Committee Appointments:
Editor, Journal Medical Internet Research, Dermatology 2021-present
Assistant Editor, Journal American Academy of Dermatology 2021-present
Reviewer, International Journal Dermatology 2019-present
Member of Patient Advocate Task Force, American Academy Dermatology 2018-2020
Consultant, Mayne Pharma, 2018- present
Consultant Aspyrian Therapeutics Inc, 2018-present
Reviewer, New England Journal of Medicine 2018- present
Reviewer, American Journal of Clinical Dermatology 2018 present
Reviewer, European Journal Pharmacology 2018-present
Reviewer, Springer Current Dermatology Reports 2018- present
Reviewer, Case Reports in Dermatologic Medicine 2017- present
Consultant, Sun Pharmaceutical, 2017- present
Consultant, Eli Lilly 2017 – present
Reviewer, British Journal Dermatology 2016-present
Reviewer, Flagship: Medical Scholarly Proceedings UCF COM, 2016- present
Member of Kansas City University of Medicine and Biosciences Scholarly Activity & Faculty Development Committee Meeting Oct 2016- present
ACORN communications section leader 2016- present
Reviewer, Journal Dermatological Treatment, 2013- present
Reviewer, JAMA –Dermatology 2013- present
Scientific Advisory Board, National Alopecia Areata Foundation, 2013- present
American Dermatoepidemiology Network (ADEN) / Epidemiology Expert Resource Group, webmaster 2011- present
Alopecia Areata Clinical Research Task Force, National Alopecia Areata Foundation, 2010- present
Epidemiology Expert Resource Group (Epi-ERG) - American Academy of Dermatology 2010- present
Reviewer, Journal American Academy of Dermatology, 2009 – Present
Scientific Advisory Board, Samumed, 2016
Member of the Clinical Guidelines Committee of the AAD. 2015-2017
Member, Scientific Advisory Board of Basal Cell Carcinoma Nevus Syndrome Life Support Network 2014-2016
Erivedge Dermatologist Advisory Board, Genentech, 2013.
Scientific Advisory Board, Xenoport, 2013
Assistant Editor, Journal American Academy of Dermatology, 2013-2018
Hedgehog Inhibitor & Advanced Basal Cell Carcinoma Advisory Board, Eli Lilly, 2012
American Academy of Dermatology, Epidemiology Expert Resource Group (Epi-ERG), & American Dermatoepidemiology Network Cooperative Effort, Chair, pSOAR (psoriasis Outcomes Assessments Review), 2012-2014
Epi-ERG/ ADEN, subcommittee to establish Dermato-epidemiology Clinical Trials Resource Cooperative, Chair– 2011- 2014

Co-editor, Cutaneous Oncology Today, 2011 to 2013
Advanced Basal Cell Carcinoma Regional Advisory Board, Genentech, 2010
S-5 Curriculum Dermatology Section Coordinator, University of Central Florida, College of Medicine, 2008-2010
Subcommittee S-1 Curriculum Development for Hematology Oncology, University of Central Florida, College of Medicine 2008- 2009
Subcommittee S-6 Curriculum Development for Muscular Skeletal (Cutaneous) Systems, University of Central Florida College of Medicine, 2008-2009
American Society for Laser Medicine and Surgery, Constitution and Bylaws Committee, 2007-2008
Graduate Medical Education Committee, Halifax Hospital, Member 2007-2008
Curriculum Committee, University of Central Florida, College of Medicine, 2007-2008
American Academy of Dermatology, EPA/NIEHS/NIOSH Liaison Task Force, Member 2003-2007
Illinois State Medical Society, Governmental Affairs Council: Member, 2002 — 2004
Massachusetts Medical Society: Delegate, 1999-2002
Hampden District Medical Society: Executive Board, 1999 —2002
Contact Dermatitis Committee of American Academy of Dermatology, January 1994-March 1998.

