We are IntechOpen, the world's leading publisher of Open Access books Built by scientists, for scientists

6,900

186,000

200M

Download

154
Countries delivered to

Our authors are among the

TOP 1%

most cited scientists

12.2%

Contributors from top 500 universities



WEB OF SCIENCE

Selection of our books indexed in the Book Citation Index in Web of Science™ Core Collection (BKCI)

Interested in publishing with us? Contact book.department@intechopen.com

Numbers displayed above are based on latest data collected.

For more information visit www.intechopen.com



Incidence of Refractive Error and Amblyopia Among Young Adults – A Hospital Based Study

Ashok Kumar Narsani¹, Shafi Muhammad Jatoi¹, Mohan Perkash Maheshwari² and Khairuddin Shah¹ ¹Department of Ophthalmology, Liaquat University Eye Hospital, Hyderabad Sindh, ²Department of Pharmacology, Baqai Medical University, Karachi Sindh, Pakistan

1. Introduction

Visual impairment remains a major public health problem world wide, with an estimated 161 million people with visual impairment, of whom 37 million are blind. Uncorrected or inadequately corrected refractive errors have been shown to be a major cause of visual impairment in population-based studies²⁻⁶. While amblyopia is a significant cause of unilateral reduced visual acuity in a population aged 40 years and older.⁷

Genetic factors are thought to play a role in development of refractive errors. It has been established that myopia clusters within families, and familial high myopia (refraction of -6 diopter {D} or less) has been linked to long arm regions on chromosomes 7, 12 and 188,9. Environmental risk factors have also been associated with refractive errors, myopia or hypermetropia. Education^{10,11} and near-work¹² are both strongly associated with increasing severity of myopia.

In different parts of Asia such as in India, the Andhra Pradesh Eye Disease Study shows 15.2%¹³ prevalence rate of myopia (spherical equivalent {SE} at least -1.0D). While study in a 15,068 Singapore military recruits aged 16 to 25 years, the prevalence rates of myopia (SE at least -0.5 D) were much higher with some racial variation, 82.2% in Chinese, 68.8% in Indians, and 65.0% in Malays¹⁴. Similar high rates of myopia (SE at least -0.25 D; 84%) were present in 16 to 18 years old Chinese children in Taiwan¹⁵. In Pakistan the prevalence rates of myopia, hypermetropia, astigmatism (with SE worse than -0.5 D, greater than +0.5D, and greater than 0.75D respectively) was 36.5%, 27.1%, and 37%, respectively in adults aged 30 years or more in the National Blindness and Visual Impairment Survey.¹⁶

In United states, the Baltimore eye survey¹⁷ and Beaver Dam study¹⁸ showed the prevalence rates of myopia (SE at least -0.5 D) were 28.1% in white adults aged 40 years or more and 26.2% in adults aged 43 to 84 years respectively.

Amblyopia is a frequent cause of unilateral or bilateral blindness. The prevalence of amblyopia ranges from 0.73% to 3.06%⁷ in previous studies. However, the epidemiology of amblyopia among this region of Asia is not well described and may be different from other because of difference in distribution of refractive error or strabismus.

AIM: To assess the incidence of refractive error and associated amblyopia among young adult.

2. Methodology

2.1 Subjects

This six months study was conducted from June 2008 to November 2008 at tertiary referral center Liaquat University Eye Hospital, Hyderabad Sindh, Pakistan. Three thousand four hundred fifty two patients were included and examined in out patients department with age ranged from 20-40 years. The proportion of men and women was 1:0.54. Both rural and urban residents were evaluated. After taking written consent all subjects underwent a comprehensive ophthalmic examination, and a standardized form was used to extract the data, including the following variables: demographic information, best corrected visual acuity, types of refractive error including myopia, hypermetropia, astigmatism and amblyopia.

2.2 Methods

Monocular visual acuity was determined with current spectacle prescription if any. Pinhole acuity was assessed in eyes with presenting visual acuity <20/20. Non-cycloplegic autorefraction followed by subjective refinements was performed in all subjects. The best corrected visual acuity was recorded. Refraction data was based on subjective refraction. Only the right phakic eye of each subject was considered for refractive error evaluation and amblyopia evaluated bilaterally.

