

DERMATOLOGY RESEARCH PORTFOLIO

SHAPING THE FUTURE OF SKIN HEALTH THROUGH DERMATOLOGY RESEARCH

MESSAGE FROM THE CEO

MAZHAR JAFFRY

MBBS, MBA, MHA | 32+ YEARS OF EXPERIENCE IN HEALTHCARE LEADERSHIP



As CEO of Revival's Site Network, I'm proud to introduce our growing Dermatology division, committed to advancing skin health through innovative clinical research.



At Revival, we understand that skin conditions go beyond the surface, they can impact self-esteem, daily life, and overall well-being. That's why dermatology is at the heart of our mission. From psoriasis and eczema to vitiligo, acne, and hidradenitis suppurativa, we're driving forward research that brings real solutions to real patients.

With a dedicated network of dermatology sites across the U.S., we're not just conducting trials, we're creating access to advanced treatments for diverse communities and pushing the boundaries of what's possible in dermatologic care.

Looking ahead, Revival remains steadfast in our commitment to leading the future of dermatology research, responding to unmet needs, and transforming patient care, one clinical trial at a time.

Thank you for being part of our mission to bring skin health to the forefront.

Sincerely,

MAZHAR JAFFRY

Chief Executive Officer

COMPANY OVERVIEW

Revival's Site Network is an integrated network of clinical research sites, committed to providing patient-centric clinical trials in dermatology and other therapeutic areas, and delivering high-quality care across a diverse patient population. We specialize in a wide range of dermatologic conditions, including psoriasis, eczema, acne, vitiligo, and hidradenitis suppurativa, with a mission to improve skin health through clinical innovation.

Our focus lies in optimizing and streamlining dermatology research operations through the use of advanced technologies, aiming to minimize risk and accelerate timelines for our sponsors. We customize each approach to meet the unique needs of every dermatology trial, ensuring protocol adherence, timely regulatory submissions, and the highest standards of patient safety.



BUILDING STRONG PARTNERSHIPS FOR BETTER CLINICAL TRIAL OUTCOMES

We emphasize fostering strong partnerships with healthcare providers and research institutions, allowing us to expand our reach and access diverse patient populations. This collaboration drives better trial outcomes and contributes to advancing medical knowledge.

DRIVING EFFICIENCY & INNOVATION IN PATIENT-CENTRIC CLINICAL RESEARCH

Our reliable and predictable processes enable us to meet critical milestones faster than the industry average. We leverage real-world insights and research expertise to develop patient-centric solutions in clinical research that can be effectively implemented on a larger scale.

WHAT WE DO: ADVANCING RESEARCH, ENHANCING CARE

At Revival's Site Network, we provide comprehensive clinical trial services across multiple therapeutic areas, designed to accelerate drug and device development while keeping patient care at the heart of every study.



OUR SERVICES INCLUDE



CLINICAL TRIAL MANAGEMENT

Full-service coordination from site selection to close-out, ensuring compliance, quality, and efficiency.



THERAPEUTIC EXPERTISE

Revival Research Institute conducts dermatology trials for conditions like psoriasis, eczema, vitiligo, alopecia areata, and more, with expert research and patient-focused care.



OUTPATIENT-BASED STUDIES

Offering convenience and accessibility for participants, while maintaining high standards of data integrity and patient safety.



PATIENT RECRUITMENT & RETENTION

Proven strategies to engage diverse populations and ensure continuity throughout the study lifecycle.



REGULATORY & COMPLIANCE SUPPORT

Navigating complex regulatory environments with precision and proactive oversight.



DATA COLLECTION & MANAGEMENT

Using advanced technologies for accurate, real-time data capture and reporting.

Through a patient-centric model and a results-driven mindset, we bridge the gap between innovation and impact bringing tomorrow's treatments closer to today's patients.

OUR SITE NETWORK SETUP IN THE UNITED STATES

Revival's Site Network is actively working towards expanding and increasing access to clinical research in various locations across the United States. Our primary objective is to make clinical trials readily available to the local population of diverse communities. To achieve this, we have formed strong collaborations with experienced and diverse healthcare professionals, as well as organizations including Accountable Care Organizations (ACOs) and Healthcare Networks that provide us with top-notch Principal Investigators and patient populations in need of clinical trials.