Community Service:

Jewish Federation of Volusia & Flagler Counties, Florida, Board, member, 2005 – Present
Co-chairman, Maimonides Medical Society, Jewish Federation of Volusia & Flagler Counties, FL 2004 – Present
Vignette Judge, 2008 Annual Research Symposium, University of Illinois College of Medicine at Urbana at Urbana Champaign, April 2008
Member, Economic Prosperity Committee, Ormond Beach Chamber of Commerce, 2006 - 2016
Member, Education Committee, Ormond Beach Chamber of Commerce, 2006 - 2016
Port Orange/South Daytona Chamber of Commerce & Economic & Government Affairs Committee
Subcommittee Chair: Health and Medical Issues. 2005-2012
Co-chairman, Israeli Bond Drive, Champaign, IL 2002— 2004
Community Relations Committee, Springfield Jewish Federation, 2000-2002
Community Relations Committee, Springfield Jewish Federation, 1995
Lubavitcher Yeshiva Academy, Board Member, Long Range Planning Committee Member, 2000— 2001
Beth Israel Synagogue, Board Member, Director of Membership - 1995-1996
Member, Young Leadership Development Program, Springfield Jewish Federation - 1990

Medical/Legal and Business Consulting:

Consultant for Vista Society of Industry Leaders 2006- present
Consultant for MedaCorp 2006- present
Consultant for Network of Experts 2005- present
Consultant for Network of Advisors, Clinical Advisors, LLC 2004 – Present
Consultant for J. Reckner Associates, Inc. 2004 – Present
Consultant for Doctor Directory 2004-present
Gerson Lehrman Group, Councils of Advisors, 2001- present
Expert Witness in areas of routine & cosmetic dermatology, hypersensitivity, immunology, and connective tissue diseases, as well as Product Liability Merit Analysis for various law firms in Massachusetts, Michigan, New York, Pennsylvania, South Carolina 1990— present
Independent Medical Examiner, Department of Industrial Accidents, Commonwealth of Massachusetts, August 1995—2001
Physician Member Medical Malpractice Tribunal, Commonwealth of Massachusetts, 1994—2001
Product Liability Merit Analysis, *Sedwick, Detert, Moran, & Arnold,* NY, NY, June 1994 - 2001

Merit Analysis Malpractice & Product Liability, Central & Western Mass Lawyers, March 1990 —2001
Trial expert witness in areas of dermatology, hypersensitivity, immunology, and connective tissue disease, *Gary .Lenehan, Esq,* Manchester, NH, 1988-89.

Grant Awards:

Beg S, Solomon JA, Kolpashchikov D, Ramshaw B, Stancescu M, Updyke KM, Lewis S, Domozych R, Stines M. Identifying the genetic sequence and source of the DNA antigenic component of the anti-DNA immune complexes in systemic lupus erythematosus (SLE) in order to establish the role of exogenous antigens evoking molecular mimicry response in human SLE disease pathogenesis. University of Central Florida College of Medicine Study Pilot Laboratory Study Internal Grant 2014-2015.

Clinical Trials:

*A Phase 2 Double-Blind, xxx-Controlled, Randomized Withdrawal and Treatment Extension Study to Assess the Long-Term Efficacy and Safety of xxx Cream in Participants With Vitiligo - 2021
*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 - 2021
*A Phase 3 Prospective, Multicenter, xxx-Term Study to Assess the Safety and Efficacy of xxx (xxx) in Subjects with Prurigo Nodularis (PN) - 2021
*Evaluation of Clinical Outcomes in Patients with Cutaneous xxx Lesions Tested with A Differential Diagnostic xxx Expression Assay - 2020
*A Phase 2 Randomized, xxx-Blind, xxx-Controlled Study to Assess the Efficacy and Safety of xxx (xxx) in Subjects with Moderate-to-Severe Atopic Dermatitis - 2020
*A Phase 3 Randomized, xxx-Blind, xxx-Controlled Study to Assess the Efficacy and Safety of xxx (xxx) in Subjects with Prurigo Nodularis - 2020
*A Phase 3, Multicenter, xxx-Label Extension Study of xxx Topical Gel, xxx in Subjects with Gorlin Syndrome (Basal Cell Nevus Syndrome) - 2020
*Gorlin Syndrome Data Analysis
*A Phase 2 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 - 2020
*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 - 2020
*Phase 2, Randomized, Double-blind, xxx-Controlled Study to Evaluate the Safety and Efficacy of xxx in Participants with Moderate to Severe Plaque Psoriasis - 2019
*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 - 2019
*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 - 2019
*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 - 2019

