#### **CURRICULUM VITAE**

# Harinderjit Singh, M.D.

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# **INSTITUTIONAL AFFLIATIONS**

Clinical Associate Professor of Ophthalmology, Medical College of Georgia. Retired

## **EDUCATION**

*Botany, Zoology, Chemistry High School		1965
Higher Secondary School, Sector 7, Bhilai (M.P.) India		
*St. Xavier's College, Ranchi, India	B.S.C.	1965-1968
*Gandhi Medical College, Bhopal, India	M.B.B.S	1968-1973
*Rotating Intern, General Hospital, Bhilai, India		1973-1974
*Resident in Nephrology, University Hospital,		1974-1975
The University of Western Ontario, London, Ontario, Canada		
*Straight Intern in Medicine. The University of Western Ontario Hospitals		1975-1976
London, Ontario, Canada		
*Rotating Intern, Reddy Memorial Hospital,		1976-1977
McGill University Teaching Hospital, Montreal, Quebec, Canada		
*Resident in Ophthalmology. The University of Western Ontario Hospitals		1977-1980
London, Ontario, Canada		
*Fellow in Diseases of the Retina and Vitreous, Retina Consultants, Ltd.,		1980-1981
Barnes Hospital, Washington University, St. Louis, Missouri		

#### MERITS AND AWARDS.

- Final BSc from Ranchi University: 9th in Merit list of Ranchi University
- First MBBS exam Vikram University: 8<sup>th</sup> in Merit List.
- Final MBBS exam, Bhopal University: 7<sup>th</sup> in Merit List.
- Awarded best teacher for Residents. Dept. of Ophthalmology, Medical College of Georgia, Augusta, GA in 1994-95.
- Best Doctors Award every year from 1999 to 2017.
- Top Ophthalmologist of US from 2014 to 2017

#### **PROFESSIONAL**

\*Instructor in Anatomy, University of Western Ontario, London, Ontario, Canada 1974-1975

\*Associate Clinical Professor of Ophthalmology, Medical College of Georgia, 1988-2017 Augusta, Georgia

## **PRACTICE EXPERIENCE**

1982-1983	Solo Practice, Saskatchewan, Canada
	Specializing in Disease and Surgery of the Retina and Vitreous
1983-1984	Group Practice, Odessa, Texas
	Specializing in Diseases and Surgery of the Retina and Vitreous
1984- 2017	Group Practice- Southeast Retina Center, P.C., Augusta, Georgia
	Specializing in Diseases and Surgery of the Retina and Vitreous –
	Partner/Surgeon. I started this practice in 1984 and retired from this practice in
	2017. In 1999 I also started a free standing Retina only outpatient Surgical center
	adjoining the office.
	2020 Jan to present: working Part-time at Augusta VA

#### **CURRENT ACTIVE MEDICAL LICENSES:**

GEORGIA 26652 TEXAS G4404 NEW YORK 145017-1 DEA FS 7706837 (REGISTERED FOR TEXAS)

# **SCIENTIFIC AND PROFESSIONAL SOCIETIES**

Member of Canadian Ophthalmology Society
Member of American Academy of Ophthalmology-1982
Fellow of the Royal College of Physicians and Surgeons (Canada)
Member of the All India Ophthalmology Society
Member of the American Society of Retina Specialist
Member of Canadian Society of Retina.
Member of the Paul Cibis Club

#### **MEETINGS, VISITING PROFESSORSHIPS**

- \* Indocyanine Green Angiography, University of South Carolina, Columbia, South Carolina
- \* Amritsar Ophthalmology Society, Keynote Speaker, December 2009
- \* Department of Ophthalmology, PGI, Chandigarh, India
- \* Grewal Eye Institute, Chandigarh, India
- \* Gandhi Medical College, Bhopal, India
- \* All India Ophthalmological Society
- \* Vitreo-Retina Society of India
- \* Punjab Ophthalmological Society, Punjab, India