Through these partnerships, we strive to ensure that dermatology clinical trials are conducted with the utmost care and precision, delivering valuable insights and potential therapies for the benefit of the communities we serve.



Our PIs work as independent contractors



30+ Principal Investigators & 27 Sub-Investigators throughout the United States



Over 25+ sites interspersed throughout the United States; Michigan, Texas, Nebraska, Georgia, Illinois, North Carolina



OUR PILLARS



FAST STUDY START-UP

As a clinical research site network, our study start-up process is designed for speed and efficiency, allowing us to activate sites quickly and begin enrollment without delays. With our streamlined workflows, we minimize start-up timelines by accelerating contract negotiations, ethics approvals, and site initiations. Our experienced team ensures rapid document collection and staff readiness. This enables sponsors to launch trials quickly and meet tight deadlines bringing therapies to patients sooner.

HIGHEST PATIENT RECRUITMENT

Our approach to patient recruitment is strategic and effective, ensuring high enrollment rates and diverse participant pools. By employing targeted recruitment strategies and engaging patients through multiple channels, we consistently meet our enrollment targets on time. Our efforts also focus on retention, ensuring participants remain engaged throughout the study duration.





QUERY-FREE DATA

At Revival's Site Network, we pride ourselves on providing query-free data, ensuring accuracy, precision, and compliance with regulatory standards. Our data undergoes meticulous review and verification processes, guaranteeing that it meets the highest quality standards. This commitment to excellence results in fewer queries, reduced delays, and a smoother path to approval for our clients.

THERAPEUTIC AREAS



DERMATOLOGY

- Atopic Dermatitis (Adolescents & Adults)
- Pediatric Hidradenitis Suppurativa
- Alopecia Areata
- Androgenetic Alopecia
- Hidradenitis Suppurativa
- **Atopic Dermatitis**
- Chronic Spontaneous Urticaria
- **Cutaneous Lupus**
- **Prurigo Nodularis**
- **Psoriasis**
- Vitiligo
- Rosacea



CARDIOLOGY

- Heart Failure with Kidney Failure
- Chronic Kidney & Cardiovascular Disease
- Pulmonary Hypertension & Heart **Failure**
- **Atrial Fibrillation**
- Hypertension
- **Heart Failure**
- Atherosclerotic Cardiovascular Disease



NEPHROLOGY

- Proteinuria Kidney Disease
- IgA Nephropathy

INTERNAL MEDICINE

- **Growth Hormone Deficiency**
- Type 1 Diabetes & Type 2 **Diabetes**
- Diabetic Peripheral Neuropathy (DPN)
- Alzheimer's Disease

THERAPEUTIC AREAS



PSYCHIATRY

- Autism Spectrum Disorder
- Pediatric Bipolar Disorder
- Bipolar Disorder
- Major Depressive Disorder (MDD)
- Obsessive-Compulsive Disorder (OCD)
- Attention Deficit Hyperactivity Disorder (ADHD)



ONCOLOGY

- Advanced Pancreatic Cancer
- Diffuse Large B-Cell Lymphoma
- Indolent Non-Hodgkin's Lymphoma (iNHL)
- Malignant Myeloid Hematologic Neoplasms (Myelofibrosis)
- Metastatic Breast Cancer
- Mutant Solid Tumors
- Relapsed/Refractory Follicular Lymphoma
- Relapsed/Refractory Follicular Non-Hodgkin's Lymphoma
- · Uterine Serous Carcinoma



WOMEN'S HEALTH

- Vaginal Yeast Infection
- · Endometrial Hyperplasia
- Oral Contraceptive
- Nocturnal Enuresis

