

Original Research Article

Study of relationship between intraocular pressure and blood pressure in different age groups

Gautam Chand sirvi¹, Jayant Kumar², Dr N D Soni³, Dr Raghuvveer Choudhary⁴

¹Department of Physiology, Dr. S.N. Medical College, Jodhpur, Rajasthan, India

²Professor, ³Head and Sr. Professor, ⁴Associate Professor, Department of Physiology, Dr. S.N. Medical College, Jodhpur, Rajasthan, India

***Corresponding author**

Gautam Chand Sirvi

Email: gautamsirvi@yahoo.com

Abstract: Elevated Intra-ocular Pressure (IOP) is one of the major risk factors for developing glaucoma or glaucomatous optic neuropathy and its progression. Glaucoma is a common ophthalmic disease in India and worldwide and it is a significant cause of visual impairment and blindness. Blindness leads not only to a reduced economical and social status, but it may also result in premature death. Hence, this work was undertaken to study the relationship between IOP and Blood Pressure (BP), so that the prediction of ocular hypertension and its consequences could be forecasted by using more common systemic parameters, i.e., BP. 300 apparently healthy subjects who were aged between 21 - 60 years were examined to find out the relationship between IOP and BP. The BP was recorded with the subjects in the supine position. The IOP was recorded by using Schiotz's tonometer. The IOP was found to increase with age in both men and women and it was statistically significant. The systolic BP and the diastolic BP were positively and significantly correlated with the IOP. As a person's IOP increases along with his/her BP; the subjects with hypertension should be monitored for ocular hypertension. Advancing age was positively associated with the IOP. Hence, in older people, screening needs to be done for high IOP. By symmetry, persons with an elevated IOP suggest that periodic BP monitoring may be indicated for these patients. Hence, a population based screening for elevated IOP and its control could reduce the number of people who are at the greatest risk of glaucoma, which is the second commonest cause for blindness and visual impairment in India and also worldwide.

Keywords: Intra-ocular Pressure, Blood Pressure, Age.

INTRODUCTION

Human aging is characterized by the progressive constriction of the homeostatic reserve of every organ system. This decline, which is often defined as homeostenosis, is evident by the third decade of life and is gradual and progressive [1]. This decreased physiologic reserve in the eye and the cardiovascular system can be manifested by the changes in the intraocular pressure (IOP) and the blood pressure (BP) with increasing age respectively. The intra-ocular pressure (IOP) is maintained by an equilibrium between the aqueous production from the ciliary body and its drainage via the trabecular complex. The mean IOP varies between 10 and 21 mm Hg (mean 16 ± 2.5) [2]. Any abnormalities in the IOP may result in the dysfunction of the eye, which in turn may affect the vision. The IOP is affected by various systemic parameters like age, gender and BP. The IOP tends to increase with the age of a person and a change in BP is directly associated with a change in the IOP [3, 4]. A medical intervention can bring the BP back to normal, hence preventing the consequences of hypertension. An elevated IOP is one of the major risk factors for

developing glaucoma or glaucomatous neuropathy and its progression [5, 6]. Glaucoma is a common ophthalmic disease worldwide and a significant cause of visual impairment and blindness. It is the second leading cause of blindness which is responsible for 23% of all the blindness cases [6]. Blindness leads not only to a reduced economical and social status, but it may also result in premature death [7]. The World Health Organization has estimated that India has a 1% prevalence of blindness. Of the estimated 8.9 million blind persons in India, 12.8% of the blind cases are caused by glaucoma. The problem is expected to reach alarming proportions by the turn of the century [8]. The World Population Projection for 2010 and 2020 derived that by 2020, India would become second in having the maximum number of glaucoma cases, surpassing Europe. From 2010 to 2020, the most detectable change in glaucoma worldwide will be its increase in India. As the proportion of those above 40 years of age increases, the proportional increase in glaucoma will challenge our resources and ingenuity [9]. If ocular hypertension or glaucoma are detected early and treated appropriately, their progression and blindness can be

prevented. Yet, in the physiologic arena, the detailed variation of the IOP and the BP with reference to different age groups affecting the visual functioning is still not available completely. Hence, this work was undertaken to study the relationship between the IOP and the BP in different age groups, so that the prediction of ocular hypertension and its consequences could be forecasted by using more common systemic parameters, i.e., BP and age.

