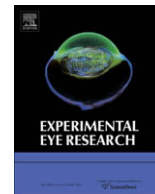


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Review

Douglas H. Johnson: A tribute

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It is both a distinct honor and yet a sad task to write a tribute for our very close friend and colleague, Dr. Douglas H. Johnson. One of us (MJ) knew Doug for nearly a quarter century, the other (MPF) for almost a decade. As many of you know, he was a rare individual, one that combined the tenacity, enthusiasm, wisdom and compassion of a dedicated scientist with the sensitivity and balance that we all treasured. Doug Johnson was a father, husband, son, scientist, clinician, traveler and friend to many of us.



1. Douglas H. Johnson: Clinician Scientist

Doug Johnson received his B.A. degree in biology from St. Olaf College, Northfield, MN in 1973 and M.D. from Mayo Medical School, Rochester, MN in 1977. He completed his medical

internship and residency in Ophthalmology at the Mayo Clinic before heading to Boston where he did his glaucoma fellowship with Dr. David Epstein at the Massachusetts Eye and Ear Infirmary (MEEI).



Meeting of Dr. David Epstein's lab group. Left to right: Levon Karaguezian, Tom Freddo, Joanne Sateriale, David Epstein, John Anderson, Mark Johnson, Doug Johnson, Mary Patterson, Jean Pei

It was at MEEI where Doug's love of research blossomed. The early 1980s saw a new team approach to the study of the pathogenesis of primary open-angle glaucoma (POAG). In Dr. David Epstein's laboratory biochemists, physiologists, engineers, morphologists and physicians were brought together to apply a new approach to solving this vexing problem. The goal was to achieve a "critical mass" of scientists that might unravel the causes of this mysterious disease. Doug loved this thrilling research environment, particularly the in-depth discussion of research topics, the open exchange of ideas, and the camaraderie of colleagues. This team oriented approach fostered his research and clinical practice throughout his professional career.

While in Boston, Doug first began to consider the possibility of developing a new method of perfusing eyes as part of these

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investigations into the cause of the obstruction of aqueous humor outflow, characteristic of POAG. Doug felt that there was a need for a new model system. As there was no generally accepted animal model that reproduces the intraocular pressure characteristic of primary open-angle glaucoma without other anterior segment changes, this suggested that human tissue needed to be used in experiments. Doug felt that cell culture was of limited utility, and he emphasized the importance of investigating the behavior of cells in situ. Vascular biology had been advanced by the isolated capillary model but no such model was available for studying aqueous humor outflow.

The best available model was the use of enucleated human eyes, but they were restricted to acute experiments (due to tissue necrosis) and some drugs that were known to affect outflow resistance in living eyes did not so affect enucleated eye (e.g. epinephrine). It was these considerations that led Doug to consider the possibility of organ culture in which anterior segments could be isolated and kept alive by perfusion of mock aqueous humor.

Doug discussed his idea with Dr. Morton Grant (developer of tonography, the method by which outflow resistance can be measured in living eyes, Grant, 1950, 1951). Dr. Grant was supportive of the idea and mentioned that Dr. Murray Johnstone had earlier at MEEI tried such a method, but found that flow through the tissue was dependent on how tightly these segments were clamped and not reproducible.

In 1983, Dr. Johnson returned to the Mayo Clinic as a Consultant in the Department of Ophthalmology (where he became full Professor in 1998), and pursued the development of an organ culture technique for the anterior segment of the eye. By 1985, with the aid of Dr. Richard Brubaker and several Mayo Clinic engineers, the perfusion organ culture of human anterior segments was established. In 1987, Doug's innovative idea had become a reality, with the publication of "Human trabecular meshwork organ culture: A new method." (Johnson and Tschumper, 1987). Over twenty years later, the perfusion organ culture of anterior segments is now used internationally to study the effects of pharmacologic agents, growth factors/cytokines, and other molecules on trabecular meshwork function and aqueous outflow resistance. It has been adapted for use with monkey, porcine, and bovine anterior segments and remains the model of choice for studying the conventional outflow pathway in human eyes.

Doug's passion was to understand the morphological architecture of the outflow pathway, particularly that of the trabecular meshwork. He felt that every image, whether taken with light, transmission or scanning electron microscopy, had the potential to teach us something about the function of the trabecular

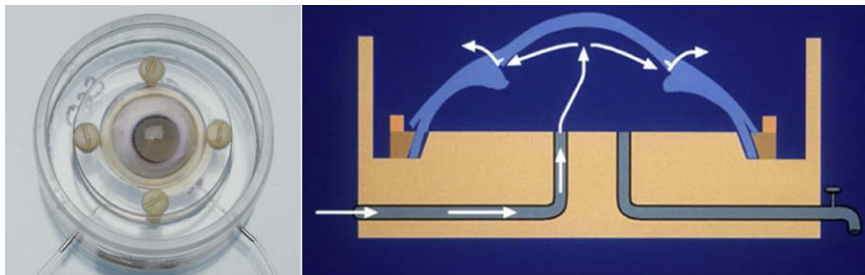


Doug encouraging Cindy Bahler (sitting) and Cheri Hann (standing). For over 15 years, both Cindy and Cheri were Doug's research technicians.

meshwork. He was a careful observer and described similarities and differences between the trabecular meshworks of normal eyes and those found in eyes with primary open-angle glaucoma, pseudoexfoliation glaucoma, pigmentary and steroid-induced glaucoma.

Recently, he described ultrastructural changes that followed nonpenetrating laser trabeculoplasty, where regions of the inner wall of Schlemm's canal and juxtacanalicular connective tissue between laser burns appeared "foamy or expanded" (Johnson, 2007). Doug felt that these "expanded" regions might represent tissue remodeling that served to lower flow resistance through the outflow pathway. He found similar structures under collector channels in normal eyes emphasizing their possible role in fluid flow. At the time when he passed, Doug was involved with characterizing the expanded regions in glaucomatous eyes, where he believed he might find pathophysiologic differences between the trabecular meshwork of normal and glaucomatous eyes, a task he had begun years earlier during an extended working vacation with Professor Elke Lutjen-Drecoll in Erlangen, Germany.

Doug's skill as a careful observer also served him well as a clinician. While evaluating the effectiveness of pilocarpine gel (Johnson et al., 1986), Doug was the first one to notice subtle, but unmistakable changes in the cornea caused by the delivery vehicle. This ability to carefully observe and then analyze data was an important tool that Doug repeatedly took advantage of.



(Left) Top view of anterior segment organ culture apparatus that Doug Johnson developed.

(Right) Schematic of fluid flow through organ culture model. (Johnson and Tschumper 1987)

Doug was a dedicated clinician, and came from a family of physicians; his father Herbert W. Johnson, MD practiced internal medicine St. Paul, MN, and his brother Stephen H. Johnson, MD is currently a practicing ophthalmologist in Newport Beach, CA. It was a pleasure to spend time with Doug in the clinic. No patient had a more caring and careful physician. While Doug's passion was to his research, his devotion was to his patients.

He was curious and wanted to find answers to questions that arose in his clinical practice, particularly why glaucomatous damage continued to progress in some patients even though intraocular pressure was significantly lowered. He spearheaded population-based studies that addressed questions ranging over the incidence of glaucoma, the probability that it would lead to blindness, and whether pigment dispersion syndrome ever converted to pigmentary glaucoma. He wanted to answer these questions both for his own scientific curiosity but also so that he could best inform and treat his patients.

Doug was able to blend his clinical work with his scientific studies. He examined the mechanism by which nonpenetrating glaucoma surgery procedures work to lower outflow resistance and concluded that such procedures unroof and expand Schlemm's canal, probably causing inadvertent ruptures of the inner wall and juxtacanalicular tissue, thus relieving the abnormal outflow resistance of glaucoma. In his own practice, he looked to improve his surgical skills and often tried new technologies such as trabecular stents to see what impact they had on aqueous outflow as well as the histological appearance of the outflow pathway.

In 2005, Doug was awarded the William and Betty MacMillan Professorship in Ophthalmology at the Mayo Clinic in recognition of his many scientific and teaching contributions. He served on the Editorial Boards for *Archives of Ophthalmology* (1994–2005) and *Experimental Eye Research* (2004–2006). At the time of his death, he was also a member of the National Advisory Eye Council.



After dinner following a glaucoma review meeting of the American Health Assistance Foundation. Left to right: Doug Johnson, Ross Ethier, David Epstein, Dan Stamer, Rand Allingham, Mark Johnson and Thom Freddo.

