Scott T. Guenthner M.D. The Indiana Clinical Trials Center, PC 824 Edwards Drive, Suite 172 Plainfield, Indiana 46168

Phone: (317) 837-6082

PROFESSIONAL EXPERIENCE:

03/2006 – Present President/CEO/Principle Investigator

The Indiana Clinical Trials Center, PC

Plainfield, Indiana

7/2002 – Present Physician/President

Dermatologist

The Dermatology Center of Indiana, PC

Plainfield, Indiana

6/1995 – 8/1995 Research Assistant

College of Medicine Student Research Fellowship

University of Iowa Iowa City, Iowa

1/1992 – 8/1994 Research Assistant

Complement Laboratory University of Iowa Iowa City, Iowa

CERTIFICATION AND LICENSURE:

7/2002 Indiana Medical License #01051021 11/2002 – Present American Board of Dermatology 03/2009- Present Good Clinical Practice Certification

HOSPITAL AFFILIATIONS:

7/2002 – Present Hendricks Regional Health

Danville, Indiana

7/2002 – Present St. Francis Hospitals and Health Services

Indianapolis, Indiana

7/2003 – Present Clarian West Medical Center

Avon, Indiana

7/2004 – Present St. Vincent Hospitals and Health Services

Indianapolis, Indiana

10/2005 - Present Community Hospitals

Indianapolis, Indiana

EDUCATION:

Graduated 5/1998 Doctor of Medicine

University of Iowa Iowa City, Iowa

Graduated 5/1994 Bachelor of Science with Highest Distinction

Psychology

University of Iowa Iowa City, Iowa

Honors: Medical: President, Alpha Omega Alpha, Iowa Chapter, 1997

The Hancher-Finkbine Medallion, 1998

Undergraduate: Phi Beta Kappa, Junior Inductee, 1993

Cumulative GPA 4.0

University of Iowa Collegiate Scholar, 1994

Omicron Delta Kappa Province XI Leader of the Year, 1993-94

Activities: Medical: President, Medical Student Ambassador Program, 1996

Medical Student Council, 1995-1998

Student Preceptor, M1 Physical Examination Skills, 1997-1998

Undergraduate: Co-President, Phi Beta Kappa

General Council, Omicron Delta Kappa

Volunteer, Johnson County Big Brothers/Big Sisters Program

POST GRADUATE TRAINING:

7/1999 - 6/2002 Residency

Dermatology Indiana University Indianapolis, Indiana

FELLOWSHIPS:

7/1998 – 6/1999 Transitional Internship

St. Vincent Hospital and Health Services

Indianapolis, Indiana

MEMBERSHIPS/COMMITTEES:

7/2002 – Present University of Iowa Alumni Association Board of Directors

PROJECTS:

5/1999 Indiana University Department of Dermatology

Skin Cancer Education Project

Coordinated a skin cancer education project in which faculty and residents in the Department of Dermatology educated individuals of the community about melanoma and non-melanoma skin cancers at a local

minor league baseball game.

PUBLICATIONS, ABSTRACTS, AND PRESENTATIONS:

- Maves, KK, Guenthner, ST, et.al.. Cloning and characterization of the cDNA encoding guinea pig properdin: a comparison of properdin from three species. Immunology, 86 (3), 475-79, Nov., 1995
- Guenthner, ST, Hurwitz, RM, Buckel, LJ, and Gray, HR. *Cutaneous Squamous Cell Carcinomas Consistently Show Histologic Evidence of In-Situ Changes: A Clinical-Pathological Correlation*. Journal of the American Academy of Dermatology, 41(3), 443-48, Sept., 1999.
- Guenthner, ST, Hurwitz, RM, Buckel, LJ, and Gray, HR. *Cutaneous Squamous Cell Carcinomas Consistently Show Histologic Evidence of In-Situ Changes: A Clinical-Pathological Correlation*. Abstract, Journal of Cutaneous Pathology, 26 (9), 465, Oct., 1999.
- Guenthner, ST, Hurwitz, RM, Buckel, LJ, and Gray, HR. *Cutaneous Squamous Cell Carcinomas Consistently Show Histologic Evidence of In-Situ Changes: A Clinical-Pathological Correlation*. Poster Presentation. The American Society of Dermatopathology Annual Meeting, San Diego, CA, Nov 4-7, 1999.
- Guenthner, ST, Draft, KS, and Hood, AF. Virtual Dermatology Case, Indiana University Department of Dermatology, May, 2000.
- Piliang MP. Guenthner ST. Treadwell PA. *Abnormal hair growth in a child with atopy*. Archives of Dermatology. 137(11), 1521-6, Nov, 2001.
- Jungers EA, Guenthner ST, Farmer ER, Perkins SM. A skin cancer education initiative at a professional baseball game and results of a skin cancer survey. Int J Dermatol. 2003 Jul; 42(7): 524-9.
- Schaefer C, Cappelleri JC, Cole J, Guenthner S, Fowler J, Johnson S, Mamolo C. Burden-of-Illness in Moderate-to-Severe Psoriasis Subjects Seeking Treatment: An Overall Descriptive Assessment (2013).
- Pariser, D., McConnehey, D., Matheson, R., Kempers, S., Guenthner, S., Bukhalo, M. A Phase 2 Study of Photodynamic Therapy (PDT) with Aminolevulinic Acid HCl (ALA) 20% Topical Solution + Blue Light vs ALA Topical Solution Vehicle + Blue Light Using Spot and Broad Area Application and Incubation Times of 1, 2 & 3 Hours for the Treatment of Multiple Actinic Keratoses (AK) on the Face or Scalp. Presented at the 71st Annual American Academy of Dermatology Meeting March 2013.

CLINICAL TRIAL EXPERIENCE:

3/2006 – 08/2006	COBRA Trial, Phase IV Trial of Clobex Spray for the Treatment of Plaque Psoriasis
6/2006-12/2009	Phase IV Study of Raptiva Safety (RESPONSE)
8/2007-3/2008	Phase III Study of Raptiva in TNF- Alpha Biologic Failures

10/2007-12/2007	Phase III Study of Brand vs Generic Benzaclin Gel Efficacy and Safety
1/2008-10/2008	Phase III Study of Brand vs. Generic Duac Gel Efficacy and Safety
3/2008-2/2010	Phase III Study of Raptiva Effectiveness for Scalp Psoriasis
9/2008-1/2009	Phase III Study of Brand vs.Generic Aldara Cream for the Treatment of Actinic Keratosis Efficacy and Safety
10/2008-4/2009	Phase III Study of Ingenol Mebutate Gel vs Placebo for the Treatment of Actinic Keratosis Efficacy and Safety
12/2008- 6/2009	Phase III Study of ABT-874 vs Etanercept vs Placebo for the Treatment of Moderate to Severe Plaque Psoriasis Efficacy and Safety
12/2008-1/2009	Phase III Study of Brand vs Generic Protopic 0.1% for the Treatment of Atopic Dermatitis
2/2009- 9/2009	Phase III Study of Brand vs. Generic Aldara Cream for the Treatment of Actinic Keratosis Efficacy and Safety
5/2009-11/2009	Phase IV Study, Open-label, 12-week Trial Assessment of Effectiveness, Safety, and Subject Satisfaction with Oracea (doxycycline, USP) Capsules 40 mg (30mg immediate release and 10 mg delayed release beads) as Monotherapy or as Add-on Therapy to Existing Topical Regimens for the Treatment of Rosacea
6/2009-1/2010	A Phase III, Multi-Center, Randomized, Parallel Group, Double-Blind, Vehicle Controlled Study to evaluate the Efficacy and Safety of PEP005 Gel, 0.015% in Patients with Actinic Keratosis ON the Head (Face or Scalp)
3/2006 04/2013	Phase IV Five year Study of Enbrel Safety (OBSERVE)
1/2009-06/2012	Phase III Study of Alitretinoin vs Placebo for the Treatment of Chronic Hand Eczema Efficacy and Safety
5/2009-02/2012	A Phase III, Multi-Center, Open-Label Continuation Study in Moderate to Severe Chronic Plaque Psoriasis Subjects who Completed a Preceding Psoriasis Study with ABT-874
8/2009-12/2010	A 12 Month, Long Term Follow-up Study of Patients with Actinic Keratosis on the Head (face or Scalp) who have Completed the Day 57 in Studies PEP005-016 or PEP005-025
9/2009- 07/2011	Double-Blind, Randomized, Phase III, Parallel Group Study Evaluating the Efficacy and Safety of CIP-Isotretinoin in patient with severe recalcitrant nodular acne
08/2010-12/2011	Phase III, A Double-Blind, Vehicle Controlled, Randomized, Parallel Design, Multiple-Site Clinical Study to Evaluate The Efficacy and Safety of Desoximetasone 0.25% Topical Spray in Patients with Moderate to Severe Plaque Psoriasis

