<u>CURRICULUM VITAE</u> JOSEPH TAUBER, M.D.

Tauber Eye Center 4400 Broadway, Suite 202 Kansas City, MO 64111 (816) 531-9100 (816) 531-9105 FAX

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$H_{C}G$	ucation	

1978 Bachelor of Sciences in Biology and Chemistry

Union College Schenectady, N.Y.

1982 Doctor of Medicine

Harvard Medical School Boston, MA

Postdoctoral Training

7/82-6/83 Internship in Internal Medicine

Northwestern Medical Center Chicago, IL

7/83-6/84 Residency in Internal Medicine

Beth Israel Hospital Boston, MA

7/84-6/87 Residency in Ophthalmology

Tufts-New England Medical Center Boston, MA

7/87-6/88 Fellowship in Ocular Immunology

Massachusetts Eye & Ear Infirmary Boston, MA

7/88-6/89 Fellowship in Corneal and External Diseases

Massachusetts Eye & Ear Infirmary Boston, MA

Licensure and Certification

1983 Diplomate National Board of Medical Examiners

1984 Massachusetts State Licensure1989 Missouri State Licensure

1989 Diplomate of the American Board of Ophthalmology

1997 Kansas State Licensure

Academic Appointments

1999-2011 Clinical Professor of Ophthalmology

Kansas University School of Medicine

1999-Present Medical Director

Saving Sight Eye Bank

1998-1999 Clinical Professor of Ophthalmology

University of Missouri-Kansas City School of Medicine

1994-1997 Clinical Associate Professor of Ophthalmology

University of Missouri-Kansas City School of Medicine

1989-1993 Clinical Assistant Professor of Ophthalmology

University of Missouri-Kansas City School of Medicine

1991-1999 Director of Clinical Research

University of Missouri-Kansas City School of Medicine

Hospital Staff Appointments

1989-Present St. Lukes Hospital of Kansas City-Kansas City, MO

Honors, Awards and Distinctions

Valedictorian, Summa Cum Laude **Union College**

William E. Lasnik Prize

Harvard

Medical School Founding Member Cabot Medical Society

Northwestern Co-chairman House Staff Committee **Medical Center** Northwestern Memorial Hospital

Tufts-New England

Medical Center Charles J. Preefer Resident Research Award

Massachusetts National Eve Institute Individual NRSA Award (EY06052) 1987-1989 Eve & Ear

Infirmary

Heed Ophthalmic Foundation Fellowship 1987-88 Heed Foundation/Knapp Fellowship 1988-89

Post - Training American Academy of Ophthalmology Achievement Award, 2005

American Academy of Ophthalmology Program Committee

Member, Cornea Subcommittee 2002 - 2006

Professional Affiliations

American Academy of Ophthalmology

Association for Research in Vision and Ophthalmology

Cornea Society

Metropolitan Medical Society of Greater Kansas City Kansas City Society of Ophthalmology & Otolaryngology

Midwest Corneal Society

Versatile Method for Evaluating a Cell Culture by Various Morphological Techniques.

Lahav M, Ishii Y, Craft J, Tauber J.

Stain Technology 57: 331-347, 1982

Cryotherapy for Iris Neovascularization and Neovascular glaucoma.

Lahav M, Tauber J, Haug S.

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New Clinical Classification for Iris Neovascularization.

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Simplified Diagnostic Coding Sheet for Computerized Data Storage and Analysis in Ophthalmology

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Dramatic Improvement Following Plasmapheresis in the Atopic Keratoconjunctivitis Associated with Hyper IgE Syndrome.

Aswad M, Tauber J, Baum J.

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Cataract Surgery in Ocular Cicatricial Pemphigoid.

Sainz de la Maza M, Tauber J, Foster CS.

Ophthalmology 95 (4): 481-486, 1988.

Lymphocyte Proliferation After HSV-1 Infection in Igh-1 Disparate Congenic Murine Strains.

Tauber J, Wells PA, Foster CS.

Investigative Ophthalmology Visual Science 29 (Suppl): 152, 1988.

Scleral Grafting in Necrotizing Scleritis.

Sainz de la Maza M, Tauber J, Foster CS.

Ophthalmology 95 (9) Suppl: 161, 1988.