*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-2018

*A Multi-Center, Randomized, Double-Blind, XXX- and XXX Comparator-Controlled
Phase 3 Study to Evaluate the Efficacy and Safety of XXX in Subjects with Moderate-to-Severe Plaque Psoriasis - 2018
*A Phase 3 Randomized, XXX-Controlled, XXX-Blind Study to Evaluate XXX in Adolescent and Adult Subjects with Moderate to Severe Atopic Dermatitis - 2018
* A Phase 3, Multicenter, Randomized, xxx-Blind, xxx-And Active xxx-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety xxx in Adult Subjects With Moderate to Severe Chronic Plaque Psoriasis - 2018
* 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 - 2018
* 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 - 2018
* A Randomized, xxx-Blind, xxx-Controlled, Efficacy Study of the xxx Receptor Antagonist xxx in Patients with Atopic Dermatitis - 2018
* A Randomized, xxx-Blind, Multicenter Study Assessing Short (16 weeks) and Long-Term Efficacy (up to 1 year), Safety, and Tolerability of Sub-Cutaneous xxx in Subjects of Body Weight xx kg or xx with Moderate to Severe Chronic Plaque-Type Psoriasis - 2018
* A Phase Ib, xxx-Blind, Randomized, xxx-Controlled Study to Assess the Safety, Tolerability, and Pharmacokinetics of Single and Repeated Doses of xxx - 2018
* A Randomized, xxx-Blind, xxx-Controlled, Phase 2B Study to Evaluate the Efficacy, Safety, Tolerability, Pharmacokinetics of xxx in Subjects with Moderate to Severe Atopic
Dermatitis - 2018
*Phase 3 A Multicenter, xxx label study to assess the safety and efficacy of xxx for maintenance in Moderate to severe plaque type psoriasis - 2017
*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 - 2017
*****Phase 3 A Randomized, Multicenter, Double-blind, xxx-controlled Study to Evaluate the Safety and Efficacy of xxx --% Topical xxx Foam Compared to Vehicle in the Treatment of Facial Papulopustular Rosacea - 2017
*Phase 3 Randomized, Double-Blind, xxx-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 - 2017
*Phase 2 Randomized, Double-Blind Dose-Ranging Study of xxx Cream in Subjects With Vitiligo - 2017
*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 - 2017
*Phase 3 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. - 2017
*****Phase 3 Double-Blind, xxx-Controlled, Randomized, Parallel Group, Multicenter,
Efficacy and Safety Study of xxx xxx X% in Adult Subjects with Actinic Keratosis on the Face of Scalp - 2017
*Phase 3 Efficacy and Safety of xxx in Field Treatment of Actinic Keratosis on Balding Scalp including 12-month follow-up 2016
*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 2016

*A Phase 3 Multi-Center, xxx Label Study Evaluating the Long Term Safety of xxx Once Daily in the Treatment of Acne Vulgaris 2016
*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 2016
*Phase 3 Xxx Versus xxx and xxx comparators in a randomized double blind for Maintenance use in Moderate to severe plaque type psoriasis – 2 (xxx) 2016 – Phase III
*Phase 2A Multicenter, Randomized, Double-Blind, xxx-Controlled, Phase 2A Study of xxx Tablets in Androgenetic Alopecia in Males with a xxx Arm - 2016
*Phase IIa A Multicenter, xxx, Double-Blind, xxx-Controlled Study to Assess The Safety And Efficacy of xxx in Treatment-Resistant Pruritus Associated with Atopic Dermatitis - 2016
*Phase 3 Efficacy and Safety of xxx in Field Treatment of Actinic Keratosis on Face or Chest including 12-month follow-up - 2016
*A Multicenter, Randomized, xxx-Blind Study Comparing the Efficacy and Safety of Xxx Dosing Regimens in Patients with Moderate-To-Severe Plaque Psoriasis
SCORING Trial: Phase IIb Open-label Trial of xxx in Subjects with Basal Cell Carcinoma Nevus Syndrome, 2015
*Phase 3 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 2016
*A Phase 3, Multicenter, Randomized, xxx-Blind, Paralled-Group, Study Followed By a Dose-Blind Period and Open-Label Follow-Up To Evaluate the Efficacy and Safety of xxx In Subjects With Moderate to Severe Chronic Plaque Psoriasis-2015
*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 2015
*Phase II. A Randomized, XXX-Blinded, XXX-Controlled, Phase II, Multicenter Study to Assess the Efficacy and Safety of XXX Different XXX Regimens in Patients with Multiple Basal Cell Carcinomas, 2014
*Phase III. An XXX-Label Study of XXX in Patients with Atopic Dermatitis Who Participated in Previous XXX Clinical Trials, 2014
*Phase 3. A Multicenter, Randomized, Double-Blind, Vehicle-Controlled Study of the Safety and Efficacy of AN2728 Topical Ointment, 2% in Children, Adolescents, and Adults (Ages 2 Years and Older) With Atopic Dermatitis 2014
*Phase 3. A Multicenter, Open-Label Study of the Long-Term Safety of AN2728 Topical Ointment, 2% in the Treatment of Children, Adolescents, and Adults (Ages 2 Years and Older) With Atopic Dermatitis 2014
*A Phase 2, Randomized, xxx-Blind, xxx-Controlled Study to Evaluate Safety and Efficacy of xxx in Subjects with Moderate to Severe Psoriasis 2014
*Phase 1 and 2, Two parts of Safety and Efficacy of Escalating doses of xxx Applied Daily for Two Consecutive Days on Approximately xxx on Trunk and Extremities in Subjects with Actinic Keratosis 2014
*Phase I. Pharmacokinetic Evaluations of XXX following Subcutaneous Administration Using Prefilled Syringe or Auto-Injector in Patients with Moderate-to-Severe Plaque Psoriasis, 2013
*Phase III. A XXX-Week, Phase 3, Randomized, XXX-Blind, XXX-Controlled, Parallel Design Study to Evaluate the Efficacy and Safety/Tolerability of Subcutaneous XXX, Followed by an Optional Long-Term Safety Extension Study, in Subjects with Moderate-to-Severe Chronic Plaque Psoriasis, 2013
*Phase IV. Pharmacokinetics and Pharmacodynamics of XXX mcg/g ointment applied XXX daily for XXX days Under Conditions of Maximal use in Pediatric Subjects (2 to 12 years of age) with Plaque Psoriasis, 2013

