

**CURRICULUM VITAE**

**SCOTT G. FOXMAN, M.D.**  
**Retinal and Ophthalmic Consultants, P.C.**

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**OFFICE LOCATIONS:**

1500 Tilton Road, Northfield, NJ 08225

2466 East Chestnut Avenue, Vineland, NJ 08361

211 South Main Street, Cape May Court House, NJ 08210

**PROFESSIONAL EXPERIENCE:**

1986 - Present      Founder, President  
Retinal and Ophthalmic Consultants, P.C.

**EDUCATION:**

1973 - 1979      B.A. Boston University Medical Honors Program  
Boston University

1975-1979      M.D. Boston University Medical Honors Program  
Boston University

**POSTDOCTORAL RESEARCH:**

1979              National Eye Institute

**POSTGRADUATE TRAINING AND FELLOWSHIP APPOINTMENTS:**

1979-1980      Intern in Medicine  
Jackson Memorial Hospital

1980-1983      Resident in Ophthalmology  
University of Minnesota, Minneapolis

1983-1985      Fellowship, Vitreoretinal Diseases  
Jules Stein Eye Institute, UCLA  
Los Angeles, California

**FACULTY APPOINTMENTS:**

1983-1985	Visiting Lecturer Jules stein Eye Institute University of California
1986-Present	Courtesy Scheie Eye Institute University of Pennsylvania School of Medicine
1986-1996	Associate Attending Ophthalmologist Division of Ophthalmology AtlantiCare Regional Medical Center
1997-Present	Chief, Division of Ophthalmology AtlantiCare Regional Medical Center
1986-Present	Assistant Attending Ophthalmologist Division of Ophthalmology Shore Memorial Hospital

**SPECIALTY CERTIFICATION:**

1979-	Diplomat of the National Board of Medical Examiners
1984-	Diplomat of the American Board of Ophthalmology

**LICENSURE:** Pennsylvania, New Jersey, California

**PROFESSIONAL SOCIETIES:**

National Societies

- American Society of Retinal Specialists
- American Academy of Ophthalmology
- American Medical Association
- Association for Research in Vision and Ophthalmology
- American Board of Ophthalmology

Local societies

- Medical Society of New Jersey
- Medical Society of Atlantic County
- Academy of Medicine of New Jersey
- New Jersey Academy of Ophthalmology
- Retina Society of New Jersey

## **CLINICAL STAFF APPOINTMENTS:**

AtlantiCare Regional Medical Center  
Shore Memorial Hospital  
Scheie Eye Institute

## **ACADEMIC APPOINTMENTS:**

University of Pennsylvania, Courtesy Staff

## **BOARD POSITIONS:**

Past President, Atlantic County Medical Society  
Past President, Lions Club Vision Care Center

## **EDITORIAL POSITION:**

Past Business Editor, American Society of Retina Specialists

## **RESEARCH:**

### BAM 114341 – Sub-Investigator

Glaxo Smith Kline  
March 2013-Present

A phase II Multicenter, Randomized, Double-Masked, Placebo-Controlled, Parallel-Group Study to Investigate the Safety, Tolerability, Efficacy, Pharmacokinetics and Pharmacodynamics of GSK933776 in Adult Patients with Geographic Atrophy (GA) Secondary to Age-Related Macular Degeneration (AMD).

### VGFTe-AMD-1124 – Sub-Investigator

Regeneron  
January 2013-Present

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.

### PROTOCOL S – Sub-Investigator

DRCR.net  
July 2012-Present

Prompt Panretinal Photocoagulation versus Intravitreal Ranibizumab with Deferred Panretinal Photocoagulation for Proliferative Diabetic Retinopathy

VIBRANT – Sub-Investigator

Regeneron

June 2012-Present

Protocol: VGFT e-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

SAKURA-Principal Investigator

Santen Incorporated

2011-Present

Protocol No. 32-007:

A Phase III, Multinational, Multi-Center, Randomized, Double-Masked Study Assessing the Safety and Efficacy of Intravitreal Injections of DE-109 (three doses) for the Treatment of Active, Non-Infectious Uveitis of the Posterior Segment of the Eye.