Glaucoma in Patients with Ocular Cicatricial Pemphigoid.

Tauber J, Melamed S, Foster CS.

Ophthalmology 96 (1): 33-37, 1989

Systemic Immunologic Responses Following Intracameral Inoculation of HSV-1 in

Susceptible and Resistant Igh-1 Disparate Congenic Murine Strains.

Tauber J, Hemady R, Ihley T, Opremcak EM, Foster CS.

Investigative Ophthalmology Visual Science 30 (Suppl): 359, 1989.

Value of Conjunctival Biopsy in Peripheral Ulcerative Keratitis.

Sainz de la Maza M, Foster CS, Tauber J, Hoang-Xuan T.

Ophthalmology 96 (9) Suppl: 121, 1989.

An Analysis of Therapeutic Decision Making in Peripheral Ulcerative Keratitis.

Tauber J, Hoang-Xuan T, Sainz de la Maza M, Foster CS.

Cornea 9 (1): 66-73, 1990.

Viral Isolation and Systemic Immune Responses After Intracameral Inoculation of Herpes

Simplex Virus type 1 in Igh-1 Disparate Congenic Murine Strains.

Hemady R, Tauber J, Ihley TM, Opremcak EM, Foster CS.

Investigative Ophthalmology Visual Science 31 (11): 2335-2341, 1990.

Ocular Cicatricial Pemphigoid (OCP) – The Potential for Cure.

Neumann R, Tauber J, Foster CS.

Ophthalmology 97 (9) Suppl: 132, 1990.

Systemic Chemotherapy for Ocular Cicatricial Pemphigoid.

Tauber J, Sainz de la Maza M, Foster CS.

Cornea 10 (3): 185-195, 1991.

Romberg's Progressive Hemifacial Atrophy: An Association with Spontaneous Scleral Melting.

Hoang-Xuan T, Foster CS, Jakobiec FA, Tauber J, Sainz de la Maza M, Krebs W.

Cornea 10 (4): 361-366, 1991.

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Hemady R, Tauber J, Foster CS.

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Investigation and Treatment of Peripheral Corneal Ulcers.

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Remission and Recurrence after Withdrawal of Therapy for Ocular Cicatricial Pemphigoid.

Neumann R, Tauber J, Foster CS.

Ophthalmology 98: 858-862, 1991.

Lipemia Retinalis

Martinez KR, Cibis GW, Tauber J.

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Improved Detection of Disease Progression in Ocular Cicatricial Pemphigoid.

Tauber J, Jabbur N, Foster CS.

Cornea 11 (5): 446-451, 1992.

The Effectiveness of Excimer Photorefractive Keratectomy for Myopia.

Piebenga LW, Deitz MR, Irvine JW, Matta CS, Tauber J, Sabates FN.

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Long Term Results of Systemic Chemotherapy for Ocular Cicatricial Pemphigoid.

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Dry Eve Syndrome and in Normal Volunteers.

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Corneal Topographic Abnormalities Following the Use of Artificial Lubricants.

Mardelli PM, Tauber J, Piebenga LW.

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Photoablation Without Nitrogen Gas Flow.

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Improvement in Squamous Metaplasia and Tear Osmolarity Following AquasiteTM Treatment.

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Matta CS, Piebenga LW, Deitz MR, Tauber J.

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Tauber J, Bealer L, Cavanaugh T, Durrie D.

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Excimer Photorefractive Keratectomy for Myopia.

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Two Zone PRK for Moderate Myopia.

Piebenga LW, Matta CS, Deitz MR, Tauber J.

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A Multicenter Comparison of Diclofenac Sodium 0.1% (DS) to Ketorolac Tromethamine 0.5% (KT)

in Patients with Acute Seasonal Allergic Conjunctivitis (SAC).

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Investigative Ophthalmology Visual Science 35 (4): 1291, 1994.

Changes in Corneal Topography After Instillation of Artificial Lubricants.

Yu JS, Wu M, Crosser V, Tauber J.

Investigative Ophthalmology Visual Science 35 (4): 1291, 1994.

Contact Lens and Spectacle Use Before and After PRK.

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Cicatricial Pemphigoid.

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With Keratoconjunctivitis Sicca.