MATERIAL & METHODS

The present study was conducted in M.D.M. Hospital associated with Dr. S. N. Medical College and associated group of hospitals, Jodhpur. 300 apparently healthy subjects aged between 21 to 60 years attending Ophthalmology OPD of M.D.M. Hospital, Jodhpur were examined. All these subjects were subdivided in four different age groups of 21-30, 31-40, 41-50 & 51-60 years. Subjects of all four groups were subjected to anthropometric measurements like height and weight. SBP & DBP of all subjects was recorded in supine position as per standard protocol. IOP was also recorded by using Schiottz's tonometer. Data thus obtained were subjected to statistical analysis for any association between IOP, age, BP and anthropometric parameters.

Inclusion criteria:

1. Subjects above 18 years.
2. Subjects who are newly diagnosed hypertensive and not on antihypertensive medication.

Exclusion criteria:

3. Subjects below 18 years.
4. Subjects who were previously diagnosed as hypertensive and were taking medications.
5. Subjects who were having closed angle glaucoma and other causes of glaucoma.
6. Subjects who are blind.
7. Subjects with any ocular surgery.
8. Subjects diagnosed as diabetic.
9. Subjects with any medications.

Selected subjects gave an informed consent after detailed procedure of the non-invasive technique was explained to them. Detailed history about name, age, occupation, personal and past history, habits of the subjects were taken. Family history of hypertension, diabetes, refractive errors and glaucoma were enquired. Vital parameter like heart rate was recorded by placing the diaphragm of the stethoscope on the mitral area of the precordium for a whole one minute. The B.P. was then recorded with a mercury sphygmomanometer, in the right upper limb by both palpatory and auscultatory method. These were recorded in resting and supine position. The pulse pressure ($PP = SBP - DBP$) and the

mean arterial pressure ($MAP = DBP + 1/3 PP$) were also calculated. A detailed clinical examination of respiratory system, cardiovascular system and central nervous system was done.

Procedure of recording the IOP:

The IOP was recorded by using Schiottz indentation tonometer. The instrument was calibrated before each use by placing it on a polished metal sphere and checking to be sure that the scale reading is 'zero'. If the reading was not zero it was readjusted to zero. Subjects were explained the nature of test and reassured that the procedure was harmless. The patient was laid in the supine position and asked to look straight upward on an over head target or a mark on the ceiling with fixed gaze. They were told to relax & breath normally and keep eyes wide open without blinking during the procedure. Cornea was anesthetized with 2 – 3 drops of 4% topical Lignocaine. The tonometer tip and footplate were wiped carefully with an alcohol swab and allowed to air dry. Subject's eye lids were retracted gently with left hand without placing tension on the globe. The footplate of the tonometer was placed directly over the cornea by holding the handle of the tonometer with right hand. The handle of the tonometer was lowered to a position midway between the top and the footplate of the cylinder; thereby the instrument will act independently by its own weight. The reading on the scale was recorded as soon as the needle became steady. The scale of the Schiottz Indentation tonometer is calibrated in such a fashion that each scale unit represents 0.05 mm protrusion of the plunger. The recording of the IOP was started with 5.5 g weight however if the scale reading was less than three, additional weight was added to the plunger to make it 7.5 g or 10 g as indicated. The IOP measurement was repeated until three consecutive readings agreed within 0.5 scale units. The average scale reading and the plunger weight were then converted into IOP in mmHg by using a conversion chart, Frieden Wald Nomogram. After each use, the tonometer plunger and footplate were rinsed with water followed by alcohol, and then wiped dry with lint – free material. After the procedure, a prophylactic antibiotic, Ciprofloxacin eye drops were instilled in both the eyes to prevent infections.

RESULTS

The ages of the subjects ranged between 21 to 60 years. The distributions of the subjects in the different age groups are shown in [Table-1]. The mean IOP in men and women in the different age groups is shown in [Table-2]. The IOP increased with the age in both men and women. After controlling the systolic BP and the diastolic BP, the age was found to be positively related to the IOP. As the age increased, the IOP of the subject also increased [Table-3],[Table-4].

Table 1: AGE AND SEX WISE DISTRIBUTION

Age groups (in yrs)	Sex		Total	%
	Male	Female		
21-30	31	23	54	18.00
31-40	39	35	74	24.66
41-50	42	35	77	25.67
51-60	48	47	95	31.67
Total	160	140	300	100

Table 2: VARIATION OF IOP WITH SEX AND AGE

Sex	Age (in yrs)	No. of Subjects	Intraocular Pressure (mmHg)	
			Mean	S.D.
Male	21-30	31	14.83	± 2.31
	31-40	39	15.67	± 2.03
	41-50	42	15.86	± 2.24
	51-60	48	16.35	± 2.01
Female	21-30	23	14.47	± 1.34
	31-40	35	15.45	± 2.2
	41-50	35	15.6	± 1.9
	51-60	47	16.53	± 2.27