VISTA – Sub-Investigator

Regeneron

2011-Present

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. PAREXEL #201601

SHORE-Principal Investigator

Genentech

2011-Present

BB-IND 8633-Protocol FVF4967g:

A Multi-Center Randomized Study Evaluating Dosing Regimens for Treatment with Intravitreal Ranibizumab Injections in Subjects with Macular Edema Following Retinal Vein Occlusion.

VIEW 1 EXTENSION - Sub-Investigator

Regeneron

2010-Present

Protocol: VGFT-OD-0910

Open-label, long-term, safety and tolerability extension study of intravitreal VEGF Trap-Eye in neovascular age-related macular degeneration

NAION - Principal Investigator

Covance

2009-Present

Protocol: 12912

Prospective case crossover study to assess whether PDE5 inhibitor exposure in men with erectile dysfunction increases the risk for the development of non-arteritic anterior ischemic optic neuropathy (NAION).

COPERNICUS - Sub-Investigator

Regeneron

2009-2011

Protocol: VGFT-OD-0819

Randomized, double-masked, controlled phase III study of the efficacy, safety and tolerability of repeated administration of VEGF Trap-Eye in subjects with macular edema secondary to central retinal vein occlusion (CRVO).

HARBOR - Principle Investigator

Genentech

2009-2012

Protocol: FVF4579G

Phase III, double-masked, multi-center, randomized, active treatment, controlled study of the efficacy and safety of 0.5 mg and 2.0 mg Ranibizumab; administered monthly or on an as-needed basis (PRN) in patients with subfoveal neovascular age-related macular degeneration.

RISE - Principal Investigator

Genentech

2008-2012

Protocol: FVF4170G

Three-year multi-center study with monthly visits for patients with diabetic macular edema comparing Lucentis with Standard of Care Laser treatments.

CRUISE-Principal Investigator

Genentech

2008-2010

Protocol: FVF41669

One-year multi-center study with monthly visits comparing Lucentis to sham procedure for patients with central retinal vein occlusion.

DAVINCI - Sub-investigator

Regeneron

2008-2010

Protocol: 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 (DME).

HORIZON - Principal Investigator

Genentech

2008-2010

Protocol: FVF3426G

An open-label, multicenter extension study to evaluate the safety and tolerability of ranibizumab in subjects with choroidal neovascularization (CNV) secondary to age-related macular degeneration (AMD) or macular edema secondary to retinal vein occlusion (RVO) who have completed a Genentech-sponsored ranizumab study.

VIEW1 - Sub-Investigator

Regeneron

2007-2010

Protocol: VGFT-OD-0605

Two-year multi-center study with monthly visits comparing VEGF-Trap against Lucentis for patients with newly-diagnosed age-related macular degeneration.

SIRIUS - Sub-Investigator

Allergan

2006-2008

Protocol: 211745-001-01

Two-year multi-center study comparing Sirna against Lucentis for age-related macular degeneration

NOTAL VISION HMP-V4 - Sub-Investigator

Notal Vision

2006-2007

Protocol: 20071801

Sensitivity of the Home Macular Perimeter (HMP) in detection of choroidal neovascularization (CNV) secondary to age-related macular degeneration (ARMD).

SAILOR-Principal Investigator

Genentech, Inc.  
2003-2007  
Protocol: FVF3689G

One-year multi-center study to evaluate the safety and tolerability of Lucentis in newly-diagnosed and previously-treated subjects with age-related macular degeneration.

Visudyne Photodynamic Therapy - Sub-Investigator

Ciba Vision  
1999-2000

Clinical Investigator for Visudyne Photodynamic Therapy study.

Flibanserin - Sub-Investigator

University of Pennsylvania and Boehringer Ingelheim (BI Trial No. 1997-1998)

Eight-week, double-blind, placebo-controlled Phase II study of various doses of flibanserin and paroxetine in patients with Major Depressive Disorder. Our role was to perform ophthalmic examinations to evaluate for potential ocular complications and side effects.