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The Effects of Instillation of Topical NSAIDs on Corneal Sensation.

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Temporary Intracanalicular Collagen Implants In Normal Volunteers.

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Epidemiological Study of Punctal Size in Normal Volunteers.

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Five and Three Year Follow-up of Photorefractive Keratectomy for Myopia of -1 to -6 Diopters.

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Excimer Retreatment for Myopic Photorefractive Keratectomy Failures. Six to 18 Month Follow-up.

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Ophthalmology 103: 444-451, 1996

Ablation Zone Centration after Photorefractiuve Keratectomy and Its Effect on Visual Outcome.

Deitz MR, Piebenga LW, Matta CS, Tauber J, Anello RD, DeLuca M

Journal of Cataract and Refractive Surgery 22(6): 696-701, 1996

The Effects of Instillation of Diclofenac sodium 0.1% on Photophobia following Pupillary Dilation.

A Pilot Study in Normal Volunteers.

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A Multicenter Comparison of the Ocular Efficacy and Safety of Diclofenac 0.1% Solution with that

Of Ketoralac Tromethamine 0.5% Solution in Patients with Acute Seasonal Allergic Conjunctivitis.

Tauber J, Raizman MB, Ostrov CS, Laibovitz RA, Abelson MB, Betts JG, Koester JM,

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Evaluation of Two Different Methods of Epithelial Removal Prior to Excimer Laser

Photorefractive Keratectomy for Myopia.

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Investigative Ophthalmology Visual Science 40 (4): S107, 1999

Refractive Outcomes Following Excimer Laser Photorefractive Keratectomy for Myopia Using

Two Different VISX Excimer Lasers.

Cable M, Matta C, Piebenga L, Polepalle S, Steward C, Tauber J.

Investigative Ophthalmology Visual Science 40 (4): S106, 1999

Safety and Efficacy of Nedocromil Sodium Ophthalmic Solution 25 BID in Patients with Ocular Itch Associated With Allergic Conjunctivitis.

Tauber J, Alocril® Community Ocular Allergy Trial Study Group

Investigative Ophthalmology Visual Science 42(4) S910, March 2001

Efficacy and Safety of Combination Therapy with Brimonidine 0.2% and Latanoprost 0.005% Versus Fixed Combination Timolol 0.5%/Dorzolamide 2%

Zabriskie N A, Ahmed I K, Cantor L B, Kent A B, Mundorf T, Tauber J, Rubin J M, Hoop J

Investigative Ophthalmology Visual Science 42 (4) S833, March 2001

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Nedocromil Sodium Ophthalmic Solution 2% Twice Daily in Patients with Allergic Conjunctivitis.

Tauber J, and the Alocril Community Allergy Trial Study Group

Advances in Therapy 19 (2) 73-84, March / April 2002

Double-Masked, Placebo-Controlled Safety and Efficacy Trial of Diquafasol Tetrasodium (INS365) Ophthalmic Solution for the Treatment of Dry Eye

Tauber J, Kellerman D

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Parasitic Keratitis and Conjunctivitis.

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Juvenile Rheumatoid Arthritis.

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Phase III Safety Evaluation of Cyclosporine 0.1% Ophthalmic Emulsion Administered Twice Daily to Dry Eye Disease Patients for Up to 3 Years.

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Followup of Penetrating Keratoplasty.

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In: Price, Ed. Curbside Consultation in Cornea and External Disease. Thoroughfare NJ, Slack, 2010 Amebiasis.

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in Intraocular Inflammation, ed. Zierhut, M, Ohno S, Orefice F, Pavesio C, Rao N. (Eds.)

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in Intraocular Inflammation, ed. Zierhut, M, Ohno S, Orefice F, Pavesio C, Rao N. (Eds.)

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Juvenile Idiopathic Arthritis.

Tauber J

In: Krachmer JH, Mannis MJ, Holland EJ, Eds. Cornea, 3rd ed., St. Louis, CV Mosby, 2011.

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Sainz de la Maza M, Molina N, Gonzalez-Gonzalez LA, Doctor PP, Tauber J, Foster CS. *Ophthalmology*, 119: 43-50, 2012.