Table 3: MEAN IOP FOR SPECIFIED LEVELS OF SYSTOLIC BP AND AGE

Age group (in yrs)	No. of Subjects	Systolic BP (mmHg)	Intraocular Pressure (mmHg)	
			Mean	S.D.
21-30	16	<120	13.81	± 1.8
	19	120-139	14.47	± 1.58
	19	140-160	15.64	± 2.09
31-40	22	<120	14.31	± 2.12
	20	120-139	15.3	± 1.75
	32	140-160	16.6	± 1.79
41-50	16	<120	14.25	± 1.39
	28	120-139	16.25	± 1.88
	33	140-160	16.03	± 2.23
51-60	16	<120	15.25	± 2.43
	36	120-139	16.53	± 1.95
	43	140-160	16.82	± 2.05

Table 4: MEAN IOP FOR SPECIFIED LEVELS OF DIASTOLIC BP AND AGE

Age group (in yrs)	No. of Subjects	Diastolic BP (mmHg)	Intraocular Pressure (mmHg)	
			Mean	S.D.
21-30	16	<80	13.82	± 1.8
	22	80-90	14.55	± 1.51
	16	91-100	15.75	± 2.23
31-40	22	<80	14.31	± 2.12
	23	80-90	15.74	± 2.11
	29	91-100	16.37	± 1.64
41-50	16	<80	14.25	± 1.39
	33	80-90	16.12	± 1.82
	28	91-100	16.14	± 2.35
51-60	16	<80	15.25	± 2.43
	41	80-90	16.71	± 1.97
	38	91-100	16.66	± 2.05

The relationship between the age, systolic BP, diastolic BP, the mean arterial pressure and the IOP and this relationship were statistically significant ($P < 0.05$)

Statistical analysis:

Results were presented as Mean \pm SD and range values. Student's t test was used for comparing means of two groups. P – Value of 0.05 or less was considered for statistical significance.

DISCUSSION

Glaucoma is the commonest cause of irreversible blindness worldwide and the second most common cause of blindness overall, after cataract. It affects approximately 70 million people and among them, 7 million are blind [10]. Glaucoma or glaucomatous optic neuropathy is characterized by a chronic, slowly progressive loss of the retinal ganglion cells and their neurons. An elevated IOP is one of the major risk factors for developing glaucomatous optic neuropathy [11]. Open angle glaucoma is a leading cause of visual impairment and blindness [12,13]. Ocular hypertension is a predisposing factor for open angle glaucoma or glaucomatous optic neuropathy. The knowledge of their cause, their natural history and the risk factors of these disorders is incomplete. The IOP is widely regarded as the most important modifiable risk factor which is associated with the development of glaucomatous optic neuropathy [14-18]. Therefore, the factors that influence the IOP and its measurement are of great relevance in understanding the pathogenesis of the disease and in reducing the burden of blindness. The correlates of this measurement include other physiologic parameters that may be needed to be considered in investigating the determinants of the IOP. We briefly investigated and described the relationship between age, BP, gender and IOP. In the present study, the IOP correlated positively with age in both men and women. The increase in the IOP with age was statistically significant. Old age has been reported as a risk factor for the development of glaucoma in patients with ocular hypertension, in multiple progression studies [16,18]. Several population based studies have found that the incidence of open angle glaucoma increased with the older age groups [18] In a study, there was a strong evidence that old age was an independent risk factor for the progression of ocular hypertension and glaucoma [18]. In elderly individuals, the onset of the structural changes in the trabecular meshwork results in a reduction in the trabecular outflow facility and the uveoscleral outflow and hence an elevated IOP in the older age group [19-21]. Aging is associated with a modest elevation of the IOP and it is also linked to a progressive decline in the cerebral and the ocular perfusion [22]. Older patients with glaucoma may have dysfunction of the ocular blood flow auto regulation [22]. Other possible risk factors for the development of ocular hypertension or glaucoma in old age are : local vasospasm, sleep apnoea,