# **CO-INVESTIGATOR IN THE FOLLOWING CLINICAL TRIALS:**

- 2000 Novartis: Phase III of Visudyne treatment with classic choroidal neovascular membrane
- 2001 Genentech, Inc.: Anti-VEGF Monoclonal antibody Fragment (rhuFab V2)
- 2003 Eli Lilly and Company, Protocol B7A-MC-MBDL: Reduction in the Occurrence of Center-Threatening Diabetic Macular Edema.
- 2003 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: A Pilot Study of Laser Photocoagulation for Diabetic Macular Edema.
- 2003 Genentech: Phase III, Placebo-Controlled, Multi-Center Study to Examine the Safety and Efficacy of Multiple-Dose Intravitreal Injections of the rhuFab V2 in Subjects with Age-Related Macular Degeneration Who Have Predominant Classic Lesion Degeneration.
- 2004 Alcon, Inc. and Bayer, Inc.: Determination of Intravitreal and Aqueous Concentrations of Moxifloxacin after Oral Administration in Scheduled Vitrectomy Patients.
- Novartis/QLT Inc.: A Randomized, Placebo-Controlled, Double Masked, Multicenter, Phase III Study of the Effect of Visudyne Therapy in Minimally Classic Subfoveal Choroidal Neovascularization (CNV) Secondary to Age-Related Macular Degeneration (AMD): Visudyne in Minimally Classic (VMC).
- 2004 EyeTech Pharmaceuticals, Anti-VEGF Pegylated Aptamer (EYE001): Intravitreal Injections for Exudative ARMD- Phase II/III Trial.
- 2004 Boehringer Ingelheim Pharmaceuticals, Inc.: A two year open label, randomized, parallel group, blinded assessment ophthalmologic safety study of pramipexole IR versus ropinirole in early Parkinson's Disease patients (Mirapex study).
- 2004 Family Investigation of Nephropathy and Diabetes, The FIND Eye Study: Identifying Genes Responsible for Diabetic Nephropathy and their Linkage Relationships to Nephropathy (FIND).
- 2005 Genaera MSI-1256F-301/302: A Phase 3 Multicenter, Randomized, Double-Masked, Controlled Study of Squalamine Lacate (MSI-1256F) for Injection for the Treatment of Subfoveal Choroidal Neovascularization Associated with Age-Related Macular Degeneration.
- 2005 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: A Randomized Trail Comparing Intravitreal Triamcinolone Acetonide and Laser Photocoagulation for Diabetic Macular Edema (Protocol B).
- 2005 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Phase 2 Evaluation of Anti-VEGF Therapy for Diabetic Macular Edema: Bevacizumab (Avastin). (Protocol H)

- 2005 EyeTech Pharmaceuticals, Inc.- A Phase 2/3 Randomized, Controlled, Double-Masked, Multi-Center, Comparative Dose-Finding Trial, in Parallel Groups, to Compare the Safety and Efficacy of Intravitreous Injections of 0.3, 0.03 or 0.003 mg Pegaptanib Sodium (Macugen), Given as Often as Every 6 Weeks for 3 years, to Sham Injections, in Subjects with Diabetic Macular Edema (DME) Involving the Center of the Macula.
- 2005 Lilly Research Laboratories, Protocol B7A-MC-MBDL: Reduction in the Occurrence of Center-Threatening Diabetic Macular Edema
- 2005 Regeneron VGFT-OD-0508: A Randomized, Controlled Study of the Safety, Tolerability and Biological Effect of Repeated Intravitreal Administration of VEGF Trap in Patients with Neovascular Age-Related Macular Degeneration.
- Genentech, Inc. FVF3689g: A Phase IIIB, Single-Masked, Multicenter, Randomized Study to Evaluate the Safety and Tolerability of Ranibizumab in Naïve and Previously Treated Subjects with Choroidal Neovascularization (CNV) Secondary to Age-Related Macular Degeneration (AMD) (SAILOR Study).
- 2005 Schering-Plough Corporation: Determination of Intravitreal Concentrations after Oral Administration of Moxifloxacin in Scheduled Vitrectomy Patients.
- Genentech, Inc. FVF3761s: Treatment of Polypoidal Choroidal Vasculopathy with Ranibizumab (Lucentis): A Phase I/II Safety Study (Poly Study).
- 2005 Jaeb Center for Health Research, National Eye Institute: Two Randomized Trials to Compare the Efficacy and Safety of Intravitreal Injection(s) of Triamcinolone Acetonide with Standard Care to treat Macular Edema Associated with Central Retinal Vein Occlusion and Branch Retinal Vein Occlusion (SCORE Study).
- Alimera C-01-05-001: A Randomized, Double-Masked, Parallel Group, Multi-center, Dose-Finding Comparison of the Safety and Efficacy of ASI-001A 0.5mg/day and ASI-001B 0.2mg/day Fluocinolone Acetonide Intravitreal Inserts to Sham Injection in Subjects with Diabetic Macular Edema (FAME).
- EyeTech Pharmaceuticals, Inc. Protocol EOP1023: A Phase IV, Open Label, Multi-Center, Trial of Maintenance Intravitreous Injections of Macugen® (Pegaptanib Sodium) Given Every 6 Weeks for 48 Weeks in Subjects with Subfoveal Neovascular Age-Related Macular Degeneration (AMD) Initially Treated with a Modality Resulting in Maculopathy Improvement (Level Study).
- 2007 Regeneron VGFT-OD-0702: An Open-Label, Long-Term, Safety, and Tolerability Study of Intravitreal VEGF Trap-Eye in Subjects with Neovascular Age-Related Macular Degeneration (Extension of Regeneron protocol 0508).