Among Doug's most cherished research moments were those involved in scientific discussion with colleagues late into the evening. They were highlights of his trips to ARVO, ICER and other scientific meetings. Doug Johnson was a long-time member of the National Glaucoma Research Scientific Advisory Committee of the American Health Assistance Foundation and served as the

committee chair since 1998. As Chairman, Doug made sure that late-night discussions and an open exchange of scientific ideas became an expected part of the annual meeting of this group. Most appropriately, and thanks to a wonderful suggestion from Thomas Freddo (Director of the School of Optometry, University of Waterloo), the American Health Assistance Foundation has now established the Dr. Douglas H. Johnson Award for Glaucoma Research.

2. Douglas H. Johnson: Teacher

Successful scientific investigation involves hard work, creativity, team management and well ... some luck. Doug Johnson taught us, and perhaps some of you, that research benefits greatly from objectivity. Doug always felt the hypothesis was the most important aspect of any research project. The best hypotheses were those crafted so that no bias was involved, avoiding personal attachment. Doug understood that you learn from an experiment whether the results support your hypothesis or not. Too many of us become wedded to our hypothesis at the exclusion of good scientific investigation. Follow the clues where they lead, not where you have planned that they ought to go. Doug felt himself a reporter of the truth rather than its architect. He felt that such objectivity could only be maintained if we kept our hypotheses at a distance. Doug succeeded at this goal. How many of us can say the same?

Doug was interested in whatever direction science led him, even though it frequently was not his planned course. Doug once ended an ARVO lecture on the penultimate slide of his presentation with the question "Does the pigmentation pattern in the trabecular meshwork of glaucomatous eyes differ from that of normal eyes?" (his thesis of that investigation). The final slide answered (in small type): "No." How many of us have couched negative results in a comfortable fashion? Doug simply followed the truth.



Doug and his family at Valerie's graduation from Whitman College in 2007. His wife Nancy (left), Valerie (middle) and Emily (far right).

3. Douglas H. Johnson: Husband and Father

In addition to his professional attributes, Doug also had many personal accomplishments. It was important for Doug to have a work-life balance, making sure to spend time with family and friends. Most important in his personal life was his family. Nancy (Nan) Johnson was the love of his life, and he felt the most fortunate man in the world to have found this love. The fruit of this wonderful relationship, Emily and Valerie, share their parent's passion for life and their father's scientific curiosity.



Doug playing his bagpipes in Edinburgh, Scotland

4. Douglas H. Johnson: The Person

Doug had several hobbies, but none more important than his bagpipes. Doug started playing bagpipes in 2001 and he never stopped. Doug performed with the Rochester Pipes and Drums band in concerts and parades. Individually, he would play for anyone that asked. He performed at laboratory functions, birthday parties, and anniversaries. He even showed up at Cindy Bahler's front door (Doug's laboratory technician for 15 years) to celebrate her 48th birthday, playing "Happy Birthday" on the bagpipes. He performed at the American Glaucoma Society (AGS), not for pleasure, but to make a point that forcibly exhaling against a closed or partially closed airway (Valsalva, demonstrated by the bagpipes) can affect intraocular pressure. We recall that for his funeral, Doug wrote a delightful and inspirational piece that was read to us, where he reminded us that bagpipes were to be expected at the event.

Doug also enjoyed traveling. Together with Nan, they traveled the world. Some trips were purely for fun such as travels with Nan, Emily, and Valerie to Europe to explore the Gothic cathedrals, Roman ruins and various wonders of history and nature. His visit to

Scotland was highlighted by playing in Pipefest 2005, a fund raiser for cancer research. Over 9000 bagpipers and drummers participated in the event, forming the largest ever pipe band and a place in the Guinness book of world records.

Most of these travels involved scientific pursuit as well, and Doug saw many of his colleagues on these trips. Before a trip they took together to Germany, Mark Johnson had ruptured his right achilles tendon and wore a cast on that trip. Doug frequently told the tale of an excessively heavy suitcase of Mark's due to unnecessary right shoes that need not have been brought. Furthermore, although Mark Johnson asked in a hotel in Erlangen, Germany for separate beds for Doug and him to sleep in (in fluent German so he thought), the presence of a double bed gave Doug more stories to expand upon as the years moved forward (it should be noted that Mark Johnson's recollection of these events varied somewhat from Doug's). However, it might be true that driving through Germany, passing round and round each Zentrum, they looked but could never find the "Zum" castle although they saw signs for it everywhere.

Doug had a wonderful sense of humor, whether on the giving or receiving end. Fortunately for Doug and for the rest of us, humor came naturally to him. Whether at a serious scientific symposium, after dinner talk, in the lab, or simply relating one of his tales to friends, Doug rarely failed to entertain. Many of us enjoyed hearing stories about ourselves, generally learning something new nearly every time he expanded on these special tales.

Doug was also a spiritual person. He taught Bible study for 15 years, helping others interpret what is meant and said in the Bible, and growing ever deeper in his own understanding and faith. In the piece that was read to us at his funeral, he described the basic but essential aspects to living a fun and worthwhile life: exercise, enthusiasm, and exhilaration. In his words "Exercise clears the mind and puts the days problems in perspective; enthusiasm about an enjoyable activity you can't wait to get to; and performing exhilarating tasks that years later you look back at and say wow, that was amazing." He wanted everyone to know that he lived life to the fullest, both personally and professionally. We can only hope that as we look back on our lives, we will be able to do what Doug did, live our lives to the fullest.

5. Doug Johnson: Our Friend

The field of Ophthalmology has lost an outstanding clinician scientist, but more importantly the world has lost a great person and we have lost a great friend. It was always Doug's dream to find the cause of glaucoma. Doug understood that multidisciplinary approaches to glaucoma, particularly POAG, were essential if we were to determine and understand the cause of the disease. It was



Sign to the Zum Castle in Germany that Doug Johnson and Mark Johnson looked for but never found. Photo courtesy of Marco Gößwein, photographer for Dr. Elke Lutjen-Drecoll.



our privilege as a scientific community to have known Doug and share our exchange of ideas. We miss his cheerful demeanor, positive analysis, and most importantly, his physical presence. We are sure that he would very much enjoy reading the diverse articles that make up this special tribute issue memorializing his many scientific investigations into the pathogenesis of glaucoma.