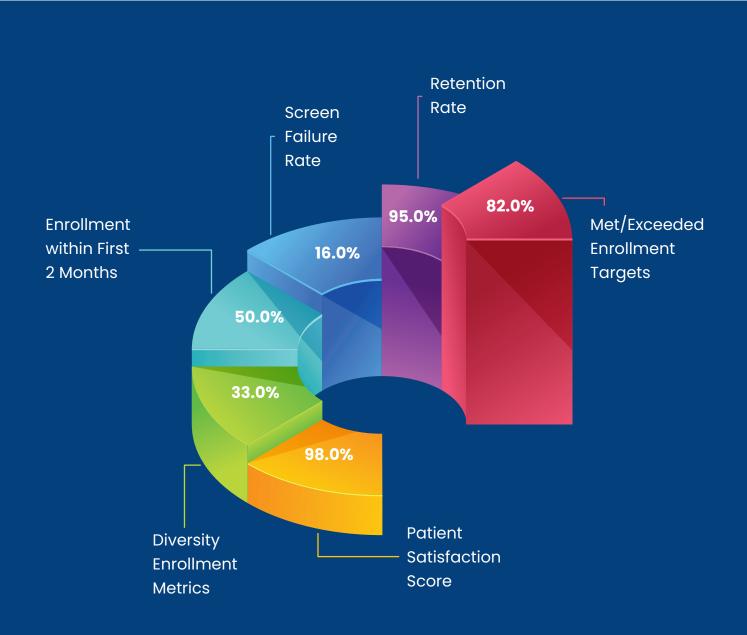


PULMONOLOGY

- Asthma
- COPD
- · Shift work disorder
- Obstructive sleep apnea

ROBUST PATIENT ENROLLMENT METRICS

Over the years, we've consistently demonstrated strong and efficient patient enrollment strategies for our clinical trials. Our historic enrollment metrics underscore our commitment to expediency without compromising quality. Detailed percentages further highlight our proficiency in this domain.



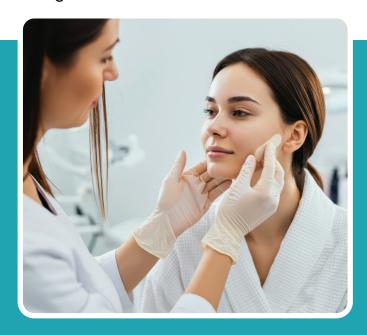


DERMATOLOGY RESEARCH

At Revival's Site Network, we've consistently demonstrated our strength in conducting high-performance dermatology trials. With a focus on improving skin health and maintaining operational excellence, our site network has played a key role in the successful execution of numerous dermatology studies. From enrolling hundreds of participants to surpassing sponsor expectations, our work reflects a strong commitment to advancing dermatologic care and innovation through clinical research.

KEY HIGHLIGHTS

- Successfully conducted studies for HS, PN, Vitiligo, Psoriasis & AD.
- Screened over 10,000 patients across various dermatology studies.
- Our dermatology site network has randomized over 6,000 patients to date, demonstrating our strong capabillities in clinical trial execution.



 We have recently expanded our footprint by adding two new sites in Georgia and North Carolina-reflecting the continued growth and success of our network.

SKILLED IN CONDUCTING TIME-SENSITIVE CLINICAL STUDIES ACROSS DIVERSE POPULATIONS

At our Dermatology site, we specialize in delivering high-quality results under tight timelines. Our team is uniquely equipped to navigate the demands of accelerated clinical trials without compromising the scientific integrity or regulatory compliance required for successful study outcomes.



ENROLLMENT GOAL METRICS

Exceeded enrollment goal for the site by up to 57% and randomized patients within a compressed timeline to meet a critical FDA submission deadline.

HOW WE DELIVERED

Driven by a "can-do" attitude and a hands-on approach, our team worked tirelessly to ensure the study's success, balancing speed with precision every step of the way.

WHAT WE DID

- Rapid Response: Stepped in as the new study site after a last-minute change and fully onboarded within days.
- Expert Training: Trained staff on the proper use of the imaging device to capture specific dermatological views relevant to pruirgo nodularis evaluation.
- Collaborative Approach: Maintained clear, consistent communication with Incyte throughout the process.

I have worked closely with both Manali and Navya on the INCYTE 206 & 319 studies, and man-o-man has it been a real pleasure to work with both of them! They have both been a Monitor's dream/ideal Study Coordinators to work with. Manali and Navya are extremely knowledgeable about the protocols, study procedures, and how to use the study platforms/devices. If there is ever an issue or error, they are proactive in letting me know, and finding solutions/corrective actions immediately, rather than me having to discover the errors when I come on-site. They are extremely receptive to any feedback/ suggestions I may have and make any necessary corrections on time.