Pioglitazone - Sub-Investigator

Takeda America, Inc.  
1996

A double-blind placebo-controlled, randomized, dose-titration study to evaluate the safety and efficacy of pioglitazone, an oral hypoglycemic agent. Our role was to perform retinal examinations and evaluate retinal photographs for evidence of diabetic retinopathy.

Adefovir Dipivoxil (bis-POM PME A) - Sub-Investigator

NJCRI  
1996-1997

A placebo-controlled study of the safety and efficacy of adefovir dipivoxil (bis-POM PME A) in prolonged survival of HIV-infected individuals with a CD4+ cell count of  $\leq 100/\text{mm}^3$ . Our role was for ophthalmic evaluation with emphasis on the presence or absence of CMV retinitis.

Silicone Oil - Principal Investigator

Richard James Corporation  
1993-1996

Principal Investigator. Clinical investigation as to the utility and safety of silicone oil injection in the vitreous cavity for patients with complicated retinal detachment. Our role was to provide patient care including vitreoretinal surgery.

Perfluorocarbon Liquid - Principal Investigator

Retinal and Ophthalmic Consultants, P.C.  
1992-1996

Clinical investigation as to the utility and safety of intra-operative use of perfluorocarbon liquid into the vitreous cavity during surgical reattachment of the retina.

Sulfur hexafluoride and Perfluoropropane - Sub-Investigator

Retinal and Ophthalmic Consultants, P.C.  
1989-1994

Clinical investigation as to the utility and safety of long-acting gases in the repair of retinal detachments.

Diabetic Retinopathy

National Institutes of Health  
Early Treatment of Diabetic Retinopathy Study (ETDRS)  
1984-1985

Dr. Scott Foxman was investigator in landmark national collaborative randomized trial of laser treatment for diabetic retinopathy.

**PUBLICATIONS:**

Original Papers

Ballantine, E., Foxman, S., Gorden, P., Rogh, J.: Rarity of Diabetic Retinopathy in Patients with Acromegaly. Arch. Intern. Med. 1981; 141:1625-1627.

Foxman, S., Cameron, J.D.: The Clinical Implications of Bilateral Microphthalmos with Cyst: Report of a Fatal Case. Am. J. Ophthalmol. 1984; 97:632-638.

Foxman, S., Heckenlively, J., Bateman, J., Wirtschafter, J.: A Classification of Congenital and Early Onset Retinitis Pigmentosa. Arch. Ophthalmol. 1985; 103:1502-1506.



Foxman, S., Heckenlively, J., Sinclair, S.: Pigmented Paravenous Retinochoroidal Atrophy and Rubeola Retinopathy. *Am. J. Ophthalmol.* 1985; 99:605-606.

Fishman, M., Kerman, B., Foxman, S.: Intraocular Cysticercus: Migratory. Ossonig, K. (ed): Proceedings of the 10<sup>th</sup> SIDUO Congress, Netherlands, W. Junk, 1986.

Heckenlively, J., Foxman, S., Parelhoff, E.: Retinal Dystrophy and Macular Coloboma. *Doc Ophthalmol.* 1988:68:257-271.

Shields, J., Perez, N., Shields, C., Foxman, S., Foxman, B.: Simultaneous Choroidal and Brain Metastasis as Initial Manifestations of Lung Cancer. *Ophthalmic Surgery and Lasers.* 2002:35:323-325.

Foxman, S.: Finding the right fit: Considerations on both sides of the employment process. *Retina Times.* 2006:24:25-27.

Foxman, S.: Office Administrator: The guide of your practice's success. *Retina Times.* 2006:24:47-51.

Ferencz L, Lang J, Yeshurn I, Pollack A, Siegel R, Lifshitz T, Karp J et al. Toward earlier detection of choroidal neovascularization secondary to age-related macular degeneration. *Retina.* 2010 March.

Brown DM, Campochiaro PA, Singh RP, Li , Gray S, Saroj N, Rundle AC, Rubio RG, Murahashi WY; CRUISE Investigators. Ranibizimab for macular edema following central retinal vein occlusion: Six month primary end-point results of phase III study. *Ophthalmology.* 2010 June; 117(6):1125-1133.e1.