- 2007 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Intravitreal Ranibizumab or Triamcinolone Acetonide in Combination with Laser Photocoagulation for Diabetic Macular Edema (Protocol I).
- 2007 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Intravitreal Ranibizumab or Triamcinolone Acetonide as Adjunctive Treatment to Panretinal Photocoagulation for Proliferative and Non-Proliferative Diabetic Retinopathy (Protocol J).
- 2007 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: The Course of Response to Focal Photocoagulation for Diabetic Macular Edema (Protocol K).
- 2007 Genentech, Inc. FVF4165g: A Phase III, Double-Masked, Multicenter, Randomized, Sham Injection Controlled Study of the Efficacy and Safety of Ranibizumab Injection Compared With Sham in Subjects with Macular Edema Secondary to Branch Retinal Vein Occlusion (Bravo Study).
- 2007 Genentech, Inc. FVF4166g: A Phase III, Double-Masked, Multicenter, Randomized, Sham Injection Controlled Study of the Efficacy and Safety of Ranibizumab Injection Compared With Sham in Subjects with Macular Edema Secondary to Central Retinal Vein Occlusion (Cruise Study).
- 2007 Genentech, Inc. FVF4168g: A Phase III, Double-Masked, Multicenter, Randomized, Sham Injection Controlled Study of the Efficacy and Safety of Ranibizumab Injection in Subjects with Clinically Significant Macular Edema with Center Involvement Secondary to Diabetes Mellitus (Ride Study).
- 2007 Eli Lilly and Company, Protocol B7A-MC-MBCU: The Effect of Ruboxistaurin on Clinically Significant Macular Edema in Patients with Diabetes Mellitus, as assessed by Optical Coherence Tomography (MBCU Study).
- Opko Health, Inc. Protocol ACU301C: A Phase 3, Randomized, Double-masked, Parallel-Assignment study of Intravitreal Bevasiranib Sodium, Administered Every 8 or 12 Weeks as Maintenance Therapy Following Three Injections of Lucentis®, Compared with Lucentis® Monotherapy Every 4 Weeks in Patients with Exudative Age-Related Macular Degeneration (AMD) (Cobalt Study).
- 2007 Regeneron Pharmaceuticals, Inc., Protocol VGFT-OD-0605: A Randomized, Double-masked, Active Controlled Phase III Study of the Efficacy, Safety and Tolerability of Repeated Doses of Intravitreal VEGF Trap in Subjects with Neovascular Age-Related Macular Degeneration (View 1 Study).
- 2007 RegenRx Biopharmaceuticals, Inc. Protocol RGN-DV-201: A Randomized, Double-Mask, Placebo-Controlled, Dose-Response, Phase 2 Study of the Safety and Efficacy of Thymosin Beta 4 in the Treatment of Diabetic Patients' Corneal Wounds Resulting from Epithelial Debridement During Vitrectomy.

- 2008 Allergan, Protocol 206207-012: A 52-Week, Masked, Multicenter, Randomized, Controlled Trial (With Up to 13 Weeks Additional Follow-Up) to Assess the Safety and Efficacy of 700 µg Dexamethasone Posterior Segment Drug Delivery System (DEX PS DDS) Applicator System in Combination with Laser Photocoagulation Compared with Laser Photocoagulation Alone in the Treatment of Subjects with Diffuse Macular Edema (DME).
- 2008 Neovista, Protocol NVI-114-08: A Randomized, Prospective, Active Controlled, Study of the Epi-Rad Ophthalmic System for the Treatment of Subfoveal Choroidal Neovascularization Associated with Wet Age-Related Macular Degeneration (Cabernet).
- 2008 Pfizer. A Phase II, Prospective, Randomized, Multi-Center, Diabetic Macular Edema Dose-Ranging, Comparator Study Evaluating the Efficacy and Safety of PF-04523655 versus Laser Therapy (DEGAS).
- Ophthotech: A Phase I, Ascending Dose and Parallel Group Trial to Establish the Safety, Tolerability and Pharmacokinetic Profile of Multiple Intravitreous Injections of Volociximab (α5β1 Integrin Antagonist) as Monotherapy or in Combinations with Lucentis 0.5 mg/eye in Subjects with Neovascular Age-Related macular Degeneration.
- Genentech FVF3426g: An Open Label, Multi-Center Extension Study to Evaluate the Safety and Tolerability of Ranibizumab in Subjects with Choroidal Neovascularization (CNV) Secondary to Age-Related Macular Edema Secondary or Retinal Vein Occlusion (RVO) who have Completed a Genentech Sponsored Ranibizumab Study (Horizon).
- 2009 Thrombogenics: A Randomized, Placebo Controlled, Double-Masked, Multi-Center Trial of Microplasmin Intravitreal Injection for Non-surgical Treatment of Focal Vitreomacular Adhesion.
- 2009 Genentech FVF4579g: A Phase III, Double-Masked, Multicenter, Randomized, Active Treatment-Controlled Study of The Efficacy and Safety of 0.5mg and 2.0mg Ranibizumab Administered Monthly or On An As-Needed Basis (PRN) In Patients with Subfoveal Neovascular Age-Related Macular Degeneration (HARBOR Study).
- 2009 Regeneron Pharmaceuticals, VGFT-OD-0706: A Double-Masked, Randomized, Controlled Study of the Safety, Tolerability and Biological Effect of Repeated Intravitreal Administration of VEGF Trap-Eye in Patients with Diabetic Macular Edema.
- 2009 Regeneron Pharmaceuticals, VGFT-OD-0819: A Randomized, Double masked, Controlled Phase 3 Study of the Efficacy, Safety, and Tolerability of Repeated Intravitreal Administration of VEGF Trap-Eye in Subjects with Macular Edema Secondary to Central Retinal Vein Occlusion.
- 2009 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Comparison of Time Domain OCT and Spectral Domain OCT Retinal Thickness Measurement in Diabetic Macular Edema (Protocol O).

