Glaucoma Prevalence

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ABSTRACT:

Glaucoma is a group of eye diseases that gradually steals sight without warning and often without symptoms. Vision loss is caused by damage to the optic nerve. This nerve acts like an electric cable with over a million wires and is responsible for carrying the images we see to the brain.

It was once thought that high intraocular pressure (IOP) was the main cause of this optic nerve damage. Although IOP is clearly a risk factor, we now know that other factors must also be involved because even people with "normal" IOP can experience vision loss from glaucoma.

The diagnosis of glaucoma is a gray area that practitioners struggle with. When do we consider the patient to have glaucoma? And who is most at risk to develop glaucoma? These are questions that I would like to have answered. There are many studies out there that provide bits and pieces of glaucoma prevalence, incidence, and risk factors. This research paper will bring many studies together to clarify the prevalence, incidence, and risk factor..

INTRODUCTION:

Glaucoma is a common eye condition in which vision is lost because of damage to the optic nerve.(14)

The optic nerve carries information about vision from the eye to the brain. (14) In most people with glaucoma, optic nerve damage is related to increased pressure of the fluid circulating inside the front portion of the eye. (14) However, glaucoma-related eye damage can occur even when the fluid pressure is normal. (14)

Glaucoma is the second leading cause of blindness in the United States, and the leading cause of blindness in African-Americans.(14) It currently affects as many as 2.5 million Americans, but up to half of people with glaucoma don't know that they have the condition. (14) Glaucoma tends to run in families and is five times more common in African-Americans than in Caucasians.(14) The risk of glaucoma also increases with age in people of all ethnic backgrounds.(14)

Glaucoma, once thought of as a single disease, is actually a broad term for a certain pattern of damage to the optic nerve (the bundle of nerve fibers that carries information from the eye to the brain).(13) This pattern usually occurs in the presence of high intraocular pressure, but contrary to popular belief, glaucoma can occur with normal or even below-normal eye pressure.(13) Worldwide, it is estimated that about 50 million people suffer from vision impairment, if not complete blindness from glaucoma.(13) In the United States, about 300,000 new cases are diagnosed each year, adding to the more than three million cases.(13)

Open-angle glaucoma, the most common form of glaucoma, affects about 3 million Americans--half of whom don't know they have it.(12) It has no symptoms at first.(12) But over the years it can steal your sight.(12) With early treatment, you can often protect your eyes against serious vision loss and blindness.(12)

THEORIES OF GLAUCOMA:

It seems that different mechanisms of damage occur in glaucoma. Schulzer et al. (1990) identified two subgroups of glaucoma patients: one group in which the degree of VF

damage was correlated with IOP level, and one group in which it was not. In glaucoma, ganglion-cell death can be mediated via apoptosis. Stimuli that may lead to apoptotic cell death include neurotrophin deprivation and glutamate toxicity (Nickells 1996). Neurotrophin withdrawal can be caused by blockage of retrograde axonal transport during periods of increased IOP or by defective neurotrophin transport by energy depletion due to ischemia. Glutamate toxicity is believed to be caused by ischemia of the optic nerve and retinal ganglion-cells.

According to the mechanical theory of glaucoma, the main cause of glaucomatous ONH damage is elevated IOP or increased susceptibility to IOP Evidence exist that IOP contributes to the pathogenesis of glaucoma

Blood flow in a tissue is determined by perfusion pressure, arterial pressure minus venous pressure, and resistance to flow between arteries and veins. Like other parts of the central nervous system, the optic nerve and ONH exhibit autoregulation of blood flow constant despite changes in the perfusion pressure, for example in cases of change in arterial pressure or when venous pressure is altered by change in IOP, Ischemia due to increased IOP may result if autoregulation is impaired, for example because of innate deficiency or vasospasm.

RISK FACTOR:

RACE/AGE	RISK FACTOR
Barbados Eye Study: With Cristina Leske PH.D	*7% of blacks have Glaucoma *1% of white minority have Glaucoma *3% mixed population have Glaucoma
	*1% of blacks in 40-50 year of age *10% of blacks in 80's year of age
Beaver Dam Eye study: With Barbara Klein MD Study consisted of 99% white population	*1% 43-54 years of age had Glaucoma *4.5% over 75 years of age have Glaucoma
II.	*1.74% increased odds for every 10 year age increment.
	*1.23% of blacks between 40-49 years of age had Glaucoma
	*11.26% of blacks over 80 years of age had Glaucoma
	*.92% of whites between 40-49 years of age had Glaucoma
	*2.16% of whites over 80 years of age had Glaucoma