Publications

1. Johnson, D.H., 1976. Long-term anticoagulation following myocardial infarction. *Minn. Med.* 59, 333–337.
2. Johnson, D.H., Brubaker, R.F., 1982. Dynamics of aqueous humor in the syndrome of exfoliation with glaucoma. *Am. J. Ophthalmol.* 93, 629–634.
3. Johnson, D.H., Bourne, W.M., Campbell, R.J., 1982. The ultrastructure of Descemet's membrane. I. Changes with age in normal corneas. *Arch. Ophthalmol.* 100, 1942–1947.
4. Johnson, D.H., Bourne, W.M., Campbell, R.J., 1982. The ultrastructure of Descemet's membrane. II. Aphakic bullous keratopathy. *Arch. Ophthalmol.* 100, 1948–1951.
5. Bourne, W.M., Johnson, D.H., Campbell, R.J., 1982. The ultrastructure of Descemet's membrane. III. Fuchs' dystrophy. *Arch. Ophthalmol.* 100, 1952–1955.
6. Johnson, D.H., Liesegang, T.J., Brubaker, R.F., 1983. Aqueous humor dynamics in Fuchs' uveitis syndrome. *Am. J. Ophthalmol.* 95, 783–787.
7. Johnson, D.H., Epstein, D.L., Allen, R.C., Bays-Smith, J., Campbell, R., Rosenquist, R., Van Buskirk, M., 1984. A one-year multicenter clinical trial of pilocarpine gel. *Am. J. Ophthalmol.* 97, 723–729.
8. Johnson, D.H., 1985. Glaucoma: the search goes on. In: *Yearbook of Ophthalmology*, pp. 105–135.
9. Johnson, D.H., Kenyon, K.R., Epstein, D.L., Van Buskirk, E.M., 1986. Corneal changes with pilocarpine gel therapy. *Am. J. Ophthalmol.* 101, 13–15.
10. Johnson, D.H., 1986. New approaches to an old problem – the medical treatment of glaucoma. In: *Yearbook of Ophthalmology*, pp. 99–124.
11. Johnson, D.H., Brubaker, R.F., 1987. Glaucoma: an overview. *Mayo Clin. Proc.* 61, 59–67.
12. Johnson, D.H., Tschumper, R.C., 1987. Human trabecular meshwork organ culture. A new method. *Investig. Ophthalmol. Vis. Sci.* 28, 945–953.
13. Johnson, D.H., 1987. Automated perimetry, the blending of art and science. In: *Yearbook of Ophthalmology*, pp. 97–124.
14. Johnson, D.H., 1988. α , β and glaucoma, 1988. In: *Yearbook of Ophthalmology*, pp. 77–99.
15. Johnson, D.H., 1989. Does pigmentation affect trabecular meshwork cellularity? *Arch. Ophthalmol.* 107, 250–254.
16. Johnson, D.H., Tschumper, R.C., 1989. The effect of organ culture on human trabecular meshwork. *Exp. Eye Res.* 49, 113–127.
17. Johnson, D.H., Richardson, T.M., Epstein, D.L., 1989. Trabecular meshwork recovery after phagocytic challenge. *Curr. Eye Res.* 8, 1121–1130.
18. Dueker, D.K., Norberg, M., Johnson, D.H., Tschumper, R.C., Feeney-Burns, L., 1990. Stimulation of cell division by argon and NA:YAG laser trabeculoplasty in cynomolgus monkeys. *Investig. Ophthalmol. Vis. Sci.* 31, 115–124.
19. Johnson, M., Johnson, D.H., Kamm, R.A., deKater, A.W., Epstein, D.L., 1990. The filtration characteristics of the aqueous outflow system. *Exp. Eye Res.* 50, 407–418.
20. Tschumper, R.C., Johnson, D.H., Bradley, J.M.B., Acott, T., 1990. Glycosaminoglycans of human trabecular meshwork in perfusion organ culture. *Curr. Eye Res.* 9, 363–369.
21. Tschumper, R.C., Johnson, D.H., 1990. Trabecular meshwork cellularity: differences between fellow eyes. *Investig. Ophthalmol. Vis. Sci.* 31, 1327–1331.
22. Johnson, D., 1990. Extracapsular cataract extraction, intraocular lens implantation, and trabeculectomy: the combined procedure. *Int. Ophthalmol. Clin.* 30, 209–214.
23. Buller, C., Johnson, D.H., Tschumper, R.C., 1990. Human trabecular meshwork phagocytosis: observations in an organ culture system. *Investig. Ophthalmol. Vis. Sci.* 31, 2156–2163.
24. Johnson, D.H., Bradley, J., Acott, T., 1990. The effect of dexamethasone on glycosaminoglycans of human trabecular meshwork in perfusion organ culture. *Investig. Ophthalmol. Vis. Sci.* 31, 2568–2571.
25. Johnson, D.H., Brubaker, R.F., 1991. Glaucoma. In: Bartley, G.B., Liesegang, T.J. (Eds.), *Essentials of Ophthalmology*. J.B. Lippincott, pp. 109–129.
26. Kimpel, M., Johnson, D.H., 1992. Factors influencing in vivo trabecular cell replication as determined by ^3H -thymidine labelling. An autoradiographic study in cats. *Curr. Eye Res.* 11, 297–306.
27. Johnson, D.H., Tschumper, R.C., 1993. Ethacrynic acid: outflow effects and toxicity in human trabecular meshwork in perfusion organ culture. *Curr. Eye Res.* 12, 385–396.
28. Fan, J.T., Johnson, D.H., Burk, R.R., 1993. Transient myopia, angle-closure glaucoma, and choroidal detachment after oral acetazolamide. *Am. J. Ophthalmol.* 115, 813–814.
29. Johnson, D.H., Yoshikawa, K., Brubaker, R.F., Hodge, D.O., 1994. The effect of long-term medical therapy on the outcome of filtration surgery. *Am. J. Ophthalmol.* 117, 139–148.
30. Johnson, D.H., Knepper, P.A., 1994. Microscale analysis of the glycosaminoglycans of the human trabecular meshwork: a study in perfusion cultured eyes. *J. Glaucoma* 3, 58–69.
31. Buller, C.R., Johnson, D.H., 1994. Segmental variability of the trabecular meshwork in normal and glaucomatous eyes. *Investig. Ophthalmol. Vis. Sci.* 35, 3841–3851.
32. Johnson, D.H., 1994. The exfoliation syndrome – a continuing challenge. In: Albert, D.M., Jakobiec, F.A., Robinson, N.L. (Eds.), *Principles and Practice of Ophthalmology*. W.B. Saunders Co., Philadelphia, pp. 1400–1413.
33. Johnson, D.J., 1996. Management of glaucoma with poor surgical prognosis. In: *American Academy of Ophthalmology Focal Points*.
34. Ten Hulzen, R.D., Johnson, D.H., 1996. Effect of fixation pressure on juxtacanalicular tissue and Schlemm's canal. *Investig. Ophthalmol. Vis. Sci.* 37, 114–124.
35. FitzSimon, J.S., Johnson, D.H., 1996. Exfoliation material on intraocular lens implants. *Arch. Ophthalmol.* 114, 355.
36. Johnson, D.H., 1996. Human trabecular meshwork cell survival is dependent upon perfusion rate. *Investig. Ophthalmol. Vis. Sci.* 37, 1204–1208.
37. Russell, P., Johnson, D.H., 1996. Enzymes protective of oxidative damage present in all decades of life in the trabecular meshwork, as detected by two-dimensional gel electrophoresis protein maps. *J. Glaucoma* 5, 317–324.
38. Tamm, E.R., Russell, P., Johnson, D.H., Piatigorsky, J., 1996. Human and monkey trabecular meshwork accumulates αB -crystallin in response to heat shock and oxidative stress. *Investig. Ophthalmol. Vis. Sci.* 37, 2402–2413.
39. Johnson, D.H., 1996. Exfoliation syndrome. In: Albert, D.M., Jakobiec, F.A. (Eds.), *Atlas of Clinical Ophthalmology*. W.B. Saunders Co., Philadelphia, pp. 284–290.
40. Tomlinson, A.J., Benson, L.M., Jameson, S., Johnson, D.H., Naylor, S., 1997. Utility of membrane preconcentration-capillary electrophoresis-mass spectrometry in overcoming limited sample loading for analysis of biologically derived drug metabolite's peptides, and proteins. *J. Am. Soc. Mass. Spectrom.* 8, 15–24.