Amblyopia was defined as best-corrected visual acuity of 20/40 or worse in the absence of any pathological cause. Hypermetropia was defined as a SE greater than +0.5 diopter sphere (DS)¹⁷⁻²¹. Emetropia was defined as a SE between -0.50 and +0.50 DS²⁰. Myopia was defined as a SE worse than -0.50 DS¹⁷⁻²¹ and a SE or worse than -5.00 DS¹⁹ was classified as high myopia. Astigmatism correction was prescribed in minus cylinder format, and astigmatism was defined as cylindrical error worse than -0.50 diopter cylinder (DC) in any axis. The axis of any cylinder component was classified as with the rule (WTR) if the minus cylinder axis was within 15° of 180°, against the rule (ATR) for minus cylinder axis within 15° of 90°, or oblique (other than WTR or ATR)²².

3. Results

Of the 3452 patients, 847 (24.54 %) patients had best corrected visual acuity 20/40 or better and remaining 2605 (75.46%) had less than 20/40 due to different anterior and posterior segment eye pathologies, or amblyopia.

Out of the 847 patients 525 (15.20 %) were phakic in right eye, and remaining 322 (9.32%) were pseudophakic. For refractive error the result was analyzed for only 15.20%

phakic ametropic patients. While for amblyopia all patients who met the criteria were evaluated.

There were 341 (64.95% of phakic ametropic patients) male and 184 (35.05%) were female. The age range from 20 to 40 years **(Table 1)**, mean age being 26±3.9 years. The mean age of men and women was 28±3.9 and 24±6.3 years respectively (statistically significant at P<0.001). The mean refractive error was 0.75D.

Hypermetropia was found in 185 (35.24% of phakic ametropic) patients (Table 1). The mean age of hypermetropia was 26.31±4.51 years. Which was not significantly different from that of entire population (P=0.5476). There were 121 men (23.04% of total ametropic) and 64 women (12.19%). Man had significantly higher prevalence of hypermetropia than women (P<0.001).

Three hundred and fifteen (60% of phakic ametropic) patients had myopia (Table 1). The mean age of myopes was 23.69±3.98 years. Which was significantly lower than the entire population (P<0.001). There were 205 (39.05%) men and 110 (20.95%) women. The man had a significantly higher prevalence of myopia than did the women (P<0.001)

Twenty five (4.76%) patients of the study population were high myopes (Table 1). Among which 15 patients were males and 10 were females. The mean age among high myopes was 22. 50±3.25 years. Which was also significantly lower than the entire population (P<0.0002). However there was no significant different between the mean age of myopes and high myopes (P=0.1632).

Two hundred and fifteen (25.38 % of phakic ametropic) patients had astigmatism worse than -0.5 DC. The males were 134 (62.32%) of total astigmatic patients and remaining were females. The man had a significantly higher prevalence of astigmatism than women (P<0.001). One hundred and forty two (66.04%) patients had ATR astigmatism, 52 (24.18%) had WTR astigmatism and 21 (9.76%) had oblique astigmatism. The prevalence of against the rule astigmatism increased significantly with age (P=0.007).

The incidence of associated unilateral amblyopia was in 19 (3.62%) of phakic ametropic patients (Table 2). Ten (52.63%) patients were male and 09 (47.37%) were females. Amblyopia was not found to be significantly different by age (p=0.1312) group and gender (p=0.1211). Anisometropia was more common in amblyopic cases (41.75%) compared to the normal population (5.91%), and 69% of amblyopic eyes had visual acuity worse than 20/60. However, two amblyopic patients had strabismus in addition to anisometropia, but the prime reason of both conditions was anisometropia. While none of the case of bilateral amblyopia were seen in this study.

Age (yrs)	Hypermetropia		Myopia		High myopia		Total (%)
	M	F	M	F	M	F	Total (%)
21-25	38	21	48	25	7	4	143 (27.23)
26-30	29	17	65	27	5	4	147 (28.00)
31-35	27	10	49	32	0	1	119 (22.68)
36-40	27	16	43	26	3	1	116 (22.09)
Total	121	64	205	110	15	10	525 (100)

Table 1. Age and type of refractive error

A 22 (2742)	Frequency	Total (0/)	
Age (yrs)	Male	Female	Total (%)
21 -25	2	2	4 (21.04)
26 -30	3	2	5 (26.32)
31 -35	2	3	5 (26.32)
36 -40	3	2	5 (26.32)
Total	10	9	19 (100)

Table 2. Demography of amblyopia

4. Discussion

The incidence of hypermetropia in this study was 35.24% of total 525 phakic ametropic patients. In contrast to Andhra Paeye disease study (APEDS)¹³, Barbados eye study²⁰, and several other studies^{17,18} hypermetropia decrease with increasing age in our study

The incidence of myopia and high myopia in this study was 60% and 4.76% of the phakic ametropic patients and decreased with age. The Attebok et al¹⁸, Wang et al¹¹, Katz et al¹⁷ reported a significant trends of decreasing myopia with age. However the Chennai glaucoma study³, Barbodos²⁰ study reported increase of myopia with age and also have an association of nuclear sclerosis with myopia. Saw²², Guggenheim et al²³ and Dan et al²⁴ reported environmental influence such as near work, night lighting and ultraviolet exposure may be responsible for ageing of the crystalline lens and associated myopia. In contrast to population based studies from Australia¹⁸, Singapore¹⁹, and Indonesia²⁵, the incidence of myopia was more in males than females in our study.