I can't speak for the 206 study since I wasn't with your site at the start of it, but for the 319 study, Manali and Navya have helped tremendously in working out the study kinks within the IRT, eDiaries, and EDC. I know you all blazed the trail in enrollment for the 319 study, so you were the first to test-run and find out when systems weren't working like they were supposed to. There were definitely hiccups with randomizing participants in IRT, eDiaries syncing/populating, and excessive query issuing in EDC. You all had the expertise and knowledge to know when things weren't working right, and always took the right actions by contacting me or the study team to inform us of the issues and find out the next best steps forward. I want to thank you all so much because I know it's never fun being the first to blaze the way, but it's because of your hard work through these starting hiccups that the study is now running smoother and more efficiently! You all were always so helpful and patient and never complained when working through those early study days.

Any time I email or call Manali or Navya, I always get a response within 24 - 48hours. No part of my email goes unanswered. I truly appreciate their attention to detail and diligent follow-up with patients and my own ad-hoc requests.

You all have also SMASHED it out of the park with recruitment! Your site is one of the highest enrollers for both the 206 and 319 studies, and you still manage to have quality data and timely EDC entry. I do not know how you all do it, but it is truly top-tier work! Your source is very detailed and captures all necessary procedures. If I put a comment in RealTime, it is always answered/resolved by the next time I come on-site. I never have to pester Manali or Navya about outstanding open queries, they always stay on top of them, and resolve them within the study agreed timelines.

Manali & Navya are both incredibly hard workers, but they are also just great people and a joy to be around and interact with! They always greet me with a smile and have a positive attitude.

I truly love coming to and working with your site! I hope to have many more studies with you all in the continued future.

Thanks!



ATOPIC DERMATITIS CLINICAL TRIAL - PFIZER

For this particular Pfizer study, our team successfully enrolled over 35 patients within a challenging 5-month enrollment window. Despite the extensive regulatory documentation and startup processes, we exceeded the Sponsor's enrollment target—demonstrating our ability to deliver rapid, high-quality recruitment under pressure.

Enrollment Goal Metrics

- Successfully randomized over 35 patients within the 5-month target window
 - 2 Exceeded the Sponsor's enrollment target

HOW DID WE DO IT?

- Targeted Community Outreach: Collaborating with dermatology clinics and local organizations to engage and identify individuals living with atopic dermatitis.
- Rigorous Pre-Screening: Utilizing strict and compliant screening protocols to ensure that only qualified and suitable participants are enrolled.
- Patient-First Strategy: Provided personalized support, flexible scheduling, and clear education to build trust and enhance participant understanding of the study's goals.

This successful execution highlights our team's agility, commitment to patient engagement, and ability to meet ambitious enrollment goals under tight timelines—reinforcing our strengths in dermatology research.

HIDRADENITIS SUPPURATIVA CLINICAL TRIAL -PFIZER

This HS clinical trial presented significant challenges due to the protocol's strict eligibility criteria, making patient recruitment particularly difficult. Despite these hurdles, our team enrolled over 53 patients, demonstrating exceptional performance in a highly selective therapeutic area.

ENROLLMENT & RANDOMIZATION HIGHLIGHTS

- 53+ patients enrolled, surpassing expectations for a complex indication
- 27 patients successfully randomized, achieving 90% of the Sponsor's target of 30
- Maintained rapid response times with regulatory and feasibility submissions completed within 48–72 hours



OPERATIONAL EXCELLENCE ACROSS TEAMS

Regulatory & Business Development

Worked diligently to ensure all IRB documents, contracts, and feasibility forms were submitted accurately and on time—often within just two to three days—allowing for swift study activation.



Clinical Operations

Maintained end-to-end control over trial execution:

- Screened and identified eligible patients under stringent criteria
- Managed timely receipt of investigational products (IPs) and lab kits
- Ensured clean, source-verified data entry into the EDC within 24 hours
- Coordinated effectively with CRAs to address data queries and compliance reviews
- Closely tracked enrollment targets to ensure Sponsor expectations were met

This case study highlights our ability to execute complex trials with strict criteria, uphold regulatory efficiency, and maintain data integrity and operational speed—even in high-pressure environments. Our success in this study reaffirms our capability to deliver results for sponsors in dermatology and other high-challenge therapeutic areas.