41. Johnson, D., Gottanka, J., Flügel, C., Hoffmann, F., Futa, R., Lütjen-Drecoll, E., 1997. Ultrastructural changes in the trabecular meshwork of human eyes treated with corticosteroids. *Arch. Ophthalmol.* 115, 375–383.
42. Johnson, D.H., 1997. Corticosteroid glaucoma. In: Epstein, D.L., Allingham, R.R., Schuman, J.S. (Eds.), *Chandler and Grants Glaucoma*. Williams & Wilkins, Baltimore, pp. 404–411.
43. Matsumoto, Y., Johnson, D.H., 1997. Trabecular meshwork phagocytosis in glaucomatous eyes. *Ophthalmologica* 211, 147–152.
44. Camp, J.J., Hann, C.R., Johnson, D.H., Tarara, J.E., Robb, R.A., 1997. Three-dimensional reconstruction of aqueous channels in human trabecular meshwork using light microscopy and confocal microscopy. *Scanning* 19, 258–263.
45. Gottanka, J., Johnson, D.H., Martus, P., Lütjen-Drecoll, E., 1997. Severity of optic nerve damage in eyes with POAG is correlated with changes in the trabecular meshwork. *J. Glaucoma* 6, 123–132.
46. Polansky, J.R., Fauss, D.J., Chen, P., Chen, H., Lütjen-Drecoll, E., Johnson, D., Kurtz, R.M., Ma, Z.-D., Bloom, E., Nguyen, T.D., 1997. Cellular pharmacology and molecular biology of the trabecular meshwork inducible glucocorticoid response gene product. *Ophthalmologica* 211, 126–139.
47. Matsumoto, Y., Johnson, D.H., 1997. Dexamethasone decreases phagocytosis by human trabecular meshwork cells in situ. *Investig. Ophthalmol. Vis. Sci.* 38, 1902–1907.
48. Gottanka, J., Flügel-Koch, C., Martus, P., Johnson, D.H., Lütjen-Drecoll, E., 1997. Correlation of pseudoexfoliative material and optic nerve damage in pseudoexfoliation syndrome. *Investig. Ophthalmol. Vis. Sci.* 38, 2435–2446.
49. Johnson, D.H., 1997. The effect of cytochalasin D on outflow facility and the trabecular meshwork of the human eye in perfusion organ culture. *Investig. Ophthalmol. Vis. Sci.* 38, 2790–2799.
50. Lütjen-Drecoll, E., May, C.A., Polansky, J.R., Johnson, D.H., Bloemendal, H., Nguyen, T.D., 1998. Localization of the stress proteins α B-crystallin and trabecular meshwork inducible glucocorticoid response protein in normal and glaucomatous trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 39, 517–525.
51. Nguyen, T.D., Chen, P., Huang, W.D., Chen, H., Johnson, D., Polansky, J.R., 1998. Gene structure and properties of TIGR, an olfactomedin-related glycoprotein cloned from glucocorticoid-induced trabecular meshwork cells. *J. Biol. Chem.* 273, 6341–6350.
52. Borrás, T., Matsumoto, Y., Epstein, D.L., Johnson, D.H., 1998. Gene transfer to the human trabecular meshwork by anterior segment perfusion. *Investig. Ophthalmol. Vis. Sci.* 39, 1503–1507.
53. Johnson, D.H., 1998. Options in the management of malignant glaucoma. *Arch. Ophthalmol.* 116, 799–800.
54. Hattenhauer, M.G., Johnson, D.H., Ing, H.H., Herman, D.C., Hodge, D.O., Yawn, B.P., Butterfield, L.C., Gray, D.T., 1998. The probability of blindness from open angle glaucoma. *Ophthalmology* 105, 2099–2104.
55. Rohde, E., Tomlinson, A.J., Johnson, D.H., Naylor, S., 1998. Protein analysis by membrane preconcentration-capillary electrophoresis – systematic evaluation of parameters affecting preconcentration and separation. *Electrophor. J. Chromatogr. B: Biomed. Sci. Appl.* 713, 301–311.
56. Rohde, E., Tomlinson, A.J., Johnson, D.H., Naylor, S., 1998. Comparison of protein mixtures in aqueous humor by membrane preconcentration-capillary electrophoresis-mass spectrometry. *Electrophoresis* 19, 2361–70.
57. Johnson, D.H., 1998. Options in the management of malignant glaucoma. *Arch. Ophthalmol.* 116, 799–800.
58. Hattenhauer, M.G., Johnson, D.H., Ing, H.H., Hodge, D.O., Butterfield, L.C., Herman, D.C., Gray, D.T., 1999. Probability of filtration surgery in patients with open-angle glaucoma. *Arch. Ophthalmol.* 117, 1211–1215.
59. Tamm, E.R., Russell, P., Epstein, D.L., Johnson, D.H., Piatigorsky, J., 1999. Modulation of myocilin/TIGR expression in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 40, 2577–2582.
60. Wang, X., Johnson, D.H., 2000. mRNA in situ hybridization of TIGR/MYOC in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 41, 1724–1729.
61. Johnson, D.H., 2000. Myocilin and glaucoma. A TIGR by the tail? *Arch. Ophthalmol.* 118, 974–978.
62. Parc, C.E., Johnson, D.H., Brilakis, H.S., 2000. Giant vacuoles are found preferentially near collector channels. *Investig. Ophthalmol. Vis. Sci.* 41, 2984–2990.
63. Johnson, D.H., Matsumoto, Y., 2000. Schlemm's canal becomes smaller after successful filtration surgery. *Arch. Ophthalmol.* 118, 1251–1256.
64. Fautsch, M.P., Bahler, C.K., Jewison, D.J., Johnson, D.H., 2000. Recombinant TIGR/MYOC increases outflow resistance in the human anterior segment. *Investig. Ophthalmol. Vis. Sci.* 41, 4163–4168.
65. Johnson, D.H., Johnson, M., 2001. How does nonpenetrating glaucoma surgery work? Aqueous outflow resistance and glaucoma surgery. *J. Glaucoma* 10, 55–67.
66. Hann, C.R., Springett, M.J., Johnson, D.H., 2001. Antigen retrieval of basement membrane proteins from archival eye tissues. *J. Histochem. Cytochem.* 49, 475–482.
67. Johnson, D.H., 2001. Glaucoma surgery – a twenty year odyssey. *Ophthalmol-Chirurgie* 13, 84–87.
68. Johnson, D.H., Johnson, M., 2001. Aqueous outflow and glaucoma surgery. How does non-penetrating glaucoma surgery work? *Ophthalmol-Chirurgie* 13, 88–89.
69. Parc, C.E., Johnson, D.H., Oliver, J.E., Hattenhauer, M.G., Hodge, D.O., 2001. The long-term outcome of glaucoma filtration surgery. *Am. J. Ophthalmol.* 132, 27–35.
70. Schoff, E.O., Hattenhauer, M.G., Ing, H.H., Hodge, D.O., Herman, D.C., Johnson, D.H., 2001. Estimated incidence of open angle glaucoma in Olmsted County, Minnesota. *Ophthalmology* 108, 882–886.
71. Brilakis, H.S., Hann, C.R., Johnson, D.H., 2001. A comparison of different embedding media on the ultrastructure of the trabecular meshwork. *Curr. Eye Res.* 22, 235–244.
72. Brilakis, H.S., Johnson, D.H., 2001. Giant vacuole survival time and implications for aqueous humor outflow. *J. Glaucoma* 10, 277–283.
73. Fautsch, M.P., Johnson, D.H., 2001. Characterization of myocilin–myocilin interactions. *Investig. Ophthalmol. Vis. Sci.* 42, 2324–2331.
74. Gottanka, J., Johnson, D.H., Martus, P., Lütjen-Drecoll, E., 2001. Beta-adrenergic blocker therapy and the trabecular meshwork. *Graefes Arch. Clin. Exp. Ophthalmol.* 239, 138–144.
75. Johnson, D.H., Johnson, M., 2001. How does non-penetrating glaucoma surgery work? In: Mermoud, A., Shaarawy, T. (Eds.), *Non-penetrating Glaucoma Surgery*. Martin Dunitz Ltd, The Livery House, London, pp. 33–55.
76. Hann, C.R., Springett, M.J., Wangm, X.F., Johnson, D.H., 2001. Ultrastructural localization of collagen IV, fibronectin, and laminin in the trabecular meshwork of normal and glaucomatous eyes. *Ophthalmic Res.* 33, 314–324.
77. Loewen, N., Fautsch, M.P., Peretz, M., Bahler, C.K., Cameron, J.D., Johnson, D.H., Poeschla, E.M., 2001. Genetic modification of human trabecular meshwork with lentiviral vectors. *Hum. Gene Ther.* 12, 2109–2119.