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Scleritis Therapy

Sainz de la Maza M, Molina N, Gonzalez-Gonzalez LA, Doctor PP, Tauber J, Foster CS. *Ophthalmology*, 119: 51-58, 2012.

Is There a Difference Between the Various Corticosteroid Preparations?

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Parasitic Keratitis and Conjunctivitis.

Tauber J, Anzaar F.

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Ed. by Sainz de la Maza M, Tauber J, Foster CS. New York, Springer-Verlag, 2012. pp. 1-307.

Scleritis Associated with Inflammatory Bowel Disease

Sainz de la Maza M, Molina N, Alonso Gonzalez-Gonzalez L, Doctor PP, Tauber J, Foster CS, *Investigative Ophthalmology Visual Science, in press.*

Clinical features and presentation of posterior scleritis: a report of 31 cases

Gonzalez-Gonzalez LA, Molina-Prat N, Priyanka Doctor P, Tauber J, Sainz de la Maza M, Foster CS, *Ocular Immunology and Inflammation 2013, in press.*

Recent Participation in Multicenter Research Studies

1990-1992	Cyclosporine-High Risk Corneal Transplantation Study
1990-1996	VISX Photorefractive Keratectomy Study
1990-1995	Fibronectin Eyedrops in the Treatment of Nonhealing Corneal Ulcers
1992-1993	Efficacy of Carbomer Gel Versus Hypotears in Dry Eye Patients
1992-1994	Efficacy and Safety of Rimexolone Ophthalmic Suspension
1,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Versus Pred Forte 1% for the Treatment of Uveitis
1992-1993	Efficacy and Safety Evaluation of Lodoxamide 0.1% Ophthalmic
1,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Solution Versus Placebo in Vernal Keratoconjunctivitis
1992-1994	Adjunctive Therapy with Galardin TM Matrix Metalloproteinase
1,,, = 1,,, :	Inhibitor in Patients with Corneal Ulcers Caused by Infection
1992-1994	Adjunctive Therapy with Galardin TM Matrix Metalloproteinase
	Inhibitor in Patients with Corneal Ulcers Caused by Rheumatoid Arthritis
1992-1993	A Clinical Study Evaluating the Ocular Safety, Comfort and Efficacy
	Of Cellufresh® Compared with Celluvisc®, Hypotears P.F.® and
	Tears Naturale II® in Subjects with Dry Eye Syndrome
1992-1993	A Prospective, Randomized, Double-Masked, Parallel Group, Two
	Week, Multicenter Comparison Trial of the Ocular Efficacy and Safety
	Of Voltaren Ophthalmic 0.1% Solution With That Of Acular 0.5%
	Ophthalmic Solution In Patients With Acute Seasonal Allergic
	Conjunctivitis
1993-1994	Safety and Efficacy of Lotoprednol Etabonate in Acute Anterior Uveitis
1994-1995	A Clinical Study Comparing the Ocular Safety and Efficacy Of
	Acular With Livostin And Vehicle In Subjects With Seasonal Allergic
	Conjunctivitis
1994-1996	A Double-Masked, Double-Dummy, Randomized, Acyclovir
	Controlled, Parallel Group Study To Compare The Efficacy And
	Safety Of Famciclovir With Acyclovir In The Treatment Of
1007 1007	Patients With Ophthalmic Zoster
1995-1996	A Dose-Ranging Study Evaluating the Safety, Tolerability, and
	Efficacy of Cyclosporine and Vehicle Ophthalmic Emulsions in
1996-1996	The Treatment of Moderate to Severe Keratoconjunctivitis Sicca
1990-1990	Comparison of Lotprednol Etabonate in Acute Anterior Uveitis and Placebo in the Treatment of Inflammation Following Cataract
	Surgery With Intraocular Lens Implantation
1996-1997	Multicenter Comparison of the Comfort and Efficacy of Genteal
1//0-1///	Ophthalmic Lubricant Solution Versus Refresh Plus Ophthalmic
	Lubricant Solution in the Treatment of Patients With Dry Eye Syndrome
1997-1998	A randomized double-masked (investigator), active-controlled, parallel
1,,,, 1,,,0	Evaluation of the safety and efficacy of Cidofovir topical ophthalmic solution in the
	treatment of viral epidemic keratoconjunctivitis
1997-1998	A Randomized, Single-Masked (Investigator), Active-Controlled, Parallel
	Evaluation Of The Safety And Efficacy Of Cidofovir Topical Ophthalmic
	Solution In The Treatment Of Herpes Epithelial Keratitis
1997-1998	A Safety And Efficacy Assessment Of Hyaluronidase Solution Injected Into
	The Vitreous Body Of The Eye To Hasten The Clearing Of Blood From The
	Vitreous