*Phase II. A Randomized, XXX-blind, XXX-Controlled, Multicenter Study to Assess the Safety, Tolerability and Preliminary Efficacy of XXX in Subjects with Moderate to Severe Hidradenitis Suppurativa, 2013
Phase I. Safety and Efficacy of Escalating Doses of XXX Formulations Applied XXX Daily for XXX Consecutive Days on Full Face or Approximately XXX on the Chest in Subjects with Actinic Keratosis, 2013
*Phase II-III. A Multicenter, Double-blind, Placebo-controlled, Adaptive Phase 3 Trial of XXX Melanoma Vaccine in Post-resection Melanoma Patients with a High Risk of Recurrence, 2012
*Phase II. XXX Compared with Calcipotriol plus Betamethasone Dipropionate Ointment, XXX Vehicle and Ointment Vehicle in Subjects with Psoriasis Vulgaris 2012.
*Phase III. A Phase 3 randomized, double-blind, 12-week vehicle-controlled, parallel-group study assessing the efficacy and safety of XXX cream versus vehicle cream in subjects with papulopustular rosacea, followed by a 40-week investigator-blinded extension comparing the long-term safety of XXX cream versus azelaic acid 15 % gel. 2012.
*Phase I. A Phase 1, xxx-Label, Multicenter Study to Evaluate the Safety, Tolerability, Pharmacodynamics, and Pharmacokinetics of XXX Foam, xxx, Applied Under Maximal-Use Conditions in Adolescent Subjects (Ages 12 to 16 Years) with Plaque Psoriasis. 2012.
*Phase III. A randomized, double-blind, vehicle-controlled, multicenter, parallel-group clinical trial to assess the safety and efficacy of XXX Foam, 15% topically applied twice daily for 12 weeks in patients with papulopustular rosacea, 2012.
*Phase III. A Multicenter Study with a Randomized, Double-Blind, Placebo-Controlled Induction Dosing Period Followed by a Randomized Maintenance Dosing Period and a Long-Term Extension Period to Evaluate the Efficacy and Safety of XXX in Patients with Moderate-to-Severe Plaque Psoriasis 2012.
*Phase III. A 12-Week Multicenter, Randomized, XXX-Blind, XXX-Controlled Study Comparing the Efficacy and Safety of XXX to XXX and XXX in Patients with Moderate to Severe Plaque Psoriasis with a Long-Term Extension Period, 2012
*Phase III. A Randomized, XXX-Blind, Parallel-Group, XXX-Controlled, Multicenter Study Comparing XXX to XXX, XXX in the Treatment of Actinic Keratosis of the Face or Balding Scalp, 2012
Phase IV. A Prospective Observational Study Of XXX Patterns and Effectiveness and Safety Outcomes in Advanced Basal Cell Carcinoma and Basal Cell Carcinoma Nevus Syndrome Patients, 2012
*Phase III – A Randomized, Double-Blind, Parallel-Group, Multicenter, Placebo-Controlled Study of the Safety and Efficacy of XXX in the Treatment of Mild to Moderate Distal Subungual Onychomycosis of the Toenail Adults 2011
*Phase II - A Multicenter, Randomized, Double-blind, Parallel-group Study to Evaluate the Safety and Efficacy of XXX Compared with Vehicle in Women with Female Pattern Hair Loss with an Additional Open-label Active Comparator (xxx 2%) Group, 2011
*Phase II - A Multicenter, Randomized, Double-blind, Parallel-group Study to Evaluate the Safety and Efficacy of XXX Compared with Vehicle in Men with Androgenic Alopecia with an Additional Open-label Active Comparator (xxx x%) Group, - 2011
Phase II – Topical treatment of adolescent (aged 12 – 17 years) subjects with scalp psoriasis, 2010
Phase III – Topical treatment of adolescent patients (aged 12 – 17 years) with psoriasis vulgaris, 2010
Phase III- Topical treatment of adult psoriasis vulgaris on the non-scalp regions of the body, 2010
Phase II –Oral treatment, Expanded Access Study of XXX in patients with locally advanced or metastatic basal cell carcinoma, 2010
Phase II – Dose-ranging oral treatment of severe acne vulgaris with nodules, 2010
Phase II – Topical treatment for moderate to severe facial acne vulgaris, 2010
Phase III – Parallel-group treatment/withdrawal and re-treatment of 2 oral doses of XXX of moderate to severe chronic plaque psoriasis, 2010
Phase III – Open label treatment of 2 oral doses of XXX of moderate to severe chronic plaque psoriasis