78. Johnson, D.H., Johnson, M., 2002. Glaucoma surgery and aqueous outflow. How does nonpenetrating glaucoma surgery work? *Arch. Ophthalmol.* 120, 67–70.
79. Oliver, J.E., Hattenhauer, M.G., Herman, D., Hodge, D.O., Kennedy, R., Fang-Yen, M., Johnson, D.H., 2002. Blindness and glaucoma: a comparison of patients progressing to blindness from glaucoma with patients maintaining vision. *Am. J. Ophthalmol.* 133, 764–772.
80. Loewen, N., Bahler, C., Teo, W., Whitman, T., Peretz, M., Xu, R., Fautsch, M.P., Johnson, D.H., Poeschla, E.M., 2002. Preservation of aqueous outflow facility after second-generation FIV vector-mediated expression of marker genes in anterior segments of human eyes. *Investig. Ophthalmol. Vis. Sci.* 43, 3686–3690.
81. Parc, C., Johnson, D.H., 2003. Physiology of aqueous humor outflow resistance: its relation to giant vacuoles. *J. Fr. Ophthalmol.* 26, 198–201.
82. Johnson, D.H., 2003. Progress in glaucoma: early detection, new treatments, less blindness. *Ophthalmology* 110, 634–635.
83. Siddiqui, Y., Ten Hulzen, R.D., Cameron, J.D., Hodge, D.O., Johnson, D.H., 2003. What is the risk of developing pigmentary glaucoma from pigment dispersion syndrome? *Am. J. Ophthalmol.* 135, 794–799.
84. Fautsch, M.P., Silva, A.O., Johnson, D.J., 2003. Carbohydrate binding proteins galectin-1 and galectin-3 in human trabecular meshwork. *Exp. Eye Res.* 77, 11–16.
85. Karger, R.A., Jeng, S.M., Johnson, D.H., Hodge, D.O., Good, M.S., 2003. Estimated incidence of pseudoexfoliation syndrome and pseudoexfoliation glaucoma in Olmsted County, Minnesota. *J. Glaucoma* 12, 193–197.
86. Johnson, D.H., 2003. Progress in glaucoma: early detection, new treatments, less blindness. *Ophthalmology* 110, 1271–1272 (Editorial).
87. Santas, A.J., Bahler, C., Peterson, J.A., Filla, M.S., Kaufman, P.L., Tamm, E.R., Johnson, D.H., Peters, D.M.P., 2003. Effect of heparin II domain of fibronectin on aqueous outflow in cultured anterior segments of human eyes. *Investig. Ophthalmol. Vis. Sci.* 44, 4796–4804.
88. Fautsch, M.P., Vrabel, A.M., Peterson, S.L., Johnson, D.H., 2004. In vitro and in vivo characterization of disulfide bond use in myocilin complex formation. *Mol. Vis.* 10, 417–425.
89. Bahler, C.K., Hann, C.R., Fautsch, M.P., Johnson, D.H., 2004. Pharmacologic disruption of Schlemm's canal cells and outflow facility in anterior segments of human eyes. *Investig. Ophthalmol. Vis. Sci.* 45, 2246–2254.
90. Loewen, N., Fautsch, M.P., Teo, W.-L., Bahler, C.K., Johnson, D.H., Poeschla, E.M., 2004. Long-term, targeted genetic modification of the aqueous humor outflow tract coupled with noninvasive imaging of gene expression in vivo. *Investig. Ophthalmol. Vis. Sci.* 45, 3091–3098.
91. Bahler, C.K., Fautsch, M.P., Hann, C.R., Johnson, D.H., 2004. Factors influencing intraocular pressure in cultured human anterior segments. *Investig. Ophthalmol. Vis. Sci.* 45, 3137–3143.
92. Christiansen, G.A., Nau, C.B., McLaren, J.W., Johnson, D.H., 2004. Mechanism of ocular hypotensive action of bimatoprost (Lumigan) in patients with ocular hypertension or glaucoma. *Ophthalmology* 111, 1658–1662.
93. Bahler, C.K., Smedley, G.T., Zhou, J., Johnson, D.H., 2004. Trabecular bypass stents decrease intraocular pressure in cultured human anterior segments. *Am. J. Ophthalmol.* 138, 988–994.
94. Hann, C.R., Bahler, C.K., Johnson, D.H., 2005. Cationic ferritin and segmental flow through the trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 46, 1–7.
95. Fautsch, M.P., Howell, K.G., Vrabel, A.M., Charlesworth, M.C., Muddiman, D.C., Johnson, D.H., 2005. Primary trabecular meshwork cells incubated in human aqueous humor differ from cells incubated in serum supplements. *Investig. Ophthalmol. Vis. Sci.* 46, 2848–2856.
96. Johnson, D.H., 2005. Trabecular meshwork and uveoscleral outflow models. *J. Glaucoma* 14, 308–310.
97. Gottanka, J., Kuhlmann, A., Scholz, M., Johnson, D.H., Lutjen-Drecoll, E., 2005. Pathophysiologic changes in the optic nerves of eyes with primary open angle and pseudoexfoliation glaucoma. *Investig. Ophthalmol. Vis. Sci.* 46, 4170–4181.
98. Fautsch, M.P., Bahler, C.K., Vrabel, A.M., Howell, K.G., Loewen, N., Teo, W.L., Poeschla, E.M., Johnson, D.H., 2006. Perfusion of His-tagged eukaryotic myocilin increases outflow resistance in human anterior segments in the presence of aqueous humor. *Investig. Ophthalmol. Vis. Sci.* 47, 213–221.
99. Fautsch, M.P., Vrabel, A.M., Johnson, D.H., 2006. Characterization of the *Felix domesticus* (cat) glaucoma-associated protein myocilin. *Exp. Eye Res.* 82, 1037–1045.
100. Fautsch, M.P., Vrabel, A.M., Johnson, D.H. The identification of myocilin-associated proteins in the human trabecular meshwork. *Exp. Eye Res.* 82, 1046–1052.
101. Park, B.-C., Shen, X., Fautsch, M.P., Tibudan, M., Johnson, D.H., 2006. Optimized bacterial expression of myocilin proteins and functional comparison of bacterial and eukaryotic myocilins. *Mol. Vis.* 12, 832–840.
102. Gottanka, J., Johnson, D.H., Grehn, F., Lutjen-Drecoll, E., 2006. Histologic findings in pigment dispersion syndrome and pigmentary glaucoma. *J. Glaucoma* 15, 142–151.
103. Fautsch, M.P., Johnson, D.H., the Second ARVO/Pfizer Institute Working Group, 2006. Aqueous humor outflow: what do we know? Where will it lead us? *Investig. Ophthalmol. Vis. Sci.* 47, 4181–4187.
104. Erie, J.C., Baratz, K.H., Mahr, M.A., Johnson, D.H., 2006. Phacoemulsification in patients with Baerveldt tube shunts. *J. Cataract. Refract. Surg.* 32, 1489–1491.
105. Jeng, S.M., Karger, R.A., Hodge, D.O., Burke, J.P., Johnson, D.H., 2007. Good MS. The risk of glaucoma in pseudoexfoliation syndrome. *J. Glaucoma* 16, 117–121.
106. Johnson, D.H., 2007. Histologic findings after argon laser trabeculoplasty in glaucomatous eyes. *Exp. Eye Res.* 85, 557–562.
107. Khare, P.D., Loewen, N., Teo, W., Barraza, R.A., Saenz, D.T., Johnson, D.H., Poeschla, E.M., 2008. Durable, safe, multi-gene lentiviral vector expression in feline trabecular meshwork. *Mol. Ther. J. Am. Soc. Gene. Ther.* 16, 97–106.
108. Sit, A.J., Nau, C.B., McLaren, J.W., Johnson, D.H., Hodge, D., 2008. Circadian variation of aqueous dynamics in young healthy adults. *Investig. Ophthalmol. Vis. Sci.* 49, 1473–1479.
109. Bahler, C.K., Howell, K.G., Hann, C.R., Fautsch, M.P., Johnson, D.H., 2008. Prostaglandins increase trabecular meshwork outflow facility in cultured human anterior segments. *Am. J. Ophthalmol.* 145, 114–119.
110. Berdahl, J.P., Allingham, R.R., Jonson, D.H., 2008. Cerebrospinal fluid pressure is decreased in primary open-angle glaucoma. *Ophthalmology* 115, 763–768.
111. Lim, K.S., Nau, C.B., O'Byrne, M.M., Hodge, D.O., Toris, C.B., McLaren, J.W., Johnson, D.H., 2008. Mechanism of action of bimatoprost, latanoprost, and travoprost in healthy subjects. A crossover study. *Ophthalmology* 115, 790–795.

Book chapters

1. Johnson, D.H., 1985. Glaucoma: the search goes on. In: *Yearbook of Ophthalmology*, pp. 105–135.
2. Johnson, D.H., 1986. New approaches to an old problem – the medical treatment of glaucoma. In: *Yearbook of Ophthalmology*, pp. 99–124.
3. Johnson, D.H., 1987. Automated perimetry, the blending of art and science. In: *Yearbook of Ophthalmology*, pp. 97–124.