- 2009 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: A Pilot Study in Individuals with Center-Involved Diabetic Macular Edema Undergoing Cataract Surgery (Protocol P).
- 2009 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: An Observational Study in Individuals with Diabetic Retinopathy without Center-Involved DME Undergoing Cataract Surgery (Protocol Q).
- 2009 Pfizer, Protocol A1451259: A Non-Treatment Study of Risk Factors for Nonarteritic Anterior Ischemic Optic Neuropathy (NAION).
- 2010 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: An Evaluation of Intravitreal Ranibizumab for Vitreous Hemorrhage Due to Proliferative Diabetic Retinopathy (Protocol N).
- 2010 Pfizer, Protocol B1181002: A Phase 1, Double-Masked, Placebo-Controlled Study Evaluating the Safety, Tolerability, Immunogenicity, Pharmacokinetics and Pharmacodynamics of Multiple Escalating Dosages of RN6G In Subjects With advanced Dry, Age-Related Macular Degeneration (AMD) Including Geographic Atrophy.
- Juvenile Diabetes Research Foundation: Ranibizumab for Edema of the Macula in Diabetes: Protocol 3 with High Dose- the Read 3 Study.
- 2010 The Macula Foundation with Genentech, Inc., Protocol FVF4926s: Extended Follow-up of Patients with Macular Edema due to Branch Retinal Vein Occlusion (BRVO) or Central Retinal Vein Occlusion (CRVO) previously treated with Intravitreal Ranibizumab (Retain).
- 2010 Regeneron Pharmaceuticals, VGFT-OD-0910: An Open-Label, Long-Term, Safety and Tolerability Extension Study of Intravitreal VEGF Trap-Eye in Neovascular Age-Related Macular Degeneration (View 1 Extension).
- 2011 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: A Phase II Evaluation of Topical NSAIDs in Eyes with Non-Central Involved DME (Protocol R).
- 2011 Genentech, Inc., Protocol FVF4967g: A Multicenter Randomized Study Evaluating Dosing Regimens for Treatment with Intravitreal Ranibizumab Injections in Subjects with Macular Edema Following Retinal Vein Occlusion (SHORE).
- 2011 Santen, Inc., Protocol 32-007: A Phase III, Multinational, Multicenter, Randomized, Double-Masked Study Assessing the Safety and Efficacy of Intravitreal Injections of DE-109 (three doses) for the Treatment of Active, Non-Infections Uveitis of the Posterior Segment of the Eye (Sakura).
- 2011 Regeneron Pharmaceuticals, Protocol VGFT-OD-1009: A Double-Masked, Randomized, Active-Controlled, Phase 3 Study of the Efficacy and Safety of Intravitreal Administration of VEGF Trap-Eye in Patients with Diabetic Macular Edema (Vista).