abnormalities of the connective tissue of the lamina cribrosa, primary ganglion cell degeneration, systemic hypertension and atherosclerosis [22, 23]. An increased age may reflect the cumulative effects of some other factors that cause the aging optic nerve head to be more vulnerable to the elevated IOP and even sometimes to the normal range of the IOP [15]. The findings from our study indicated that systolic BP, diastolic BP, PP and MAP were positively independently correlated to the IOP and that the correlations were statistically significant. Some studies have found that a change in the IOP was directly and significantly associated with a change in the BP [14, 15, 24-28]. A positive association between the systolic BP and a raised IOP has constantly been shown in both cross sectional and longitudinal studies [24, 29-32]. Some studies have shown that the diastolic BP was positively associated with a raised IOP [4, 33, 34]. The IOP may have been increased in patients with an increased BP due to an increased retinal blood volume after a rise in the central retinal vein pressure because of an increased pressure in the adjacent central retinal artery [34]; an increased blood volume in the ciliary body and a decreased facility of the aqueous outflow, owing to an increase in the resistance in the episcleral and the anterior ciliary veins [34]; an increased ultra filtration of the aqueous fluid in the ciliary body, owing to the increased perfusion pressure in the ciliary arteries; [4,19,33-36] obstruction to the aqueous drainage at the anterior chamber angle due to the increasing episcleral venous pressure [3,4,34]. The IOP rises and falls by 1 mmHg with every heart beat; during systole, the central retinal artery compress the accompanying vein to increase the vascular resistance in this vessel [34]. Follow-up studies which were done for five years have shown that the cumulative probability of untreated patients developing glaucoma was calculated to be greater than twice the rate of that in the treated patients [16]. The findings from this study indicate that the IOP increased with age in both men and women and that it was statistically significant. The systolic BP and the diastolic BP were positively and significantly correlated with the IOP.

CONCLUSION

It can be concluded that persons with hypertension and advancing age need to be monitored for high IOP. By symmetry, in persons with an elevated IOP, periodic BP monitoring may be indicated. Hence, a population based screening for an elevated IOP and its control could reduce the number of people who are at the greatest risk of glaucoma, which is the second commonest cause for blindness and visual impairment in India and also worldwide.

REFERENCES

1. Kasper DL, Braundwald E, Fauci AS, Hauser SL, Longo DL, Jame son JL; Harrison's Principles of Internal Medicine. 16th ed. United States of