Phase III – Topical treatment of psoriasis vulgaris on the non-scalp regions of the body (trunk and/or limbs), 2010
Outpatient Registry – 2010 Multicenter, open registry of patients with psoriasis who are candidates for systemic therapy including biologics
Phase III – Subcutaneouse Injectable Treatment of Type II Diabetes, 2009
Phase III - Subcutaneouse Injectable Treatment of Type II Diabetes, 2009
Phase III - Subcutaneouse Injectable Treatment of Type II Diabetes, 2009
Phase III - Subcutaneouse Injectable Treatment of Type II Diabetes, 2009
Phase III - Subcutaneouse Injectable Treatment of Type II Diabetes, 2009
Phase II – Topical Treatment of Atopic Dermatitis,2009
Phase II – Topical Treatment of Atopic Dermatitis, 2009
Phase II – Topical Treatment of Atopic Dermatitiis 2009
Phase II - Topical Treatment of Atopic Dermatitis 2009
Phase II – Topical Treatment of Psoriasis of Intertriginous Areas 2009
Phase III- Topical treatment of pediculosis capitis, 2009
Phase III – Oral Treatment of severe recalcitrant nodular acne, 2009
Phase III - Topical Treatment of Mild to Moderate Onychomycosis, 2009
Phase III – Oral Treatment of Severe Chronic Hand Eczema, 2009
Phase III – Topical Treatment for Mild to Moderate Acne Vulgaris, 2008
Phase III – Topical Treatment for Inflammatory Rosacea 2008
Phase III- Topical Treatment for Atopic Dermatitis 2008
Phase III- Topical Treatment of Mild to Moderate Psorasis 2008
Phase II – Topical Treatment of Moderate to Severe Psoriasis 2008
Phase II- Oral Treatment for Atopic Dermatitis 2008
Phase III- Topical Treatment of Actinic Keratosis,2008
Phase II- Topical Treatment of Actinic Keratosis 2008
Phase II – Oral Treatment for Advanced Basal Cell Carcinoma, 2008
Phase III – Topical Treatment for Actinic Keratosis, 2007
Phase III – Topical Treatment for Psoriasis, 2007
Phase III – Topical Treatment for Actinic Keratosis, 2007
Phase II – Oral Treatment for Cutaneous T-Cell Lymphoma, 2007
Phase III – Topical Treatment for Moderate to Severe Acne Vulgaris, 2007
Phase III – Topical Treatment forModerate to Severe Acne Vulgaris, 2007
Phase II -Validation Study for Female Alopecia, 2007
Phase II – Topical Treatment for Moderate to Severe Psoriasis, 2007
Phase III – Topical Treatment for Moderate to Severe Psoriasis, 2007
Phase II – Oral Treatment for Herpes Zooster, 2007
Phase III – Topical Treatment for Pediculosis Capitis, 2007
Phase III _- Topical Tretment for Actinic Keratosis, 2007
Phase III- Topical Treatment for inflammatory Rosacea 2007
Phase III – Oral medication for treatment of Osteoarthritis 2007
Phase II – Topical Treatment for Actinic Keratosis, 2006
Phase III – Topical Treatment for Actinic Keratosis, 2006
Phase III – Oral Treatment for Onychomycosis, 2006
Phase III – Topical Treatment for Acne Vulgaris, 2006
Phase III – Topical Treatment for Acne Vulgaris, 2006 / 2007
Phase III – Topical Treatmnt for Impetigo, 2006/ 2007
Phase III – Topical Treatment for Tinea Pedis, 2006 / 2007