Baltimore Eye Study:	*1% of blacks between the ages of 40-50 had Glaucoma *10% over the age of 80 had Glaucoma
Framingham Eye Study:	*4x-5x the risk of blacks to have Glaucoma over the age of 40

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AGE	RISK FACTORS
Beaver Dam Study	*2.1% prevalence in Caucasian population *8.8% Prevalence in St. Lucia population (African-Caribbean)
African-Caribbean Eye Study	*African Caribbean have a 3.7 fold odd on developing Glaucoma over the Roscommon Caucasians

GENDER	RISK FACTORS
Baltimore Eye Study	* No difference between sexes
Beaver Dam Study	* No difference between sexes
Roscommon Study	* No difference between sexes
Barbados Eye Study	*Males are 1.4 times more succeptible to get Glaucoma
Rotterdam Study	*Males are 3 times more succeptible to get Glaucoma
Dalby Study	*Females are more succeptible to get Glaucoma

INTRAOCULAR PRESSURE	RISK FACTOR
Baltimore Eye Study	*Prevalence of POAG increases, with
	increased IOP
	*With decreased IOP, decrease visual
	field loss
	*Normal IOP is between 10-22 mmHg
	*IOP readings fluctuate with different
	corneal thickness.

OPTIC NERVE HEAD	RISK FACTORS

	*7% of the normal population had C/D
Study - Optic disc, cup and neuroretinal	of 0.5 or greater.
rim size, configuration and correlations	*The average horizontal C/D ratio was
in normal eyes.	found to be 0.74 in Caucasians and .057
-	in African-Americans.
	*C/D 's less than 0.74 horizontal and
	0.64 vertical should be considered
	normal.

× (v)

HYPERTENTION	RSK FACTORS
*A number of studies have noted a dire	ct relationship between rise in blood
pressure and a rise in intraocular press	ure.
*younger subjects (< 60 years of age) wi	ith raised blood pressure have a lower riskof
POAG than the age-matched normal po	pulation.
*Older subjects (>70 years of age) high	er risk than their aged-matched population.
Baltimore Eye Study	*Patients who are nocturnal dippers,
	increase the risk of POAG

GENETICS	RISK FACTORS
*Increased risk with family history, 13	-47%
Baltimore Eye Study	*Family history increases risk for POAG
	*Odds are:
	1. Siblings 3.69
	2. Parents 2.17
	3. Children 1.12

OTHER FACTORS	RISK FACTORS
Myopia	*Increase risk of POAG with increase myopia
Diabetes mellitus	*Increase risk of Glaucoma with diabetes mellitus
Smoking	*No associations
Alcohol	*No associations

Conclusion:

Glaucoma of some type is found in about 2% of the population over the age of 40.(11) It can also affect children and young adults, though much less frequently.(11)

It is estimated that over 3 million Americans have glaucoma but only half of those know they have it. (3) Approximately 120,000 are blind from glaucoma, accounting for 9% to 12% of all cases of blindness in the U.S. (3) About 2% of the population ages 40-50 and 8% over 70 have elevated IOP.*(3) Glaucoma is the second leading cause of blindness in the U.S. and the first leading cause of preventable blindness.(3) Glaucoma is the leading cause of blindness among African-Americans. (3) Glaucoma is 6 to 8 times more common in African-Americans than Caucasians. (3) African-Americans ages 45-65 are 14 to 17 times more likely to go blind from glaucoma than Caucasians with glaucoma in the same age group.(3) The most common form, Open Angle Glaucoma, accounts for 19% of all blindness among African-Americans compared to 6% in Caucasians. (3) Other high-risk groups include: people over 60, family members of those already diagnosed, diabetics, and people who are severely near sighted. (3) Estimates put the total number of suspected cases of glaucoma at around 65 million worldwide. (3)

Its very important to take into consideration the corneal thickness, which can determine and explain certain situation where a case does not make sense.

Another factor is central corneal thickness. Normal central corneal thickness varies from one study to another. The studies referenced here find the CCT to average from 524 to 563 um, with standard deviations of from 29 to 38 um.(16) Our experience is that measurements of CCT below 490 um or well above 630 um are frequent in our practices. (16) On average every 20 um account for 1 mmHg of pressure. Prevalence of POAG increases, with increased IOP. With decreased IOP, decrease visual field loss.