09/2010-01/2012	Phase II, Multicenter, Randomized, Double-Blind, Dose-Ranging Study to Evaluate IDP-107 Versus Placebo in the Treatment of Severe Acne Vulgaris with Nodules
10-2010-07/2011	Phase III, A National, Multi-center, Prosepctive, Randomized, Double-Blind, 4-arm, Parallel Group, 8-week Study in Subjects with Psoriasis Vulgaris on the Non-Scalp Regions of the Body (trunk or limbs)
10-2010- 07/2011	Phase III, A Randomized, Controlled Evaluation of the Safety and Efficacy of a Topical Treatment for Moderate- Severe Facial Acne Vulgaris
03/2011-01/2012	A Multicenter, Randomized, Double-Blind, Vehicle Controlled, Parallel Group Comparison Study to Determine the Therapeutic Equivalence of Generic Imiquimod Cream 3.75% And Zyclara (Imiquimod) Cream3.75% In Subjects with Actinic Keratoses
06/2011-02/2012	A Multicenter, Double-Blind, Randomized, Vehicle Controlled, Parallel-Group Study Comparing Adapalene and Benzoyl Peroxide Topical Gel 0.1%/2.5% to Epiduo and Both Active Treatments to Placebo in the Treatment of Mild to severe Acne Vulgaris
09/2011-04/2012	A Randomized, Double-Blind, Placebo-Controlled, Multiple-Site, Study Comparing Metronidazole Topical Gel 1% to Metrogel 1% in the Treatment of Moderate to Severe Rosacea
12/2011- 07/2012	A Multicenter, Randomized, Double-Blind, Placebo-Controlled Evaluation of Rosacea-Specific Inflammatory Biochemical markers in the Skin of Adults with Papulopustular Rosacea Treated with Daily Doxycycline 40 mg (30 mg Immediate Release/10mg Delayed Release) capsules
12/2011-12/2012	A Phase II Study of Photodynamic Therapy with Levulun Topical Solution+ BLU light vs. Levulun Topical Solution Vehicle + BLU Light Using Spot and Broad Area Application and Incubation Times of 1,2, and 3 Hours For the Treatment of Multiple Actinic Keratoses on the Face and Scalp
12/2011-08/2012	An Evaluation of the Burden of Illness Among Adults in the United States with Moderate to Severe Plaque Psoriasis
03/2012- 09/2013	Phase 3 Randomized, Double-Blind, 12- week, Vehicle controlled, parallel-group study assessing the efficacy and safety of CD5024 1% cream versus vehicle cream in subjects with papulopustular rosacea, followed by a 40- week investigator blinded extension comparing the long- term safety of CD5024 1% cream versus azelaic acid 15% gel
03/2012- 11/2013	A Sequential Treatment Regimen of Cryotherapy and Picato (Ingenol Mebutate) gel, 0.015% Field Therapy Compared to cryotherapy alone for the treatment of Actinic Keratoses on the Face and Scalp
04/2012- 12/2012	A Phase II Study Comparing Treatment with LEO 90100 with Betamethasone Dipropionate in LEO 90100 Vehicle and Calcipotriol in LEO 90100 Vehicle in Subjects with Psoriasis Vulgaris