Phase III – Topical Treatment for Actinic Keratosis, 2005 / 2006
Subjects with Psoriasis for Inclusion in a repository and use in Genomic (from DNA and RNA), Serologic and Metabolic (from Serum) and Proteomic (from Protein) Research Studies, 2005 / 2006
Healthy Subjects for inclusion in a repository and use in Genomic (from DNA and RNA), Serologic and Metabolic (from Serum) and Proteomic (from Protein) Research Studies, 2005
Phase III - Topical Treatment for Actinic Keratosis, 2005
Phase II – Topical Treatment for Acne Vulgaris, 2005
Phase IV - Topical Gel for Acne Vulgaris, 2000
Phase IV - Topical gel for Acne Vulgaris, 1999
Medical Device Research - Assay Process for Determining Mitogen or Antigenic Response, 1983
Phase III- Retrospective analysis to determine common allergens and interrupted metabolic pathways in 'treatment resistant' dermatologic patients with atopic dermatitis, psoriasis, alopecia areata, chronic vaginitis, and urticaria 1982-1999
Phase III - Immunostimulation by low dose Cytoxan, 1977
Phase III - Cancer immunotherapy using DNCB, 1976-1983
Phase III - Cancer immunotherapy using Nitrogen Mustard, 1976-1983
Phase III - Cancer immunotherapy using purified protein derivative, 1976-1983
Phase III - Viral augmented melanoma antigen, RPMI, 1976-1979
Pre-Clinical: Animal toxicity studies, viral augmented melanoma antigen, Roswell Park Memorial Institute, (RPMI), 1976-1977
Phase I-II - Vesicular stomatitis virus augmented melanoma antigen, RPMI, 1976-1977
Phase III - BCG as immunoaugmentation agent in conjunction with topical immunotherapy 1976

The Clinical Trails Listed Above Were Conducted By The Following Companies:

Abbvie, Allergan, Altana, Anacor, Apotex, Asana, AstraZeneca, Asubio, Barrier, Basilea, Bayer, Boehringer Ingelheim, Biocryst, Braintree, Catawba, Centocor, Celtic, Chilter, Cipher, Clynsis, Concentrics, Covance, CuTech, Dermira, Dow, Eli Lilly, Encorium, Epithany, Galderma, Genentech, Genomics, GlaxoSmithKline, Glenmark, Health Decisions, HedgePath Pharmaceutical, Hill, ICON, Incyte, Inventiv, Kendle International, Leo Pharmaceuticals, Manhattan, Maruho, MAVIS, Merck, Novartis, Noven, Novum, Omnicare, ParaPro Inc, Parexel, PellePharm, Peplin, Pfizer, PharmaNet, Polynoma, PPD Development, PRA, Premier, Pro Trials, Quintiles, Regeneron, Research Sample Bank, Rho, Roche, Sanofi-Aventis, SciQuus, Serentis, SGS, Steifel,, Sterling Bio, Sun Pharmaceutical, Symbio, Taisho, Taro, Teva, Theraputics, TKL research, Tolmar, Topaz, Vanda, Worldwide Clinical Trials

Indications:

Alopecia Areata, Pattern Alopecia, Actinic Keratosis, Advanced Basal Cell Carcinoma, Basal Cell Carcinoma Nevus Syndrome, Squamous Cell Carcinoma, Melanoma, Lymphangiosarcoma, Cutaneous T-Cell Lymphoma, Cosmetic Dermatology, Acne Scarring, Acne Vulgaris, Rosacea, Atopic Dermatitis,, Chronic Hand Eczema, Hidradenitis Supprativa, Psoriasis, Pediculosis Capitis, Tinea Pedis, Onychomycosis, Impetigo, Herpes Zoster, Type II Diabetes, Osteoarthritis, Healthy Volunteer

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