- America: The McGraw Hill Companies; 2005; 44(2).
2. Khurana AK, Khurana I; Anatomy and Physiology of the Eye. 2nd ed. New Delhi: CBS Publishers and Distributors; 2006; 44-81.
 3. Kaufman PL, Alm A; Alder's Physiology of the Eye: clinical application. 10th ed. Missouri: Mosby (Inc); 2003; 235-71.
 4. Klein BEK, Klein R, Knudtson MD; Intra-ocular pressure and systemic blood pressure: a longitudinal perspective: the Beaver Dam eye study. *Br J Ophthalmol* 2005; 89: 284-7.
 5. Pache M, Flammer J; A sick eye in a sick body? Systemic findings in patients with primary open-angle glaucoma. *Surv Ophthalmol* 2006; 51(3): 179-212.
 6. Mansberger SL, Demirel S; Early detection of glaucomatous visual field loss: why, what, where and how? *Ophthalmol Clin N Am* 2005; 18: 365-73.
 7. Park K; Park's Textbook of Preventive and Social Medicine. 18th ed. Jabalpur: Banarsidas Bhanot Publishers; 2005; 319-23.
 8. Jacob A, Thomas R, Koshi SP, Braganza A, Muliylil J; Prevalence of primary glaucoma in an urban south – Indian population. *Indian J Ophthalmol* 1998; 46: 81-6.
 9. Quigley HA, Broman AT; The number of people with glaucoma worldwide in 2010 and 2020. *Br J Ophthalmol* 2006; 90: 262-7.
 10. Yip JLY, Aung T, Wong TY, Machin D, Khaw PT, Khaw KT, et al. The socio-economic status, the systolic blood pressure and the intraocular pressure: the Tanjong Pagar study. *Br J Ophthalmol* 2007; 91: 56-61.
 11. Shiose Y, Kawase Y. A new approach to the stratified normal intraocular pressure in a general population. *Am J Ophthalmol* 1986 Jun; 101: 714-21.
 12. Leske MC, Connell AMS, Schachat AP, Hyman L; The Barbados eye study: prevalence of open angle glaucoma. *Arch Ophthalmol* 1994; 112: 821-9.
 13. Leske MC; The epidemiology of open – angle glaucoma: a review. *Am J Epidemiol* 1983 Aug; 118 (2): 166-91.
 14. Leske MC, Connell AMS, Wu SY, Nemesure B, Li X, Schachar A; Incidence of open-angle glaucoma. *Arch Ophthalmol* 2001; 119: 89-95.
 15. Le A, Mukesh BN, McCarty CA, Taylor HR; Risk factors which are associated with the incidence of open – angle glaucoma: the visual impairment project. *Invest Ophthalmol Vis Sci* 2003; 44 (9): 3783-9.
 16. Gordon MO, Beiser JA, Brandt JD, Heuer DK, Higginbotham EJ, Johnson CA, *et al.*; The ocular hypertension treatment study. *Arch Ophthalmol* 2002; 120: 714-20.
 17. Kass MA, Heuer DK, Higginbotham EJ, Johnson CA, Keltner JL, Miller JP, *et al.*; The ocular hypertension treatment study. *Arch Ophthalmol* 2002; 120: 701-13.
 18. Friedman BS, Wilson MR, Liebman JM, Fechtner RD, Weinreb RN; An evidence – based assessment of the risk factors for the progression of ocular hypertension and glaucoma. *Am J Ophthalmol* 2004; 138 suppl: S19-31.
 19. Nemesure B, Hennis A, Wu SY, Leske MC; Factors which are related to the 4 – year risk of a high intraocular pressure. *Arch Ophthalmol* 2003; 121: 856-62.
 20. Carol TB, Scott KA, Michael YE, Carl CB; Aqueous humor dynamics in ocular hypertensive patients. *J Glaucoma* 2002; 11 (3): 253-8.
 21. Gaasterland D, Kupfer C, Milton R, Ross K, McCain L, MacLellan H; Studies on the aqueous humor dynamics in man VI. Effect of age on the parameter of intraocular pressure in normal human eyes. *Exp Eye Res* 1978; 26 (6): 651-6.
 22. Harris A, Rechtman E, Siesky B, Cuypers CJ, McCranorl, Garzozzi HJ; The role of the optic nerve blood flow in the pathogenesis of glaucoma. *Ophthalmol Clin N Am* 2005; 18: 345-53.
 23. Tielsch JM, Katz J, Sommer A, Quigley HA, Javitt JC; Hypertension, perfusion pressure and primary open – angle glaucoma. *Arch Ophthalmol* 1995; 113: 216-21.
 24. Klein BB, Klein R; The intra-ocular pressure and the cardiovascular risk variables. *Arch Ophthalmol* 1981; 99: 837-9.
 25. Dielemans I, Vingerling JR, Algra D, Hofman A, Grobbee DE; Primary open -angle glaucoma, the intraocular pressure and the systemic blood pressure in the general elderly population. *Ophthalmology* 1995; 102 (1): 54-60.
 26. Bonomi L, Marchini G, Marraffa M, Bernardi P, Morbio R, Varotto A; Vascular risk factors for primary open angle glaucoma. *Ophthalmology* 2000; 107 (7): 1287-93.
 27. Hennis A, Wu SY, Nemesure B, Leske MC; Hypertension, diabetes and longitudinal changes in the intra-ocular pressure. *Ophthalmology* 2003; 110 (5): 908-14.
 28. Lee JS, Kim SM, Choi HY, Oum BS; A relationship between the intraocular pressure, age and the body mass index in a Korean population. *J Korean Ophthalmol Soc* 2003; 44 (7): 1559-66.
 29. McLeod SD, West SK, Quigley HA, Fozard JL; A longitudinal study of the relationship between the intraocular and the blood pressures. *Invest Ophthalmol Vis Sci* 1990; 31: 2361-6.
 30. Nomura H, Shimokata H, Ando F, Miyake Y, Kuzuya F; Age related changes in the intra-ocular pressure in a large Japanese population. *Ophthalmology* 1999; 106 (10): 2016-22.33.
 31. Rohtchina E, Mitchell P, Wang JJ; Relationship between age and intra-ocular pressure: the Blue Mountains eye study. *Clin Exp Ophthalmol* 2002; 30 (3): 173-5.

32. Foster PJ, Machin D, Wong TY, Ng TP, Kirwan JF, Johnson GJ *et al.*; Determinants of the intraocular pressure and its association with glaucomatous optic neuropathy in Chinese Singaporeans: the Tanjong Pagar study. *Invest Ophthalmol Vis Sci* 2003; 44: 3885-91.
33. Hiller R, Sperduto RD, Krueger DE. Race; iris pigmentation and intraocular pressure. *Am J Epidemiol* 1982; 115 (5): 674-83.
34. Bulpitt CJ, Hodes C, Everitt MG; The intra-ocular pressure and the systemic blood pressure in the elderly. *Br J Ophthalmol* 1975; 59: 717-20.
35. Klein BEK, Klein R, Linton KLP; The intra-ocular pressure in an American community. *Invest Ophthalmol Vis Sci* 1992; 33: 2224-8.
36. Wu SY, Nemesure B, Hennis A, Leske MC; Nine – year changes in the intra-ocular pressure. *Arch Ophthalmol* 2006; 124: 1631-6.