Recent Participation in Multicenter Research Studies

1997-1998 1998-1999	A Multicenter, Double-Masked, Randomized, Vehicle-Controlled, Parallel-Group Study of the Safety and Efficacy of Cyclosporine 0.05% And 0.1% Ophthalmic Emulsions Used Twice Daily for Up To One Year In Patients With Moderate to Severe Keratoconjunctivitis Sicca A Six Month, Multicenter, Triple-masked, Placebo-Controlled Adjunctive Therapy Study of the Safety and Efficacy of AL-6221 0.0015% and AL-6221 0.004% Ophthalmic Solution in Patients With Open Angle Glaucoma or Ocular Hypertension Maintained on Timoptic 0.5%
1998-2000	A Phase III Study of MDX-RA Compared With Placebo Administered in Patients Undergoing Extracapsular Extraction with Phacoemulsification For Cataract
1998-1999	A Multicenter, Open-Label, Phase 2 Study Extension Evaluating the Safety of Cyclosporine 0.1% Ophthalmic Emulsion Used Twice Daily for Up to One Additional Year in Patients With Moderate to Severe
1998-2000	Keratoconjunctivitis Sicca A Multicenter, Open-Label, Phase 3 Study Extension Evaluating the Safety of Cyclosporine 0.1% Ophthalmic Emulsion Used Twice Daily for One Additional Year in Patients With Moderate to Severe Keratoconjunctivitis Sicca Previously Dosed for 6 to 12 Months with
1998-2001	Cyclosporine 0.05% or 0.1% Safety and Efficacy Study of Vitrase (Hyaluronidase) for Ophthalmic Intravitreal Injection for Clearance of Severe Vitreous Hemorrhage Study
1998-1999	RUF366 A Pilot Study of Zenarestat 0.3% (RUF366) For the Treatment of Corneal Epitheliopathy Associated With Diabetes Mellitus
1998-1999	Genteal-Gel Ophthalmic Lubricant in the Treatment of Patients with Moderate to Severe Dry Eye Syndrome
1999-Present	A 48 Month, Multicenter, Randomized, Double-Masked, Placebo-Controlled, Clinical Study To Evaluate The Effectiveness And Safety Of Oral Memantine In Daily Doses Of 20mg And 10mg In Patients With Chronic Open Angle Glaucoma At Risk For Glaucomatous Progression
2000-2001	Cyclosporine 0.05% Ophthalmic Emulsion vs Tears Naturale Free Used Twice Daily In Increasing Daily Lens Comfort and Wear Time Joseph Tauber, M.D., Primary Investigator
2000-2001	Memantine Oral Doses of 20mg and 10mg In Patients with Chronic Open-Angle Glaucoma At Risk for Glaucomatous Progression
2001-2002	Inspire 365 Ophthalmic Solution vs Placebo In Patients with Moderate to Severe Dry Eye Disease
2001-2002	A Multicenter Evaluation of Strategies to Optimize the Tolerability of Restasis 0.05% Ophthalmic Emulsion
2001-2004	Randomized, 5 Year Postmarketing Safety Study of Xalatan Compared to "Usual Care" in Patients with Open-Angle Glaucoma or Ocular Hypertension
2001-2004	A Three Month Clinical Study Comparing the Efficacy and Tolerability of Alphagan and Xalatan in subjects with Chronic Open-Angle Glaucoma or Chronic Ocular Hypertension Presently on Topical Beta Blockers