In a study 7% of the normal population had C/D of 0.5 or greater, therefore any C/D's over 0.5 should be considered a glaucoma suspect until proven otherwise. African Americans normally have C/D's larger than Caucasians. Consider 0.5 C/D or larger on a Caucasian to have a larger risk than an African American.

Genetics have a role in glaucoma diagnosing. Increased risk with family history of 13-47%, Odds from the Baltimore eye study states that the odds are as follows. Siblings 3.69, parents 2.17, children 1.12. Therefore as a practioner its crucial to include genetic desposition as part of the decision making.

Systemic hypertension has an effect on glaucoma, a number of studies have noted a direct relationship between rise in blood pressure and a rise in intraocular pressure.

Other factors that could effect glaucoma are myopia and diabetes mellitus. Increased risk of POAG with increase myopia. Increase risk of glaucoma with diabetes mellitus. No association was found with smoking and alcohol.

The best way to protect yourself from loss of vision due to glaucoma is with regular, thorough eye exams. You can't treat a disease you don't know you have. Loss of vision from glaucoma is irreversible. Glaucoma usually has no signs or symptoms until serious loss of vision occurs. Most cases of glaucoma are controlled with medication or surgery. Therefore early detection is crucial.

BIBLIOGRAPHY:

- 1. http://www.glaucoma.org/learn/
- 2. Nada J. Lingel, Dennis L. Smith et al. Evaluating the optic nerve head in glaucoma. Pacific college of Optometry 2001 jan.
- 3. Ed Edelson et al. Race, ethnicity revisited in glaucoma symposium (commemorating the Baltimore Eye Survey). Ophthalmology Times 2002 nov; v27 i21 p1.
- 4. James M. Tielsch, Alfred Sommer, Joanne Katz, Richard M. Royall, Harry A. Quigley, Jonathan Javitt et al. The journal of the American Medical Association 1991 july; v266 n3 p369(6).
- 5. LeAnn M. Weih, Bickol N. Mukesh, Catherine A. McCarty and Hugh R. Taylor et al. Association of Demographic, Familial, Medical, and Ocular Factors with Intraocular Pressure. Arch Ophthalmol 2001; 119:875-880
- 6. Michael A. Kass, Dale K. Heuer, Eve J. Higginbotham, Chris A. Johnson, John L. Keltner, J. Philip Miller, Richard K. Parrish 2, M. Roy Wilson and Mae O. Gordon et al. The ocular hypertension treatment study: a randomized trial determines that topical ocular hypotensive medication delays or prevents the onset of primary open-angle glaucoma. Arch Opthalmol 2002; 120:701-713
- 7. Raan S. Ramrattan, Roger C.W. Wolfs, Songhomitra Panda-Jonas, Jost B. Jonas, Douwe Bakker, Huibert A. Pols, Albert hofman, Paulaus T.V.M. de jong et al. Prevalence and causes of visual field loss in the elderly and associations with

- impairment in daily functioning: the Rotterdam study. Archives of Opthalmology 2001 dec; v119 i12 p1788(7)
- 8. Harry A. Quigley, Sheila K. West, Jorge Rodriguez, Beatriz Munoz, Ronald Klein, Robert Snyder et al. The prevalence of glaucoma in a population-based study of hipanic subjects: proyecto VER. Arch Ophthalmol 2001 dec; 119:1819-1826
- 9. Alan P. Rotchford, Gordon J. Johnson et al. Glaucoma in Zulus: a population-based cross-sectional survey in a rural district in South Africa. Arch Opthalmol april 2002; 120:471-478
- 10. M. Kroese, H. Burton, S. Vardy, T. Rimmer and D. McCarter et al. Prevalence of primary open angle glaucoma in general opthalmic practice in the United Kingdom. British Medical Association 2002 sept; v86 i9 p978(3)
- 11. http://193.128.182.48/servlet/dycon/iga/iga/live/en/uk/AboutGlaucoma index
- 12. http://www.nei.nih.gov/health/glaucoma/glaucoma facts.htm
- 13. http://www.ahaf.org/glaucoma/about/glabout.htm
- 14. http://www.intelihealth.com/IH/ihtIH?t=9938&p=
- 15. S. Fraser, R. Worwald et al. Epidemiology of Glaucoma. Opthalmology 1999, 12.1.1-12.1.5
- 16. L. Phillips, C. Cakanac, M. Eger, M. Lilly et al. Central corneal thickness and measured IOP: a clinical study. Journal of the American Optometric Association 2003 april; v74 p219