STRATEGIC SITE CAPABILITIES FOR DERMATOLOGY STUDIES

In addition to our strong enrollment metrics, our sites are well-equipped to support large-scale dermatology clinical trials. These high-capacity locations enable us to reach a wide and diverse patient population efficiently.

DERMATOLOGY CLINICAL SITES LED BY EXPERIENCED INVESTIGATORS



ALIMOIN, MD

DERMATOLOGY | YEARS OF EXPERIENCE: 30+YEARS

TROY, MICHIGAN

BACKGROUND AND EXPERTISE

Dr. Ali Moiin, M.D., is a board-certified dermatologist serving the Metro Detroit area. With over twenty years of experience, Dr. Moiin holds medical licenses in both Michigan and California. He completed his undergraduate studies at the University of California, Santa Barbara, where he also earned a Master of Arts in Biology. In 1987, Dr. Moiin obtained his Doctor of Pharmacy degree from the University of California, San Francisco. He continued his education at the University of California, Davis, where he earned his medical degree.

PROFESSIONAL TRAINING & EXPERIENCE

Dr. Moiin completed his internship and residency at the Wayne State University School of Medicine, Department of Dermatology, in Detroit, Michigan. He furthered his expertise with a rotation in cosmetic surgery at Tulane University Medical Center. Since 1995, Dr. Moiin has held multiple faculty appointments at Wayne State University School of Medicine and continues to serve as a faculty member. While his extensive clinical experience speaks volumes, it is his dedication to the community that truly distinguishes him. Dr. Moiin is actively involved with the Grosse Pointe Park Foundation and the Wayne State University School of Medicine's LGBT People in Medicine organization, earning the 2016 WSU Service Award for his contributions.

CONTRIBUTIONS TO REVIVAL RESEARCH INSTITUTE, LLC

Since joining Revival Research Institute, LLC, in 2019, Dr. Moiin has participated in more than 50 dermatology clinical trials, covering a range of indications such as atopic dermatitis, nail psoriasis, and prurigo nodularis, among others. His research site is the top recruiter within the Revival Site Network and has been recognized as one of the highest-enrolling sites in the United States. The site is supported by a team of highly experienced and motivated clinical coordinators and laboratory technicians.

COMMITMENT TO QUALITY AND PATIENT CARE

Our site is fully equipped to support any clinical trial, with a comprehensive suite of essential equipment and a team of highly qualified staff. Together, we are committed to achieving Revival's goal of providing sponsors with the highest quality, query-free data while delivering exceptional patient care to all research participants.

EDUCATION

- Detroit Medical Center/Wayne State University Residency, Dermatology, 1995
- University of California San Francisco Medical Centers Internship, Internal Medicine, 1992
- University of California, Davis, School of Medicine Medical School, 1991

BOARD CERTIFICATIONS

- American Board of Dermatology (Recertified)
- Michigan Board of Medicine (Recertified)

FRANCES FLORENTINO, MD

DERMATOLOGY | YEARS OF EXPERIENCE: 31+ YEARS | EVANS. GEORGIA

BACKGROUND AND EXPERTISE

Dr. Frances Florentino is a board-certified dermatologist currently practicing in Evans, Georgia. Her dermatology training was completed at Henry Ford Health System in Detroit, Michigan.



Dr. Florentino treats both pediatric and adult patients for various skin, nail, and hair disorders. With over 31 years of experience in the medical field, she is actively involved in conducting clinical trials for multiple skin conditions.

PROFESSIONAL MEMBERSHIPS AND DERMATOLOGY EXPERTISE

Dr. Florentino is a member of prestigious organizations such as the American Board of Dermatology, the American Academy of Dermatology, and the Georgia Society of Dermatologists. Her extensive experience allows her to treat both pediatric and adult patients with a wide range of skin, nail, and hair conditions. She is proficient in skin cancer surgeries and the removal of various types of skin lesions. Additionally, Dr. Florentino is skilled in cosmetic dermatology, offering treatments such as Botox® Cosmetic, Dysport®, Restylane®, and various laser therapies.



MULTILINGUAL CAPABILITIES & COMPREHENSIVE PATIENT CARE

Dr. Florentino's ability to speak both English and Spanish enhances her communication with a diverse patient population. She is dedicated to providing comprehensive care addressing both medical and cosmetic dermatological needs. Her combination of clinical expertise, academic experience, and compassionate care ensures that patients receive top-quality treatment for their dermatological concerns.