- 2011 Genentech, Inc., Protocol CFD4870g: A Phase Ib/II, Multicenter, Randomized, Single-Masked, Sham Injection-controlled Study of Safety, Tolerability, and Evidence of Activity of FCFD4514S Intravitreal Injections Administered Monthly or Every Other Month to Patients with Geographic Atrophy (Mahalo).
- Alimera, Protocol C-01-11-008: An Open-label, Multi-center Extension Study of the Safety and Utility of the New Inserter of Iluvien (fluocinolone acetonide intravitreal insert) 0.19mg in Subjects and the Safety of Iluvien in Subjects with Macular Edema (FAME Extension).
- Genentech, Inc., Protocol FVF4916s: Treatment of Polypoidal Chorodial Vasculopathy with High Dose Ranibizumab (Lucentis): A Phase I Safety Study (High Tide Poly).
- 2011 Lpath, Inc., Protocol LT1009-Oph-003: A Phase 2A, Multi-center, Masked, Randomized, Comparator-Controlled Study Evaluating iSONEP (Sonepcizumab [LT1009]) as either Monotherapy or Adjunctive Therapy to Lucentis of Avastin Alone for the Treatment of Subjects with Choroidal Neovascularization Secondary to Age-Related Macular Degeneration.
- 2012 Acucela Inc., Protocol 4429-202: A Phase 2b/3 Multicenter, Randomized, Double-Masked, Dose-Ranging Study Comparing the Efficacy and Safety of Emixustat Hydrochloride (ACU-4429) with Placebo for the Treatment of Geographic Atrophy Associated with Dry-Age-Related Macular Degeneration.
- 2012 Alcon Research Ltd., Protocol C-12-006: A Prospective, Randomized, Double-Masked, Multicenter, Two Arm Study Comparing the Efficacy and Safety of ESBA1008 versus EYLEA in Subjects with Exudative Age-Related Macular Degeneration (OSPREY).
- 2012 Genentech, Protocol GX28198: A Multicenter, Open-Label Extension Study to Evaluate the Long-term Safety and Tolerability of FCFD4514S in Patients with Geographic Atrophy (OLEi).
- 2012 Genentech, Protocol ML28713: Ranibizumab For Persisent Diabetic Macular Edema After Bevacizumab (ROTATE Trial).
- GlaxoSmithKline, Protocol BAM114341: A Phase 2, Multi-centre, Randomised, Double-masked, Placebo-controlled, Parallel-group Study to Investigate the Safety, Tolerability, Efficacy, Pharmacokinetics and Parmacodynamics of GSK933776 in Adult Patients with Geographic Atrophy (GA) Secondary to Age-related Macular Degeneration (AMD).
- 2012 Jaeb Center for Health Research: A Validation Study of the Electronic ETDRS Test Protocol for Low-Contrast Visual Acuity Testing in Adults and Older Children (EVA Low-Contrast VA)
- 2012 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: A Comparative Effectiveness Study of Intravitreal Aflibercept, Bevacizumab and Ranibizumab for Diabetic Macular Edema (Protocol T).

- 2012 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Prompt Panretinal Photocoagulation versus Intravitreal Ranibizumab with Deferred Panretinal Photocoagulation for Proliferative Diabetic Retinopathy (Protocol S).
- 2012 Pfizer: Protocol B1181003: A Phase 2 Multi-Center, Randomized, Double-Masked, Placebo-Controlled, Multidose Study to Investigate the Efficacy, Safety, Pharmacokinetics and Pharmacodynamics of RN6G (PF-04382923) in Subjects with Geographic Atrophy Secondary to Age-Related Macular Degeneration.
- 2012 QDR, Protocol 2010-00703 DME: A Randomized, Multi-center, Phase II Study of the Safety, Tolerability, and Bioactivity of Repeated Intravitreal Injections of iCo-007 as Monotherapy or in Combination with Ranibizumab or Laser Photocoagulation in the Treatment of Diabetic Macular Edema with Involvement of the Foveal Center (iDeal).
- 2012 Regeneron: Intravitreal Aflibercept Injection for the treatment of CHoroidAl Neovascularization seconDary to presumed ocuLar histoplasmosis syndromE (the HANDLE study). Principal Investigator.
- 2012 Regeneron, Protocol VGFTe-AMD-1124: An Open-Label Study of the Efficacy, Safety, and Tolerability of Intravitreal Administration of VEGF Trap-Eye (Intravitreal Aflibercept Injection) in Patients with Neovascular Age-Related Macular Degeneration (RE-VIEW).
- 2012 Regeneron: Protocol VGFTe-RVO-1027: A Double-Masked, Randomized, Active-Controlled Study of the Efficacy, Safety and tolerability of Intravitreal Administration of VEGF Trap-Eye (Intravitreal Aflibercept Injection [IAI]) in Patients with Macular Edema Secondary to Branch Retinal Vein Occlusion (Vibrant).
- 2012 Quark Pharmaceuticals, Protocol QRK202: An Open-Label Dose Escalation Study of PF-04523655 (Stratum I) Combined with a Prospective, Randomized, Double-Masked, Multi-Center, Controlled Study (Stratum II) Evaluating the Efficacy and Safety of PF-04523655 Alone and in Combination with Ranibizumab Versus Ranibizumab Alone in Diabetic Macular Edema (Matisse).
- 2013 Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Treatment for Central-Involved Diabetic Macular Edema in Eyes with Very Good Visual Acuity (Protocol V).
- 2013 Pfizer, Protocol B1261009: A Phase 2 Randomized, Double-Masked, Placebo-Controlled, Parallel Group, Multi-Center Study to Compare the Efficacy and Safety of a Chemokine CCR2/5 Receptor Antagonist (PF-04634817) with that of Ranibizumab in Adult Subjects with Diabetic Macular Edema.
- Ophthotech, Protocol OPH1003: A Phase 3 Randomized, Double-Masked, Controlled Trial to Establish the Safety and Efficacy of Intravitreous Administration of Fovista<sub>TM</sub> (Anti PDFT-B Pegylated Aptamer) Administered in Combination with Lucentis® Compared to Lucentis® Monotherapy in Subjects with Subfoveal Neovascular Age-Related Macular Degeneration.