4. Johnson, D.H., 1988. Alpha, beta, and glaucoma. In: Yearbook of Ophthalmology, pp. 77–99.
5. Johnson, D.H., Brubaker, R.F., 1992. Glaucoma. In: Bartley, G.B., Liesegang, T.J. (Eds.), *Essentials of Ophthalmology*. J.B. Lippincott Company, New York, pp. 109–129.
6. Johnson, D.H., 1994. The exfoliation syndrome – a continuing challenge. In: Albert, D.M., Jakobiec, F.A. (Eds.), *Principles and Practice of Ophthalmology*. W.B. Saunders Company, Philadelphia, pp. 1400–1413.
7. Johnson, D.H., 1996. Exfoliation syndrome. In: Albert, D.M., Jakobiec, F.A. (Eds.), *Atlas of Clinical Ophthalmology*. W.B. Saunders Co., Philadelphia, pp. 284–290.
8. Johnson, D.H., 1997. Corticosteroid glaucoma. In: Epstein, D.L., Allingham, R.R., Schuman, J.S. (Eds.), *Chandler and Grant's Glaucoma*, fourth ed. Williams & Wilkins, Baltimore, pp. 404–411.
9. Johnson, D.H., 1999. Pseudoexfoliation syndrome. In: Morrison, J.C., Pollack, I.P. (Eds.), *Essentials of Glaucoma*. Thieme, New York.
10. Johnson, D.H., 2000. The exfoliation syndrome: a continuing challenge. In: Albert, D.M., Jakobiec, F.A. (Eds.), *Principles and Practice of Ophthalmology*, second ed., vol. 4. W.B. Saunders Company, Philadelphia, pp. 2718–2730.
11. Johnson, D.H., Johnson, M., 2001. How does non-penetrating glaucoma surgery work. In: Mermoud, A., Shaarawy, T. (Eds.), *Non-penetrating Glaucoma Surgery*. M. Dunitz Ltd., London, pp. 33–55.
12. Johnson, D.H., 2003. Pseudoexfoliation syndrome and glaucoma. In: Morrison, J.C., Pollack, I.P. (Eds.), *Glaucoma: Science and Practice*. Thieme Medical Publishers, New York, pp. 215–225.
13. Johnson, D.H., Lutjen-Drecoll, E., 2008. Glaucomatous changes in the trabecular meshwork. In: Tombran-Tink, J., Barnstable, C.J., Shields, M.B. (Eds.), *Mechanisms of the Glaucomas: Disease Processes and Therapeutic Modalities*. Ophthalmology Res., Humana Press, Totowa, pp. 99–116.
14. Sit, A.J., Johnson, D.H., 2008. The exfoliation syndrome: a continuing challenge. In: Albert, D.M., Miller, J.W. (Eds.), *Albert & Jakobiec's Principles and Practice of Ophthalmology*, third ed., vol. 2. Saunders Elsevier, Philadelphia, pp. 2581–2593.
5. Johnson, D.H., 1985 Mar. Does pigmentation of the trabecular meshwork affect cellularity? *Investig. Ophthalmol. Vis. Sci.* 26 (Suppl.), 4 (abstract 3).
6. Johnson, D.H., Tschumper, R.C., 1986. An in situ method for culture of human trabecular meshwork. In: *Int. Congr. Eye Res. IV*, p. 184.
7. Johnson, D.H., 1986 Mar. Do trabecular meshwork cells replicate? *Investig. Ophthalmol. Vis. Sci.* 27 (Suppl.), 210 (abstract 2).
8. Johnson, D.H., Tschumper, R.C., 1987 Mar. The effect of organ culture on human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 28 (Suppl.), 283 (abstract 3).
9. Tschumper, R.C., Johnson, D.H., 1987 Mar. A new method for organ culture of human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 28 (Suppl.), 133.
10. Bourne, W.M., Brunette, I., Johnson, D.H., Maguire, L.J., 1989. Long-term corneal perfusion: a research model for donated human eyes unsuitable for transplantation. In: 28th Annual Scientific Session, Eye Bank Association of America, p. 5.
11. Buller, C.R., Johnson, D.H., 1989. Human trabecular meshwork phagocytosis. *Investig. Ophthalmol. Vis. Sci.* 30 (Suppl.), 223.
12. Johnson, D.H., Bradley, J.M., Acott, T.S., Fauss, D.J., Polansky, J.R., 1989. The effect of steroids on human trabecular meshwork in perfusion organ culture. *Investig. Ophthalmol. Vis. Sci.* 30 (Suppl.), 223.
13. Tschumper, R.C., Johnson, D.H., Acott, T.S., Knepper, P.A., 1989. Glycosaminoglycans of human TM in organ culture. *Investig. Ophthalmol. Vis. Sci.* 30 (Suppl.), 224.
14. Johnson, D.H., 1990. Studies on human trabecular meshwork in organ culture. In: *Int. Congr. Eye Res. VI*, p. 81.
15. Tschumper, R.C., Sloan, L.M., Rosenblatt, J.E., Johnson, D.H., 1990. A human in vitro model for acanthamoeba keratitis. *Investig. Ophthalmol. Vis. Sci.* 31 (Suppl.), 419.
16. Buller, C.R., Johnson, D.H., Campbell, R.J., 15 Mar 1991. Do segmental variations exist in the TM of glaucomatous eyes? *Investig. Ophthalmol. Vis. Sci.* 32, 943 (abstract 1347-85).
17. Johnson, D.H., Tschumper, R.C., 15 Mar 1991. Ethacrynic acid: outflow effects and toxicity in human trabecular meshwork in perfusion organ culture. *Investig. Ophthalmol. Vis. Sci.* 32, 1257 (abstract 2885).
18. Bahler, C.K., Johnson, D.H., 1992. The porcine eye as a model for aqueous outflow perfusion studies. *Investig. Ophthalmol. Vis. Sci.* 33 (Suppl.), 1161.
19. Buller, C.R., Johnson, D.H., 1992. Computerized morphometric analysis of JCT: light microscopy vs transmission electron microscopy. *Investig. Ophthalmol. Vis. Sci.* 33 (Suppl.), 1165.
20. Flugel, C., Wiesand, M., Lutjen-Drecoll, E., Johnson, D., 1992. Regional differences in primary open angle glaucomatous eyes. In: Tenth International Congress of Eye Research X, p. 67.
21. Johnson, D.H., Knepper, P., 1992. The effects of ascorbate and dexamethasone on GAGs, IOP, and ultrastructure of human TM in organ culture. *Investig. Ophthalmol. Vis. Sci.* 33 (Suppl.), 1164.
22. Johnson, D.H., Knepper, P.A., 1992. The effects of dexamethasone on GAGs, IOP, and ultrastructure of human TM in organ culture. In: Tenth International Congress of Eye Research X, p. 81.
23. Kimpel, M.W., Johnson, D.H., 1992. Factors influencing in vivo trabecular cell replication in cats. *Investig. Ophthalmol. Vis. Sci.* 33 (Suppl.), 1164.
24. Buller, C.R., Johnson, D.H., 15 Mar 1993. Computerized morphometric analysis of JCT: circumferential differences in normal and glaucomatous eyes. *Investig. Ophthalmol. Vis. Sci.* 34, 1384 (abstract 3362-8).
25. Johnson, D.H., 1994. Cytochalasin D increases outflow facility in the human eye in perfusion organ culture. In: Joint European Research Meetings in Ophthalmology and Vision, p. 51.

Editorials

1. Johnson, D.H., 1998 Jun. Options in the management of malignant glaucoma. *Arch. Ophthalmol.* 116 (6), 799–800.
2. Johnson, D.H., Johnson, M., 2002 Jan. Glaucoma surgery and aqueous outflow – how does nonpenetrating glaucoma surgery work? *Arch. Ophthalmol.* 120 (1), 67–70.
3. Johnson, D.H., 2003. Progress in glaucoma: early detection, new treatments, less blindness. *Ophthalmology* 110, 1271–1272.

Abstracts and letters

1. Johnson, D.H., Bourne, W.M., Campbell, R.J., 1981 Mar. Ultrastructure of Descemet's membrane in Fuchs' dystrophy. *Investig. Ophthalmol. Vis. Sci.* 20, 114 (abstract 1).
2. Allen, R.C., Johnson, D.H., Epstein, D.L., 1982. A long-term clinical trial of betaxolol in glaucoma patients. *Ophthalmology* 89, 191.
3. Johnson, D.H., Richardson, T.M., Epstein, D.L., 1983. Trabecular meshwork phagocytosis: long-term response. *Investig. Ophthalmol. Vis. Sci.* 24 (Suppl.), 136.
4. Johnson, M.C., Johnson, D.H., Kamm, R.D., 1983. Filtration characteristics of the aqueous outflow system. *Investig. Ophthalmol. Vis. Sci.* 24 (Suppl.), 86.