04/2012- 12/2012	A Phase II Study Comparing Treatment with LEO 90100 with Calcipotriol Plus Betamethasone Ointment, LEO 90100 Vehicle and Ointment Vehicle in Subjects with Psoriasis Vulgaris
04/2012- 12/2012	A Randomized, Double-Blind Placebo Controlled, 4-Week trial of IMO-3100 in Patients with Moderate to Severe Plaque Psoriasis
04/2012-05/2014	Randomized, Double-Blind, Vehicle-Controlled, Multicenter, parallel-Group Clinical Trials to assess the Safety and Efficacy of Azelaic Acid Foam, 15% Topically Applied Twice Daily for 12 weeks in Subjects with papulopustular Rosacea
05/2012-01/2013	A randomized, Double-Blind, Placebo-Controlled, Parallel Design, Multiple Site, Clinical Study Comparing Naftifine HCL Cream 1% to Naftin Cream 1% in the Treatment of Tinea Pedis
08/2012- 07/2013	A randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Vehicle Controlled, Multicenter Study Comparing Imiquimod 3.75% to Zyclara
10/2012- 05/2013	A randomized, Double-Blind, Placebo-Controlled, Parallel Design, Multiple Site, Clinical Study Comparing Clindamycin 1%/Benzoyl Peroxide 5% Topical Gel to Duac
10/2012- 07/2013	A randomized, Double-Blind, Placebo-Controlled, Parallel Design, Multiple Site, Clinical Study Comparing Diclofenac Sodium Gel 3% to Solaraze
08/2012-06/2014	A Phase 2, Multicenter, Randomized, Double-Blind, Vehicle Controlled, Dose Escalating Study to Evaluate Coretexalone 17a- Propionate Once Daily or Twice Daily in Subjects with Facial Acne Vulgaris.
04/2013- 12/2013	A randomized, Double-Blind, Placebo-Controlled, Parallel Design, Multiple Site, Clinical Study Comparing Diclofenac Sodium Gel 3% to Solaraze
02/2013-02/2013	A Randomized, Controlled Evaluation of the Safety and Efficacy of Topical Treatments for Moderate-Severe Facial Acne Vulgaris-Braintree
03/2013-09/2013	A Multicenter, Randomized, Double-Blind, Phase 3 Study of the Safety, Efficacy, Systemic Exposure, and Pharmacodynamics of Calcipotriene Foam, 0.005% Versus Vehicle Foam in Pediatric Subjects (Ages 2-11) with Plaque Psoriasis
04/2013-04/2014	A Double-Blind, Randomized, Parallel-Group, Vehicle Controlled, Multicenter Study Comparing TOLMAR Azelaic Acid Gel, 15% to Reference Listed Drug in the Treatment of Rosacea
04/2013-06/2014	A Double-Blind, Randomized, Parallel Group, Vehicle-Controlled, Multicenter Study Comparing TOLMAR Naftifine HCl cream, 2% to Reference listed Drug in the treatment of Tinea Pedis