- 2013 Regeneron: Treat and Extend Therapy Using Intravitreal Aflibercept (IAI) for Previously Treated Patients Exiting the Wet Age-Related Macular Degeneration Extension Study (0910) RANGE.
- Jaeb Center for Health, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Short-term Evaluation of Combination Corticosteroid+Anti-VEGF Treatment for Persistent Central-Involved Diabetic Macular Edema Following Anti-VEGF Therapy in Pseudophakic Eyes (Protocol U).
- 2014 Regeneron- Investigator Initiated Study: Intravitreal Aflifercept for Neovascular Polypoidal Choroidal Vasculopathy (RIVAL TRIAL).
- 2014 National Eye Institute: Study of COmparative Treatments for REtinal Vein Occlusion 2 (SCORE2).
- 2014- ThromboGenics: Ocriplasmin Research to Better Inform Treatment (ORBIT): A Phase IV Observational Study.
- 2014- Genentech: A Phase III, Multicenter, Randomized, Double-Masked, Sham-Controlled Study of Efficacy and Safety of 10 mg Lampalizumab Intravitreal Injections Administered Every 30 or 45 Days to Patients with Geographic Atrophy Secondary to Age-Related Macular Degeneration (CHROMA).
- 2014- Allergan: Evaluation of Abicipar Pegol (AGN-150998) in Patients with Decreased Vision Due to Diabetic Macular Edema.

- Ophthotech, Protocol OPH1005, Sub-Retinal Fibrosis in Neovascular AMD: A 24 Month Phase 2A Open Label Safety Study of Fovista® (Anti-PDGF-BB Pegylated Aptamer) Regimen Administered in Combination with Anti-VEGF Therapy (Avastin® or Eylea®) During the Induction and Maintenance Phase of Therapy.
- 2015- Alimera Sciences, Iluvien® (Fluocinolone Acetonide Intravitreal Implant) M-01-15-004, A Phase 4 Safety Study of IOP Signals in Patients Treated with Iluvien® 0.19 MG (PALADIN).
- A Phase 2 Randomized, Double-masked, Multicenter, Active-controlled Study Evaluating Administration of Repeated Intravitreal Doses of hI-con1™ in Patients with Choroidal Neovascularization Secondary to Age Related Macular Degeneration (EMERGE).
- 2015- 2016 Alcon Research, Ltd., Protocol RTH258-C001, Efficacy and Safety of RTH258 versus Aflibercept: A Two-Year, Randomized, Double-Masked, Multicenter, Three Arm Study Comparing the Efficacy and Safety of RTH258 versus Aflibercept in Subjects with Neovascular Age Related Macular Degeneration (HAWK).
- Ophthotech, Protocol OPH1006, Effect of Anti-VEGF Agents Administered on a Quarterly Maintenance Regimen in Subjects with Neovascular AMD Receiving Anti-PDGF Therapy: An 18 Month Phase 2A Open Label, Randomized Study of Avastin, Lucentis, or Eylea (Anti-VEGF Therapy) Administered in Combination with Fovista (Anti-PDGF BB Pegylated Aptamer).
- 2015- Allegro, Protocol DME 202B, A Phase 2 Multicenter, Randomized, Controlled, Double Masked Clinical Trial Designed to Evaluate the Safety and Exploratory Efficacy of Luminate (Alg-1001) As Compared to Avastin And Focal Laser Photocoagulation in The Treatment of Diabetic Macular Edema.
- 2015- Regeneron Pharmaceuticals, Protocol R2176-3-AMD-1417, A Phase 2, Double-Masked, Randomized, Controlled, Multiple-Dose, Regimen-Ranging Study of the Efficacy and Safety of Intravitreal REGN2176-3 In Patients with Neovascular Age-Related Macular Degeneration (CAPELLA).
- 2015- Genentech, Inc., Protocol GX28228, A Phase II, Multicenter, Randomized, Active Treatment-Controlled Study of The Efficacy and Safety of The Ranibizumab Port Delivery System for Sustained Delivery of Ranibizumab in Patients with Subfoveal Neovascular Age-Related Macular Degeneration (LADDER).
- 2015- Roche, A Multiple-Center, Multiple-Dose and Regimen, Randomized, Active Comparator Controlled, Double-Masked, Parallel Group, 36-Week Study to Investigate the Safety, Tolerability, Pharmacokinetics, and Efficacy of RO6867461 Administered Intravitreally in Patients with Choroidal Neovascularization Secondary to Age-Related Macular Degeneration (AVENUE).