26. Buller, C.R., Camp, J.J., Johnson, D.H., 15 Mar 1994. Three dimensional analysis of trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 35 (4), 1557 (abstract 1417) Annual Meeting Abstract Issue.
27. Johnson, D.H., 15 Mar 1994. Human trabecular cell survival is dependent upon perfusion rate. *Investig. Ophthalmol. Vis. Sci.* 35 (4), 2082 (abstract 3825) Annual Meeting Abstract Issue.
28. Hann, C.R., Camp, J.J., Johnson, D.H., 1995. 3-D analysis of channel architecture in the trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 36 (4), S195.
29. Russell, P., Johnson, D.H., 1995. Two dimensional gel electrophoresis of segments of human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 36, S129.
30. Ten Hulzen, R.D., Johnson, D.H., 1995. Quantitative analysis of the juxtacanalicular tissue and Schlemm's canal using two different fixation pressures. *Investig. Ophthalmol. Vis. Sci.* 36 (4), S730.
31. Johnson, D.H., 1996. Cytochalasin D increases outflow facility in the human eye in perfusion organ culture. *Investig. Ophthalmol. Vis. Sci.* 37 (3), S894.
32. Lutjen-Drecoll, E., Johnson, D., Gottanka, J., Flugel-Koch, C., 1996. Changes in the trabecular meshwork of human eyes after long term treatment with steroids. *Investig. Ophthalmol. Vis. Sci.* 37, S444.
33. Matsumoto, Y., Bahler, C.K., Hann, C.R., Johnson, D.H., 1996. Dexamethasone decreases phagocytosis in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 37 (3), S825.
34. Tamm, E.R., Russell, P., Johnson, D.H., Piatigorsky, J., 1996. Trabecular meshwork accumulates alpha B-crystallin in response to heat shock oxidative stress and culture conditions. *Investig. Ophthalmol. Vis. Sci.* 37, S152.
35. Borrás, T., Masumoto, Y., Johnson, D.H., 15 Mar 1997. Gene transfer into human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 38 (4), S452 (abstract 2122).
36. Gottanka, J., Flugel-Koch, J., Martus, P., Johnson, D.H., Lutjen-Drecoll, E., 15 Mar 1997. Correlation of severity of optic nerve damage with changes in the trabecular meshwork in different forms of glaucoma. *Investig. Ophthalmol. Vis. Sci.* 38 (4), S160 (abstract 794).
37. Hattenhauer, M.G., Johnson, D.H., Herman, D.C., Yawn, B.P., Butterfield, L., Ing, H.H., Hodge, D.O., Gray, D.T., 15 Mar 1997. Probability of blindness from open angle glaucoma. *Investig. Ophthalmol. Vis. Sci.* 38 (4), S729 (abstract 3362).
38. Johnson, D.H., 15 Mar 1997. Schlemm's canal in POAG and PEX at varying stages of severity. *Investig. Ophthalmol. Vis. Sci.* 38 (4), S562 (abstract 2616).
39. Bintareef, R.M., Hattenhauer, M.G., Herman, D.C., Johnson, D.H., Ing, H.H., Hodge, D.O., 1998. Probability of success of trabeculectomy with and without mitomycin C. *Investig. Ophthalmol. Vis. Sci.* 39 (4), S248.
40. Brilakis, H.S., Johnson, D.H., 1998. What is the lifespan of a giant vacuole? *Investig. Ophthalmol. Vis. Sci.* 39 (4), S487.
41. Matsumoto, Y., Mori, K., Kinoshita, S., Johnson, D.H., 1998. Dexamethasone increases the volume and DNA content of the nucleus in human trabecular meshwork cells. *Investig. Ophthalmol. Vis. Sci.* 39, S704.
42. Rohde, E., Tomlinson, A., Naylor, S., Johnson, D.H., 1998. Mass spectrometric analysis of aqueous humor and pseudoexfoliation material. *Investig. Ophthalmol. Vis. Sci.* 39 (4), S490.
43. Welge-Luben, U., May, C.A., Polansky, J.R., Nguyen, T.D., Bloemendal, H., Johnson, D.H., Lutjen-Drecoll, E., 1998 Jul. Stress proteins and TIGR in normal and glaucomatous trabecular meshwork. *Exp. Eye Res.* 67 (Suppl. 1), S134 (abstract 429).
44. Hann, C.R., Johnson, D.H., Springett, M.J., 15 Mar 1999. Immunogold labeling of type IV collagen in the trabecular meshwork of normal and glaucomatous eyes. *Investig. Ophthalmol. Vis. Sci.* 40 (4), S667.
45. Johnson, D.H., Bahler, C.K., 15 Mar 1999. Heparitinase increases outflow facility in the human eye. *Investig. Ophthalmol. Vis. Sci.* 40 (4), S504.
46. Parc, C.E., Johnson, D.H., 15 Mar 1999. Do anatomic factors influence the formation of giant vacuoles? *Investig. Ophthalmol. Vis. Sci.* 40 (4), S505.
47. Wang, X., Bahler, C.K., Johnson, D.H., 15 Mar 1999. mRNA in situ hybridization of TIGR/MYOC in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 40 (4), S667.
48. Hattenhauer, M.G., Johnson, D.H., Herman, D.C., Hodge, D.O., Gray, D.T., 2000. Half empty or half full? *Arch. Ophthalmol.* 118, 861–862.
49. Herndon, L.W., Johnson, D.H., Coleman, A.L., Lee, P.P., 2000. Factors associated with vision loss in glaucoma. *Investig. Ophthalmol. Vis. Sci.* 40, S280.
50. Johnson, D.H., Wang, X., 2000 Feb. Localization of the TIGR-producing cells in human trabecular meshwork by in situ hybridization. *J. Glaucoma* 9 (1), 109 (abstract 16).
51. Fautsch, M.P., Bahler, C.K., Jewison, D.J., Johnson, D.H., 15 Mar 2000. rTIGR/MYOC increases outflow resistance in the human anterior segment. *Investig. Ophthalmol. Vis. Sci.* 41 (4), S762.
52. Hann, C.R., Johnson, D.H., Springett, M.J., 15 Mar 2000. Immunogold labeling of fibronectin and laminin in the trabecular meshwork of normal and glaucomatous eyes. *Investig. Ophthalmol. Vis. Sci.* 41 (4), S504.
53. Johnson, D.H., Parc, C.E., Oliver, J., Hattenhauer, M.G., Hodge, D.O., 15 Mar 2000. The long-term outcome of glaucoma filtration surgery. *Investig. Ophthalmol. Vis. Sci.* 41 (4), S518.
54. Oliver, J.E., Johnson, D.H., Hattenhauer, M.G., Hodge, D.O., Herman, D.C., 15 Mar 2000. Cumulative patient survival rates after diagnosis of open angle glaucoma, and legal blindness secondary to open angle glaucoma in Olmsted County, Minnesota. *Investig. Ophthalmol. Vis. Sci.* 41 (4), S280.
55. Wang, X., Johnson, D.H., 15 Mar 2000. mRNA in situ hybridization of perlecan and laminin gamma1 in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 41 (4), S504.
56. Fautsch, M.P., Bahler, C.K., Jewison, D.J., Johnson, D.H., 16 Oct 2000. MYOC increases outflow resistance in the human anterior segment. In: Fourteenth International Congress of Eye Research, p. S51.
57. Johnson, D.H., Bahler, C., 19 Oct 2000. Glycosaminoglycans, the extracellular matrix, and aqueous outflow. In: Fourteenth International Congress of Eye Research, p. S156.
58. Wheelock, R.H., Johnson, D., Borrás, T., Russell, P., 19 Oct 2000. Microarray comparison of gene expression changes in human trabecular meshwork cells and perfused trabecular meshwork after chronic glucocorticoid treatment. In: Fourteenth International Congress of Eye Research, p. S172.
59. Gottanka, J., Johnson, D.H., Lutjen-Drecoll, E., 2001. Why is intraocular pressure elevated in pigmentary glaucoma? *Investig. Ophthalmol. Vis. Sci.* 42, S145.
60. Peters, D.M., Santas, A.J., Lui, X., Polansky, P.L., Kaufman, E.R., Tamm, D., Johnson, D., 2001. The heparin II domain of fibronectin modulates aqueous outflow: effect on trabecular meshwork matrix and adhesions. *Investig. Ophthalmol. Vis. Sci.* 42, S730.
61. Fautsch, M.P., Johnson, D.H., 15 Mar 2001. Evidence of MYOC–MYOC interactions. *Investig. Ophthalmol. Vis. Sci.* 42 (4), S144.
62. Good, M.S., Krebsbach, V., Loewen, N., Poeschla, E.M., Johnson, D.H., 15 Mar 2001. Transduction of retinal ganglion cells in culture using feline immunodeficiency virus. *Investig. Ophthalmol. Vis. Sci.* 42 (4), S411.
63. Loewen, N., Fautsch, M.P., Xu, R., Cameron, J.D., Johnson, D.H., Poeschla, E.M., 15 Mar 2001. Comparison of lentiviral, type C