Recent Participation in Multicenter Research Studies

2001-2004	A Four-Month, Randomized, Double-Masked, Parallel Group, Two-Arm
2001-2004	Study Comparing the Safety and Efficacy of Alphagan™ 0.2%/Xalatan™ 0.005%
	Combination Therapy and Cosopt [™] Combination Therapy for the Reduction
	Of Intraocular Pressure in Patients with Glaucoma or Ocular Hypertension
2001-2003	A Multi-Center, Parallel, Randomized, Double-Masked, Vehicle-Controlled, Dose Ranging
2001 2002	Study to Evaluate the Safety, Tolerability and Efficacy of Testosterone 0.01%, 0.1%, and
	0.3% Ophthalmic Solutions Administered Twice Daily for 16 weeks in Patients with
	Keratoconjunctivitis Sicca
2001-2003	A Multi-Center, Double- Masked, Randomized, Placebo-Controlled, Dose-Ranging Study of
	Multiple Ocular Instillations of INS365 Ophthalmic Solution vs. Placebo in Subjects with Dry
	Eye Disease.
2001-2002	A Prospective, Randomized, Parallel-group, Multi-center, Double-masked Trial Comparing
	the Efficacy and Safety of 1.5% Levofloxacin Ophthalmic Solution with 03.% Ofloxacin
	Ophthalmic Solution for Treating Bacterial Keratitis.
2004-2005	A 21 Day, Pilot, Multicenter, Investigator-masked Randomized Parallel Comparison
	of the Safety and Efficacy of Gatifloxacin 0.3% Ophthalmic Solution Compared with
	Ciprofloxacin 0.3% Ofloxacin Ophthalmic Solution in Patients with Acute Bacterial Corneal
2004-2005	Ulcers. A Multi-Center, Parallel Group, Double- Masked, Randomized, Placebo-Controlled Study of
2004-2003	Multiple Ocular Instillations of Diquafasol Tetrasodium Ophthalmic Solution, 2% in Subjects
	with Dry Eye Disease.
2004-2005	Rebamipide Ophthalmic Suspension in the Treatment of Dry Eye: A Multi-Center, Phase 3,
	Randomized, Double-Masked, Placebo-Controlled, Parallel Group, 26 Week Study.
2005-2007	A 24 week randomized, double-blind, multicenter, parallel-group, placebo-controlled
	evaluation of the safety and efficacy of 0.3% and 1% pimecrolimus ophthalmic suspensions
	used twice daily in patients with moderate to sever keratoconjunctivitis sicca.,
2005-2006	Optimization of Clinical Testing in Patients with Dry Eye.
2005-2007	A Study to Evaluate the Clinical and Microbial Efficacy of 0.6% ISV-403 compared to
2005 2005	Vigamox [™] in the Treatment of Bacterial Conjunctivitis.
2005-2007	Randomized, Double-Masked Study of Rimexolone 0.2% and Vehicle in Treatment of Dry
2006-2009	Eye. A Double-Masked, Placebo-Controlled, Parallel Group, Multi-center, Dose-ranging Study to
2000 2007	Assess the Efficacy and Safety of LX211 as Therapy in Subjects with Active Sight
	Threatening, Non-Infectious Intermediate-, Anterior and Intermediate-, Posterior- or Pan-
	Uveitis
2006-2009	A Double-Masked, Placebo-Controlled, Parallel Group, Multi-center, Dose-ranging Study to
	Assess the Efficacy and Safety of LX211 as Therapy in Subjects with Clinically Quiescent
	Sight Threatening, Non-Infectious Intermediate-, Anterior and Intermediate-, Posterior- or
2007 2000	Pan- Uveitis
2006-2009	A Double-Masked, Placebo-Controlled, Parallel Group, Multi-center, Dose-ranging Study to Assess the Efficacy and Safety of LX211 as Therapy in Subjects with Active Sight
	Threatening, Non-Infectious Anterior Uveitis
2007-2009	A Multi-center, Placebo-Controlled, Randomized, Parallel-Group Dose-Ranging
2007-2007	Study to Assess the Efficacy and Safety of LX201 for Prevention of Corneal Allograft
	Rejection Episodes and Graft Failure Following Penetrating Keratoplasty with LX201
	Implantation In Subjects Who Are At Increased Immunological Risk
2007-2009	A Multi-center, Placebo-Controlled, Randomized, Parallel-Group Dose-Ranging
#UU -#UUJ	Study to Assess the Efficacy and Safety of LX201 Implantation for the Prevention of
	Corneal Allograft Rejection Episodes or Graft Failure in Subjects Who Have
	Experienced One or More Rejection Episodes Following Penetrating Keratoplasty
	Experienced One of More Rejection Episodes Following Fenetiating Relatioplasty