EDUCATION & EXPERIENCE

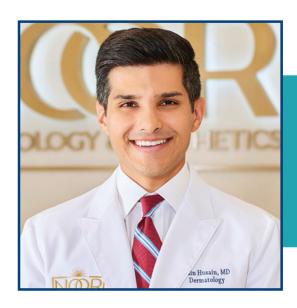
With over 31 years of experience in the medical field and significant contributions to research, Dr. Florentino is a vital part of Revival Research's network of investigators. She is affiliated with Doctors Hospital, Piedmont Augusta, and Piedmont McDuffie.

EDUCATION

- Henry Ford Health/Henry Ford Hospital Residency, Dermatology, 1993-1996
- Henry Ford Health/Henry Ford Hospital Internship, Transitional Year 1992–1993
- University Of Michigan Medical School Medical School, 1992

BOARD CERTIFICATIONS

- American Board of Dermatology
- MI State Medical License (Active through 2000)
- GA State Medical License (Active through 2005)
- OH State Medical License (Active through 2007)



ZAIN HUSSAIN, MD

DERMATOLOGY | YEARS OF EXPERIENCE: 10+ YEARS | CARY, NORTH CAROLINA

BACKGROUND & EXPERIENCE

Dr. Zain Husain is a dual board-certified dermatologist and Mohs micrographic surgeon with more than a decade of clinical expertise. He earned his medical degree from Rutgers New Jersey Medical School, completed his dermatology residency at Georgetown University Hospital, and pursued advanced fellowship training in dermatologic surgery and cosmetic dermatology at Montefiore Medical Center. Throughout his career, Dr. Husain has held leadership positions in prominent dermatology practices and served as a Clinical Instructor at George Washington University, where he contributed to the training of future dermatologists.

SPECIALIZED CLINICAL FOCUS

Dr. Husain specializes in the diagnosis and treatment of complex skin cancers, with advanced expertise in Mohs micrographic surgery and reconstructive techniques. He also has a strong background in cosmetic dermatology, offering a wide range of laser treatments and aesthetic procedures. This unique combination allows him to deliver truly comprehensive, patient-centered care - providing both life-saving treatments and confidence-restoring cosmetic solutions.

ACADEMIC CONTRIBUTIONS & PUBLICATIONS

Dr. Husain has authored more than 20 peer-reviewed publications covering dermatologic surgery, melanoma, laser therapy, and complex reconstructions, with his research featured in leading journals such as Dermatologic Surgery and JAAD. He has presented his work at national conferences and has contributed to medical education as a Clinical Instructor. With his strong academic background, surgical expertise, and dedication to advancing patient care, Dr. Husain brings exceptional value to Revival's Investigators Network as a Principal Investigator.

EDUCATION & EXPERIENCE

Dr. Husain received his medical degree from Rutgers New Jersey Medical School, followed by dermatology residency training at Georgetown University and advanced fellowship at Montefiore Medical Center. As a board-certified Mohs surgeon, he brings extensive expertise in both dermatologic oncology and cosmetic dermatology.

EDUCATION

Fellowship - Micrographic Surgery and Dermatologic Oncology/Cosmetic Dermatology

Montefiore Medical Center - Albert Einstein College of Medicine

Residency - Dermatology 2015-2016

Georgetown University Hospital

Internship - Transitional Year 2012-2015

Albert Einstein Medical Center

• Medical Degree 2011–2012

Rutgers New Jersey Medical School

BOARD CERTIFICATIONS

Mohs Micrographic Surgery
 2007-2011

Micrographic Dermatologic Surgery

DERMATOLOGY

· American Board of Dermatology





THE WAY FORWARD

Revival's Site Network is committed to advancing clinical research through innovation, collaboration, and community engagement. Our focus is on optimizing trial procedures, integrating cutting-edge technologies, and enhancing efficiency. We aim to increase diversity in clinical trials by 80% within the next year, expanding access to underrepresented communities while upholding the highest ethical standards.

COMMITTED TO REDEFINING PATIENT CARE, **ONE CLINICAL** TRIAL AT A TIME.

FOR MORE INFORMATION

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SCAN HERE