- retroviral, and adenoviral vectors for transduction of human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 42 (4), S731.
64. Bahler, C.K., Fautsch, M.P., Hann, C.R., Johnson, D.H., 2002. H-7 increases outflow facility in human cultured anterior segments. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 1031).
 65. Fautsch, M.P., Silva, A.O., Johnson, D.H., 2002. Expression of galectin-1 and galectin-3 in the human trabecular meshwork. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 1057).
 66. Hann, C.R., Bahler, C.K., Fautsch, M.P., Johnson, D.H., 2002. Effect of RGD peptide on outflow facility in human anterior segments in perfusion organ culture. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 1024).
 67. Jeng, S.M., Karger, R.A., Johnson, D.H., Hodge, D.O., Burke, J., Good, M.S., 2002. Conversion rate of pseudoexfoliation syndrome to pseudoexfoliation glaucoma in a population-based study. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 2947).
 68. Johnson, D.H., 2002. Histologic findings after laser trabeculoplasty in glaucomatous eyes. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 3958).
 69. Karger, R.A., Jeng, S.M., Johnson, D.H., Hodge, D.O., Burke, J., Good, M.S., 2002. Incidence of pseudoexfoliation syndrome in Olmsted County, Minnesota. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 3321).
 70. Loewen, N.A., Fautsch, M.P., Bahler, C.K., Johnson, D.H., Poeschla, E.M., 2002. Effects of FIV-based lentiviral vector transduction of trabecular meshwork on aqueous outflow facility and in vivo monitoring of expression in cats. In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 3960).
 71. Siddiqui, Y., Hodge, D.O., Cameron, J.D., Johnson, D.H., 2002. What is the risk of developing pigmentary glaucoma from pigment dispersion syndrome? In: Annual Meeting of the Association for Research in Vision and Ophthalmology 2002 (abstract 285).
 72. Quiram, P.A., Hodge, D.O., Burke, J., Johnson, D.H., Good, M.S., 8 May 2002. Congenital glaucoma: a population-based study and analysis of cases at the Mayo Clinic. *Investig. Ophthalmol. Vis. Sci.* (abstract 3320).
 73. Bahler, C.K., Fautsch, M.P., Acott, T.S., Johnson, D.H., 2003. Factors influencing outflow facility in cultured human anterior segments. In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 1156).
 74. Fautsch, M.P., Vrabel, A.M., Bahler, C.K., Johnson, D.H., 2003. How do protein expression profiles from the human anterior segment culture model compare to the in vivo state? In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 3165).
 75. Karger, R.A., Jeng, S.M., Hodge, D.O., Johnson, D.H., Good, M.S., 2003. Cataract surgery in patients with pseudoexfoliation syndrome in Olmsted County, Minnesota. In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 3422).
 76. Loewen, N.A., Teo, W., Fautsch, M.P., Bahler, C.K., Johnson, D.H., Poeschla, E.M., 2003. Transgene-specific toxicity in lentiviral vector – transduced feline tm in vivo. In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 1147).
 77. Poeschla, E.M., Fautsch, M.P., Loewen, N., Teo, W., Johnson, D.H., 2003. Wild-type and mutant myocilin co-expression causes heterodimer formation and intracellular retention of wild-type protein without impairing proteasome function. In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 1177).
 78. Vrabel, A.M., Fautsch, M.P., Peterson, S.L., Johnson, D.H., 2003. Characterization of disulfide bonds in myocilin complex formation. In: Arvo Annual Meeting Abstract Search & Program Planner 2003 (abstract 1150).
 79. Fautsch, M.P., Vrabel, A.M., Peterson, S.L., Johnson, D.H., 2003. Mar. Characterization of disulfide bond use in myocilin complex formation. *FASEB J.* 17 (4–5) (abstract 629.12).
 80. Bahler, C.K., Johnson, D.H., Smedley, G.T., 2004 Apr. Trabecular bypass stents increase outflow facility in cultured human anterior segments. *Investig. Ophthalmol. Vis. Sci.* 45 (Suppl. 2), U434.
 81. Fautsch, M.P., Vrabel, A.M., Bahler, C.K., Johnson, D.H., 2004 Apr. Aqueous humor supplementation enhances cultured trabecular cell protein expression profiles. *Investig. Ophthalmol. Vis. Sci.* 45 (Suppl. 2), U467.
 82. Hann, C.R., Bahler, C.K., Johnson, D.H., 2004 Apr. Cationic ferritin and segmental flow through the trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* 45 (Suppl. 2), U441.
 83. Johnson, D.H., 27 Apr 2004. Trabecular meshwork and uveoscleral outflow models. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 2235).
 84. Howell, K.G., Vrabel, A.M., Leontovich, A.A., Charlesworth, M.C., Muddiman, D.C., Raghavakaimal, S., Johnson, D.H., Fautsch, M.P., 2 May 2005. A comparison of gene and protein expression in primary human trabecular meshwork cells cultured with human aqueous humor or fetal bovine serum. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1345).
 85. Scholz, M., Gottanka, J., Johnson, D.H., Lutjen-Drecoll, E., 2 May 2005. Pathophysiologic changes in the optic nerves of eyes with primary open angle and pseudoexfoliation glaucoma. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1274).
 86. Fautsch, M.P., Bahler, C.K., Vrabel, A.M., Howell, K.G., Loewen, N., Poeschla, E.M., Johnson, D.H., 3 May 2005. Recombinant myocilin purified from human trabecular meshwork cells increases outflow resistance in human anterior segments but only in the presence of aqueous humor. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 2374).
 87. Hann, C.R., Johnson, D.H., 4 May 2005. Focal adhesions are most prevalent in Schlemm's canal and juxtacanalicular regions of the trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 3699).
 88. Park, B.C., Shen, X., Fautsch, M.P., Tibudan, M., Johnson, D.H., Yue, B.Y.J.T. 4 May 2005. Optimized bacterial expression of myocilin proteins and functional comparison of bacterial and eukaryotic myocilins. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 3707).
 89. Vrabel, A.M., Fautsch, M.P., Charlesworth, M.C., Madden, B., Muddiman, D.C., Johnson, D.H., 4 May 2005. Analysis of myocilin interactions in human trabecular meshwork. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 3690).
 90. Lim, K.S., Nau, C.B., Toris, C.B.A., Hodge, D., McLaren, J.W., Johnson, D.H., 5 May 2005. Mechanism of ocular hypotensive action of bimatoprost (Lumigan), Latanoprost (Xalatan) and Travoprost (Travatan) in normal subjects – a comparison of measurement techniques. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 4725).
 91. Bahler, C.K., Howell, K.G., Hann, C.R., Fautsch, M.P., Johnson, D.H., 1 May 2006. Prostaglandins increase outflow facility in

- cultured human anterior segments. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1845).
92. Fautsch, M.P., Howell, K.G., Leonovich, A.A., Raghavakaimal, S., Johnson, D.H., 1 May 2006. Global gene expression in primary trabecular cell monolayer culture is altered following incubation in human aqueous humor. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1843).
93. Howell, K.G., Vrabel, A.M., Johnson, D.H., Fautsch, M.P., 1 May 2006. Myocilin levels are elevated in the aqueous of patients with POAG. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1847).
94. Khare, P.D., Teo, W., Loewen, N., Barraza, R., Fautsch, M., Johnson, D.H., Poeschla, E.M., 4 May 2006. Long-term genetic modification of trabecular meshwork by lentiviral vectors: an 18 cat study. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 4770).
95. Sit, A.J., Nau, C.B., McLaren, J.W., Johnson, D.H., Hodge, D.O., 7 May 2006. Circadian variation of aqueous dynamics in adults 18–45 years old. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1141).
96. Fautsch, M.P., Howell, K.G., Bahler, C.K., Leontovich, A.A., Raghavakaimal, S., Johnson, D.H., 7 May 2006. Time-course analysis of gene expression in trabecular meshwork following infusion of TGF β in organ culture. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1143).
97. Lei, Y., Garrahan, N., Johnson, D.H., Becker, D., Hernandez, M., Albon, J., Boulton, M., Morgan, J.E., 7 May 2006. Quantification of retina transneuronal changes in human glaucoma using a novel multiphoton-DAPI method. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1555).
98. Howell, K.G., Leontovich, A.A., Raghavakaimal, S., Johnson, D.H., Fautsch, M.P., 7 May 2006. Elevated levels of TGF β cause unique gene expression changes in trabecular meshwork cells incubated in human aqueous humor. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 2060).
99. Hann, C.R., Johnson, D.H., 7 May 2006. Trabecular meshwork pigmentation and collector channels – areas of higher flow? *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 2053).
100. Dzau, J.R., Lee, P.P., Herndon, L.W., Coleman, A.L., Johnson, D.H., 28 April 2008. Perceived causes of loss of vision in glaucoma from patient, family member, and eye care provider perspectives: a focus group study. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1588).
101. Berdahl, J.P., Johnson, D.H., Allingham, R.R., 28 April 2008. Cerebrospinal fluid pressure is lower in primary open angle glaucoma. *Investig. Ophthalmol. Vis. Sci.* Available from: ARVO (<http://www.arvo.org>) (E-abstract 1599).

References

- Grant, W.M., 1950. Tonographic method for measuring the facility and rate of aqueous flow in human eyes. *Arch. Ophthalmol.* 44, 204–214.
- Grant, W.M., 1951. Clinical measurements of aqueous outflow. *Arch. Ophthalmol.* 46, 113–131.
- Johnson, D.H., 2007. Histologic findings after argon laser trabeculoplasty in glaucomatous eyes. *Exp. Eye Res.* 85 (4), 557–562.
- Johnson, D.H., Kenyon, K.R., Epstein, D.L., Van Buskirk, E.M., 1986. Corneal changes during pilocarpine gel therapy. *Am. J. Ophthalmol.* 101 (1), 13–15.
- Johnson, D.H., Tschumper, R.C., 1987. Human trabecular meshwork organ culture. A new method. *Investig. Ophthalmol. Vis. Sci.* 28 (6), 945–953.