- 2015- Tyrogenex, Protocol X82-OPH-201, A Randomized, Double-Masked, Placebo-Controlled, Dose-Finding, Non-inferiority Study of X-82 Plus Prn Eylea Compared to prn Eylea Monotherapy in Neovascular AMD.
- 2015- Regeneron Pharmaceuticals, Investigator Initiated Research Grant, Intravitreal Aflibercept Injection (IAI) for Persistent Diabetic Macular Edema (DME) after Treatment with Bevacizumab and/or Ranibizumab (ROTATED).
- 2016- Apellis Pharmaceuticals, A Phase II, Multicenter, Randomized, Single-Masked, Sham-Controlled Study of Safety, Tolerability and Evidence of Activity of Intravitreal APL-2 Therapy in Patients with Geographic Atrophy (GA). (FILLY).
- 2016- Regeneron Pharmaceuticals, A Phase 3, Double-Masked, Randomized Study of the Efficacy and Safety of Intravitreal Aflibercept Injection in Patients with Moderately Severe to Severe Nonproliferative Diabetic Retinopathy (PANORAMA).
- 2015-2016 Neurotech, A Multi-Center, Two-Stage, Open-Label Phase 1 and Randomized, Active Controlled, Masked Phase II Study to Evaluate the Safety and Efficacy of Intravitreal Implantation of NT-503-3 Encapsulated Cell Technology Compared with Eylea for the Treatment of Recurrent Subfoveal Choroidal Neovascularization (CNV) Secondary to Age-Related Macular Degeneration (AMD).
- Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network, Anti-VEGF for PDR/DME Prevention Study (PROTOCOL W).
- 2016- Astellas, A Phase 2, Double-Masked, Randomized, Active Controlled Study to Evaluate the Efficacy and Safety of ASP8232 in Reducing Central Retinal Thickness in Subjects with Diabetic Macular Edema (VIDI.) Principal Investigator.
- 2016- Regeneron Pharmaceuticals, Investigator Initiated Research Grant, Endolaserless Vitrectomy with Intravitreal Aflibercept Injection for Proliferative Diabetic Retinopathy-Related Vitreous Hemorrhage (LASER LESS).
- 2016- OHR Pharmaceuticals, A Phase III Study of the Efficacy and Safety of Squalamine Lactate Ophthalmic Solution, 0.2% Twice Daily in Subjects with Neovascular Age-Related Macular Degeneration (MAKO).
- 2016- Regeneron Pharmaceuticals, A Randomized, Double-Masked, Active-Controlled, Phase 2 Study of the Efficacy, Safety, and Tolerability of Repeated Doses of Intravitreal REGN910-3 in Patients with Diabetic Macular Edema (RUBY).
- 2016- Regeneron Pharmceuticals, A Randomized, Double-Masked, Active-Controlled Phase 2 Study of the Efficacy, Safety, and Tolerability of Repeated Doses of Intravitreal REGN910-3 in Patients with Neovascular Age-Related Macular Degeneration (ONYX).

2016-	Roche, A Multiple-Center, Multiple-Dose, Randomized, Active Comparator—Controlled, Double-Masked, Parallel Group, 36-Week Study to Investigate the Safety, Tolerability, Pharmacokinetics, and Efficacy of RO6867461 Administered Intravitreally in Patients with Diabetic Macular Edema (BOULEVARD).
2014-	Roche, A Phase III, Multicenter, Randomized, Double-Masked, Sham-Controlled Study of Efficacy and Safety of 10 mg Lampalizumab Intravitreal Injections Administered Every 30 or 45 Days to Patients with Geographic Atrophy Secondary to Age-Related Macular Degeneration (CHROMA).
2014- 2016	Thrombogenics, Ocriplasmin Research to Better Inform Treatment (ORBIT).
2014- 2015	Allergan, Evaluation of Abicipar Pegol (AGN-150998) in Patients with Decreased Vision Due to Diabetic Macular Edema (PALM).
2014-2015	Retina Assocaiates of Hawaii, Intravitreal Aflibercept Injection (Eylea) for Polypoidal Choroidal Vasculopathy with Hemorrhage or Exudation (EPIC).
2015-	Jaeb Center for Health Research, National Eye Institute, Diabetic Retinopathy Clinical Trial Network: Peripheral Diabetic Retinopathy (DR) Lesions on Ultrawid field Fundus Images and Risk of DR Worsening Over Time (PROTOCOL AA).

#### ABSTRACTS/PRESENTATIONS/PUBLICATIONS

Changes in Diabetic Retinopathy Through 2 years: Secondary Analysis of Randomized clinical trial Comparing Aflibercept, Bevacizumab, and Ranibizumab.

Susan B Bressler MD, Danni Liu MSPH; Adam Glassman; Barbara A Blodi; Alessandro A Castellarin, MD; Lee M, Jampool, MD; Paul Kaufman, MD; Michele Melia ScM; Harinderjit Singh, MD; John A. Wells MD; for DRCR network.

JAMA Ophthalmology April 2017,135(6) 558-568.