2007-2008 A Phase III Study of the Efficacy and Safety of Cyclosporine Ophthalmic Solution, 0/1% (STA-603) Compared to Vehicle in Subjects with Moderate to Severe Dry Eye Syndrome A Phase 2b Multicenter, Randomized, Double-Masked Study of the Safety and Efficacy of 2007-2008 Difluprednate 0.05% Ophthalmic Emulsion Compared to Prednisolone Acetate 1% Ophthalmic Suspension in the Treatment of Endogenous Anterior Uveitis 2007-2008 CR-1485CJ Evaluation of Multifocal Soft Contact Lens Designs A Multi-Center, Open-Label, Randomized, Pilot Study of the Safety and Efficacy of Azasite 2008-2009 Ophthalmic Solution 1% in Combination With Mechanical Therapy vs. Mechanical Therapy Alone in Subjects with Blepharitis 2008-2009 Efficacy and Safety of COL-101 for the Treatment of Blepharitis in Patients with Facial Rosacea A Phase II Randomized, Masked, Parallel-Group Study of Safety and Preliminary 2008-2009 Efficacy of the Latanoprost Punctum Plug Delivery System in Subjects with Primary Open Angle Glaucoma or Ocular Hypertension 2008-2009 An Open-Label Proof-Of-Concept Study To Assess The Effects of Two AIN457 Doses (on Day 1 and Day 22) In Patients With Noninfectious Uveitis A Multicenter, Parallel Group, Double-masked, Randomized, Placebo-controlled Study of The 2009-2009 Effects of Diquafasol tetrasodium Ophthalmic Solution, 2% In Subjects With Dry Eye Disease and a Central Corneal Staining Score of 3 (NEI Scale) An Evaluation of the Safety and Efficacy of Moxifloxacin AF Ophthalmic Solution 0.5% 2009-2009 for the Treatment of Bacterial Conjunctivitis in the USA A Device Evaluation Study to Assess the Physical and Clinical Performance of Prototype 2009-2012 Punctal Plug Design Iterations An Open-Label, Phase 2 Study of Punctal Placement of the Latanoprost Punctal Plug 2009-2009 Delivery System (L-PPDS in Subjects With Ocular Hypertension (OH) or Open-Angle Glaucoma (OAG) 2009-2009 A Phase I/II Prospective, Randomized, Double Masked, Vehicle And Comparator Controlled, Dose Ranging Study Of Cp-690,550 In Subjects With Dry Eye Disease 2009-2009 A Multi-Center, Open Label Randomized Pilot Study of the Safety and Efficacy of Azasite Ophthalmic Solution, 1% in Combination with Mechanical Therapy versus Mechanical Therapy Alone in Subjects with Blepharitis A Clinical Safety and Efficacy Evaluation of Zylet® (loteprednol etabonate 0.5% 2009-2010 and tobramycin 0.3% ophthalmic suspension) compared to Lotemax® (loteprednol etabonate ophthalmic suspension 0.5%), Tobramycin Ophthalmic Solution USP, 0.3%, and the Vehicle of Zylet® for the Treatment of Blepharoconjunctivitis in Pediatric Subjects A Multi-Center, Double Masked, Placebo-Controlled, Parallel-Group, Randomized Study 2009-2010 of the Safety and Efficacy of Azithromycin Ophthalmic Solution, 1% versus Placebo for Four Weeks in Subjects with Dry Eye Disease A Randomized, Double-Masked, Parallel-Group, Vehicle-Controlled, Multicenter, 2010-2011 Exploratory Study Assessing Safety and Efficacy of BOL-303242-X Ophthalmic Suspension in Dry Eye Syndrome A Randomized, Multi-Center, Double-Masked, Vehicle-Controlled, Parallel-Group, 2011-2011 Safety and Efficacy Study of Azithromycin Ophthalmic Solution, 1% versus Vehicle for Four Weeks in Subjects with Blepharitis A Prospective, Multi-Center, Randomized, Double-Masked, Positive-Controlled Phase 3 2011-2013 Clinical Trial Designed to Evaluate the Safety and Efficacy of Iontophoretic Dexamethasone Phosphate Ophthalmic Solution Compared to Prednisolone Acetate (1%) in Patients with Non-Infectious Anterior Segment Uveitis