05/2013-09/2014	A randomized, Vehicle-Controlled, Double-blind, Parallel Group, Multi-center Phase III Study to Evaluate the Safety and Efficacy of M518101 in Subjects with Plaque Psoriasis
09/2013-02/2015	A Multicenter, Double-blind, Randomized, Parallel-group, Vehicle-controlled Study to Evaluate the Safety and Efficacy and Clinical Equivalence of a Generic Azelaic Acid Gel, 15% and the Reference listed Finacea Gel 15% in patients with Moderate Facial Rosacea
09/2013-02/2015	A Phase II, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled, Dose Finding, and Efficacy Study of VPD-737 in the Treatment of Subjects with Chronic Pruritis
11/2013-02/2015	Safety and Efficacy of Escalating Doses of Ingenol Mebutate Once Daily for Two or Three Consecutive Days When Used on Full Face, Full Balding Scalp or Approximately 250 cm2 on The Chest in Subjects with Actinic Keratosis
11/2013-08/2014	A Multicenter, Randomized, Double-Blind, Vehicle Controlled, Parallel Group Comparison Study to Determine the Therapeutic Equivalence of Generic Imiquimod Cream, 2.5% and Zyclara cream 2.5% in Subjects with Actinic Keratoses
11/2013-06/2015	A Randomized, Double-Blind, Multicentric, Parallel-group, Active and Placebo Controlled, Three Arm Clinical Study to Compare the Efficacy and Safety of Clindamycin Phosphate 1.2%/Benzoyl Peroxide 5% gel versus DUAC Gel versus Placebo in the ratio of 2:2:1 respectively in Patients with Acne Vulgaris
12/2013-12/2014	A Randomized, Double-Blind, Vehicle-Controlled, Multicenter, Parallel Group Study of the Safety of Betamethasone Dipropionate Spray 0.05% versus Diprolene Lotion 0.05% and the Efficacy of Bethamethasone Dipropionate Spray 0.05% versus Vehicle Spray in the Treatment of Moderate Plaque Psoriasis
02/2014-10/2014	A randomized, Placebo-Controlled, Parallel Group Study to Evaluate the Efficacy and Safety of An Anticholinergic Agent Agent for the Treatment of Primary Axillary Hyperhydrosis
02/2014-01/2015	A Safety and Efficacy Study to Compare Dapsone Dermal Gel with Vehicle Control in Patients with Acne Vulgaris
02/2014-09/2014	A Randomized, Double-Blind, Multiple Site, Placebo-Controlled, Parallel Design Study Comparing Adapalene and Benzoyl Peroxide Gel 0.1%/2.5% to Epiduo Topical Gel in the Treatment of Acne Vulgaris
05/2014-07/2015	A Phase 3 study of photodynamic therapy with Levulan Kerastick Topical solution + Blue light versus topical solution vehicle+ blue light for the treatments of actinic keratoses on the upper extremities
06/2014-02/2015	Efficacy and Safety of Oxymetazoline HCl Cream 1.0% for the Treatment of Persistent Erythema Associated with Rosacea
11/2014-05/2015	Safety and efficacy of escalating doses of LEO 43204 applied once daily for two consecutive days on approximately 250 cm2 on trunk and extremities in subjects with actinic keratosis