Intravitreal Aflibercept injection (IAI) for Persistent DME after treatment with Bevacizumab and /or Ranibizumab: ROTATED Trial 52 weeks Results; By Caitlen Taylor BS; Heather Frazier, BS; Priscilla Rex; William Marcus; David Starnes, BS; Harveen Walia, MS; Evin M Samy; Harinderjit Singh, MD; Robert A Lalane III, MD; Dennis M Marcus, MD. Presented at ARVO 2019.

Intravitreal Aflibercept for Neovascular Ployploidal Choroidal Vasculopathy in a Predominantly Non Asian Population (RIVAL Trial) Dennis M Marcus, MD; Farrooq; Frazier H; Marcus WM; Harinderjit Singh, MD; Published in Ophthalmic Surgery, laser and Imaging Retina Jan 2017 Vol 46 No 1.

# Endolaserless Vitrectomy With Intravitreal Aflibercept Injection for Proliferative Diabetic Retinopathy-Related Vitreous Hemorrhage (LASER LESS TRIAL)

Dennis M. Marcus, MD1, Harinderjit Singh, MD1, Davis C. Starnes, BS1, Harveen Walia, BS1, Amina Farooq, MD2, Heather Frazier, BS1, William B. Marcus1, and Robert A. Lalane, MD Published in Journal Of Vitreo Retinal Disease, April 2018

Endolaserless Vitrectomy with Intravitreal Aflibercept Injection (IAI) for Proliferative Diabetic Retinopathy (PDR)-Related Vitreous Hemorrhage (LASER LESS TRIAL): 44 Week Results. Marcus DM, Singh, Harinderjit; Farooq, Amina; Starnes, Davis; Walia, Harveen; Frazier, Heather, Lalane, RA. Presented at the Macula Society Meeting, Beverly Hills, CA. February 2018

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Endolaserless Vitrectomy with Intravitreal Aflibercept Injection (IAI) for Proliferative Diabetic Retinopathy (PDR)-Related Vitreous Hemorrhage: 1 Year Results (LASER LESS TRIAL). Dennis M. Marcus, MD, Harinderjit Singh, MD, Robert A. Lalane III, MD, Southeast Retina Center, Augusta, GA. Presented at ASRS annual meeting July 2018 Vancouver, British Columbia, Canada.

# Wide-Field Fluorescein Angiographic-Guided Aflibercept (WFFAGA) Monotherapy for Proliferative Diabetic Retinopathy (PDR)

Lindsay Williamson, BS¹; Davis Starnes, BS¹; Caitlen Taylor, BS¹; Rachel Levy, BA¹; Venkatkrish Kasetty, BS¹; Priscila Rex¹; Harinderjit Singh, MD¹; Robert A. Lalane, MD¹; Dennis M. Marcus, MD¹ Published in Journal of Vitro retinal Disease 2019.

Intravitreal Aflibercept for Neovascular Polypoidal Choroidal Vasculopathy In A Predominantly Non-Asian Population (RIVAL Trial). Marcus DM, Farooq A, Frazier H, Fechter CM, Marcus WB, Singh H. To be presented at the Annual Meeting of the Association for Research in Vision and Ophthalmology (ARVO), Seattle, Washington, May 2016.

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# **PUBLICATIONS IN NON-REFERRED JOURNALS**

Humoral and Cellular Presensitization in Transplantation: (unpublished data)

#### **CHARITABLE CONTRIBUTIONS**

2011- Present

Guru Granth Sahib Sikh Society, Chandigarh, India

- \* Attended Eye Camp in Chandigarh, India, seeing patients for consultation, February, 2011 to present . Visiting twice a year.
- \* Attended Eye Camp in Nanded, Maharashtra, seeing patients for consultation, performing surgery, injections, and laser for vitreoretinal and other ophthalmologic issues, and assessing patients for cataract surgeries.

  In this eye camp 150 Cataract surgeries were performed. From December 18-December 22, 2011

2008- Present Grewal Eye Institute, Chandigarh, India

- \* Visiting physician twice a year for consultation and vitreoretinal surgeries
- 2006 Jabalpur, Madhya Pradesh, India
  - \* Attended Eye Camp organized by the Lion's Club for two days. Performed eye exams and patients who needed vitreoretinal surgery were operated on in Chandigarh, India
- 1998 Al-Shifa Trust Eye Hospital, Rawalpindi Pakistan
  - \* Examined patients in consultation and examined for vitreoretinal diseases
- 1994- Present Guru Harkrishan Charitable Eye Hospital, Sohana Near Chandigarh, India.
  - \* Initial promoting member of hospital
  - \* Served as Trustee of the organization and was instrumental in designing the facility. Also supplied equipment worth millions of dollars to this charitable organization during this time period

\* Visited twice a year for 10 to 14 days to work in the hospital by examining patients, consulting and performing vitreoretinal surgeries

1994, 1997 Tribhuvan University Teaching Hospital, Kathmandu, Nepal

\* Participated in Eye Camps, seeing pts and performing Vitreo retina surgeries.