2012-2013 A Randomized, Double-Masked, Parallel –Group, Comparative Study to Evaluate the Clinical Efficacy and Safety of ISV-502 (1.0% Azithromycin and 0.1% Dexamethasone) Compared to Azasite alone, Dexamethasone 0.1% alone and Vehicle in the Treatment of Patients with Non-Bacterial Blepharitis A Randomized, Double-masked, Placebo-controlled Study of the Safety and Efficacy of 2012-2013 2% Rabamipide versus placebo in the treatment of Dry Eyes. A Randomized, Double-masked, Placebo-controlled, Dose-Ranging Study of EBI-005 2012-2013 Ophthalmic Solution versus Placebo in subjects with Dry Eye Syndrome. 2013-2013 A Phase 3, Multicenter, Randomized, Double Masked and Placebo Controlled Study Evaluating the Efficacy of a 5.0% Concentration of Lifitegrast Ophthalmic Solution Compared to Placebo in Subjects with Dry Eye Currently Using Artificial Tears (OPUS-2) 2013-2013 A Randomized, Double-masked, Placebo-controlled Study of the Safety and Efficacy of GevokizumAb in the Treatment of Active Non-infectious Intermediate, Posterior, or Pan-Uveitis 2013-2013 A Randomized, Double-masked, Placebo-controlled Study of the Safety and Efficacy of GevokizumAb in the Treatment of Subjects with Non-infectious Intermediate, Posterior, or Pan- uveitis Currently Controlled with Systemic Treatment Ophthalmic AGN195263 for the Treatment of Meibomian Gland Dysfunction 2013-2014 A Multi-Center, Double-Masked, Randomized, Placebo Controlled Study of EBI-005 2014- present Ophthalmic Solution versus Placebo in Subjects with Dry Eve Disease 2014-present A Double-Masked, Randomized, Multicenter, Placebo Controlled, Parallel-Group Study of SJP-0035 Ophthalmic Solution Compared With Placebo to Assess Safety and Efficacy of SJP-0035 Ophthalmic Solution for Corneal Epithelial Wound Healing in Patients With Moderate to Severe Corneal Epithelial Disorders A Phase III, Double-Masked, Randomized, Controlled Trial of KPI-121 in 2014-present Postsurgical Inflammation A Phase 2, Double-Masked, Randomized, Vehicle Controlled Study to Evaluate 2014-present the Effect of KPI-121 0.25% Ophthalmic Suspension on Signs and Symptoms of Inflammatory Meibomian Gland Disease 2014-present Dry Eye Assessment and Management (DREAM) Study (NIH Sponsored Trial) 2014-present A Phase 3, Multicenter, Randomized, Double-masked, and Placebo-controlled Study Evaluating the Efficacy and Safety of a 5.0% Concentration of Lifitegrast Ophthalmic Solution Compared to Placebo in Subjects with Dry Eye Disease 2014-present A Multicenter, Double-Masked, Parallel-Group, Vehicle Controlled Study to Assess the Efficacy and Safety of Rx-10045 Nanomicellar Ophthalmic Solution for Treatment of Ocular Inflammation And Pain In Subjects Undergoing Cataract Surgery A Randomized, Multicenter, Double-Masked, Vehicle Controlled, Dose-Ranging 2014-present Study of the Safety and Efficacy of OTX-101 in the Treatment of Keratoconjunctivitis Sicca 2014-present Evaluation of a Lubricant Eye Drop on Tear Lipid Layer Thickness in Contact Lens 2015-present A Phase 2 Randomized, Investigator-Masked, Comparator-controlled Trial to Evaluate the Safety and Efficacy of NS2 Eye Drops in Patients with Anterior Uveitis 2015-present A Phase 3, Multicenter, Randomized, Double-masked, and Placebo Controlled Study Evaluating the Efficacy and Safety of a 5.0% Concentration of Lifitegrast Ophthalmic Solution Compared to Placebo in Subjects with Dry Eye Disease and History of Recent Artificial Tear Use