09/2014-09/2015	A Randomized, Double-Blind, Placebo-Controlled, Study Investigating Vaccine Responses in Adults with Moderate to Severe Atopic Dermatitis Treated with Dupilumab
03/2012-12/2015	A Phase 3B, Randomized, Double-Blind, Active-Controlled, Multicenter Study to Evaluate a "Subject Tailored" Maintenance Dosing Approach in Subjects with Moderate-to-Severe Plaque Psoriasis-PSTELLAR
05/2015-02/2016	A Multicenter, Randomized, Double Blind, Vehicle-Controlled Study to Evaluate the Safety and the Effect on Sweat Production of Three Concentrations of Topically Applied BBI-4000 gel in Subjects with Axillary Hyperhidrosis
05/2015-12/2015	A Randomized, Double-Blind, Parallel-Design, Multiple-Site Study to Evaluate the Therapeutic Equivalence of Diclofenac Sodium Gel 3% Compared to Solaraze 3%, gel in the Treatment of Actinic Keratosis
05/2015-12/2015	A Phase 2, Multicenter, Randomized, Double-Blind, Vehicle-Controlled Study of the Safety, Tolerability, and Efficacy of 0.15% and 0.25% Concentrations of Topical SM04554 Solution in Male Subjects with Androgenetic Alopecia (AGA)
Current Trials	
3/2006 – Present	APPLES Trial, Phase IV Study of Protopic Ointment Safety and Outcomes in the Treatment of Atopic Dermatitis
9/2008- Present	Phase IV Study of Remicade Safety (PSOLAR)
12/2008-Present	Phase IV Study of Humira Safety for the Treatment of Plaque Psoriasis (ESPRIT Registry)
10/2012-Present	A Phase 3 Study to Evaluate the Efficacy and Safety of Induction and Maintenance Regimens of Brodalumab Compared with Placebo and Ustekinumab in Subejcts with Moderate to Severe Plaque Psoriasis
03/2013-Present	A 64-Week, Phase 3, Randomized, Double-Blind, Placebo Controlled, Parallel Design Study to Evaluate the Efficacy and Safety/Tolerability of Subcutaneous SCH900222/MK-3222, Followed by an Optional Long-Term Safety Extension Study, in Subjects with moderate-to-Severe Chronic Plaque Psoriasis
12/2013-Present	I1F-MC-RHBA (b): A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study Comparing the Efficacy and Safety of LY2439821 to Etanercept and Placebo in Patients with Moderate-to-Severe Plaque Psoriasis
03/2014-Present	RegiSONIC: A Prospective Observational Study of Treatment Patterns and Effectiveness and Safety Outcomes in Advanced Basal Cell Carcinoma and Basal Cell Carcinoma Nevus Syndrome Patients
04/2014-Present	A Double-Blind, Randomized, Parallel-Group, Active-Control Study to Compare the Efficacy and Safety of CHS-0214 Versus Enbrel® in Subjects With Chronic Plaque Psoriasis (CHS-0214-04) (RaPsOdy)

08/2014-Present	Twelve Month Follow-up Evaluation of Subjects Participating in Dusa-CP0108 (A Phase 3 Study of Photodynamic Therapy with Levulan® Kerastick® Topical Solution + Blue Light Versus Topical Solution Vehicle + Blue Light for the Treatment of Actinic Keratoses on the Upper Extremities)
02/2015-Present	A Phase 2 Study Comparing The Occurrence Of Actinic Keratoses On The Face In High-Risk Individuals After Cryotherapy + Photodynamic Therapy With Levulan® Topical Solution + Blue Light Versus Cryotherapy + Vehicle Topical Solution + Blue Light
11/2014-Present	A Phase 3, Multicenter, Randomized, Double-blind Study to Evaluate the Efficacy and Safety of Guselkumab for the Treatment of Subjects With Moderate to Severe Plaque-type Psoriasis and an Inadequate Response to Ustekinumab
11/2014-Present	A Phase 3, Multicenter, Randomized, Double-blind, Placebo and Active Comparator-controlled Study Evaluating the Efficacy and Safety of Guselkumab in the Treatment of Subjects with Moderate to Severe Plaque-type Psoriasis Incorporating Randomized Withdrawal and Retreatment
11/2014-Present	A Phase 3, Multicenter, Randomized, Double-blind, Placebo and Active Comparator-controlled Study Evaluating the Efficacy and Safety of Guselkumab in the Treatment of Subjects with Moderate to Severe Plaque-type Psoriasis
11/2014-Present	A Randomized, Multicenter, Double-blind, Placebo-controlled Study to Evaluate the Efficacy and Safety of 1.5 mg/kg per Day of Sarecycline Compared to Placebo in the Treatment of Acne Vulgaris
11/2014-Present	A Multi-Center Open-Label Evaluation of the Safety of Sarecycline Tablets in the Treatment of Acne Vulgaris
02/2015-Present	A Multicenter, Randomized, Double-Blind, Parallel Group Comparison of Halobetasol Propionate Foam 0.05% versus Vehicle Foam in Subjects with Plaque Psoriasis
02/2015-Present	A Phase 3 Confirmatory Study Investigating the Efficacy and Safety of Dupilumab Monotherapy Administered to Adult Patients with Moderate-to-Severe Atopic Dermatitis
02/2015-Present	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
02/2015-Present	An Open-Label study of Dupilumab in Patients with Atopic Dermatitis Who Participated in Previous Dupilumab Clinical Trials
02/2015-Present	A Multicenter, Randomized, Double-Blind, Vehicle-Controlled, Parallel Group Comparison Study To Determine The Therapeutic Equivalence Of Generic Fluorouracil Cream, 0.5% And Carac® (Fluorouracil) Cream, 0.5% In Subjects With Actinic Keratoses

04/2015-Present	A Multicenter, Randomized, Double-Blind, Vehicle controlled, Parallel Group Comparison Study To Determine The Therapeutic Equivalence Of A Generic Ingenol Mebutate Gel, 0.05% And Picato® Gel, 0.05% In Subjects With Actinic Keratosis On The Trunk Or Extremities
03/2015-Present	An Open-label Study Evaluating the Long-term Efficacy, Quality of Life, and Safety of ABSORICA® (isotretinoin) Capsules Administered Without Food in Subjects With Severe Recalcitrant Nodular Acne
07/2015-Present	A Phase 3, Randomized, Double-Blind, Vehicle-Controlled Efficacy and Safety Study of DRM04 in Subjects with Axillary Hyperhydrosis
07/2015-Present	An Open-Label Study Assessing Long-Term Safety Of Drm04 In Subjects With Primary Axillary Hyperhidrosis
08/2015-Present	Efficacy and Safety of Ingenol Mebutate Gel in Field Treatment of Actinic Keratosis on Full Face, Balding Scalp or Approximately 250 cm ² on the Chest
08/2015-Present	A Double-Blind, Randomized, Parallel-Group, Active Control Study to Compare the Efficacy and Safety of CHS-1420 Drug Product Versus Humira® in Subjects With Chronic Plaque Psoriasis
08/2015-Present	A Double-blind, randomized, placebo-controlled study, Evaluating the efficacy and safety of Once Weekly High Dose Oral Finasteride in the Treatment of Severe Nodulocystic Acne
08/2015-Present	A Randomized, Double-Blind, Placebo-Controlled Ascending Multiple Dose Study to Assess the Safety, Tolerability, Pharmacokinetics, and Pharmacodynamics of VTP-43742 in Healthy Volunteers and Psoriatic Patients and Clinical Proof-of-Concept in Psoriatic Patients
12/2015-Present	A Phase 3, Multi-Center, Randomized, Double-Blind, Vehicle-Controlled, 2-Arm, Parallel Group Comparison Study Comparing the Efficacy and Safety of IDP-121 and IDP-121 Vehicle Lotion in the Treatment of Acne Vulgaris
12/2015- Present	A Phase 3, Multicenter, Double-Blind, Randomized, Vehicle Controlled Clinical Study to Access the Safety and Efficacy of IDP-122 in the Treatment of Plaque Psoriasis
12/2015-Present	A Phase 3, Randomized, Vehicle-Controlled, Double-Blind, Multicenter Study to Evaluate the Safety and Efficacy of a Once-Daily CLS001 Topical Gel Versus Vehicle Administered for 12 Weeks to Subjects with Papulopustular Rosacea with a 4 Week Follow-up Period
02/2016-Present	A Phase 1b Open-Label, Randomized Study Evaluating the Absorption and Systemic Pharmacokinetics and HPA Axis Suppression Potential of Topically Applied IDP-118 Lotion and HP Monad Lotion in Subjects with Moderate to Severe Plaque Psoriasis
02/2016-Present	A Phase 2, Multicenter, Evaluator-Blinded Randomized, Vehicle-Controlled Study to Compare the Safety and Efficacy of IPD-118 Lotion with Tazorac ® (tazarotene) Cream, 0.05% in the Treatment of Plaque Psoriasis

02/2016-Present BI 655066 versus Ustekinumab and placebo comparators in a

randomized double blind trIal for Maintenance use in Moderate to

severe plaque type psoriasis

02/2016-Present BI 655066 versus placebo In a Multicenter randomized double-blind

study in patients with Moderate to severe chronic plaque psoriasis evaluating the efficacy and safety with randomized withdrawal and re-

treatment