The incidence of astigmatism in this study was 25.38% and increased significantly with age. The same has been reported from Chenni³, Australia¹8, Singapore¹9, and Indonesia²5. ATR astigmatism in this study was predominant that made the 66.04% of the total ametropic patients. In relation to the Chennai³ study the incidence of ATR astigmatism significantly increased with age and WTR astigmatism decreased with age in our study. Gudmundsdottir et al²6, Pensyl et al²² and Goss et al²8 reported same and the reason for this could be increased lid laxity with age causing flattening of vertical corneal meridian thereby decreasing WTR astigmatism and increased ATR astigmatism.

In this study the incidence, of associated amblyopia was 3.62% of phakic ametropic patients, which was less than Goel B.S²⁹ study in which amblyopia was reported 5.97%. One thing common in both were higher rate of amblyopia in ametropic then general population. In contrast to Karki JKD³⁰ study amblyopia was not found significantly different by age and gender in this study.

In conclusion, 15.20% of people had refractive error and 3.62% has the amblyopia. The prevalence of myopia was 60% and decreased with age. Hypermetropia was more common among men. The prevalence of astigmatism was 25.38%. It was interesting to note that against the rule astigmatism in contrast to other studies was observed more often (66.04%) in this study. Though the above study represent the regional population of limited age (20-40 years), but the differences in the pattern of refractive error in this study leads us to believe that genetic, racial, environmental and occupational influences may play an important role.

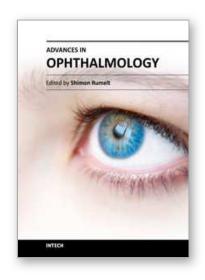
5. References

- [1] Resnikoff S, Pascolini D, Etya'ale D, et al. Global data onvisual impairment in the year 2002. *Bull World Health Organ*.2004;82:844 –51.
- [2] Munoz B, West SK, Rubin GS, Schein OD, Quigley HA, Bressler SB, Bandeen-Roche K. Causes of blindness and visual impairment in a population of older Americans.

 The Salisbury Eye Evaluation Study. *Arch Ophthalmol*. 2002;118:819–825.
- [3] Foran S, Wang JJ, Mitchell P. Causes of incident visual impairment. *Arch Ophthalmol*. 2002;120:613–619.
- [4] Weih LM, VanNewkirk MR, McCarty CA, Taylor HR. Age-specific causes of bilateral vision impairment. *Arch Ophthalmol*. 2000;118:264–269.
- [5] VanNewkirk MR, Weih L, McCarty CA, Taylor HR. Cause-specific prevalence of bilateral visual impairment in Victoria, Australia. The Visual Impairment Project. *Ophthalmology*. 2001;108:960–967.
- [6] Munoz B, West SK, Rodriguez J, Sanchez R, Broman AT, Snyder R, Klein R. Blindness, visual impairment and the problem of uncorrected refractive error in a Mexican-American population: Proyecto VER. *Invest Ophthalmol Vis Sci.* 2002;43:608–614.
- [7] Brown SA, Weih LM, Fu CL, Dimitrov P, Taylor HR, McCarty CA. Prevalence of amblyopia and associated refractive errors in an adult population in Victoria, Australia. *Ophthalmic Epdimol*. 2000;7:249-58.
- [8] Naiglin L, Gazagne C, Dallongeville F, Thalamas C, Idder A, Rascol O, Malecaze F, Calvas P. A genome wide scanfor familial high myopia suggests a novel locus on chromosome7q36. *J Med Genet*. 2002;39:118–124.
- [9] Young TL, Ronan SM, Drahozal LA, Wildenberg SC, Alvear AB, Oetting WS, Atwood LD, Wilkin DJ, King RA. Evidence that a locus for familial high myopia maps to chromosome 18p. *Am J Hum Genet*.1998;63:109–119.
- [10] The Framingham Offspring Eye Study Group. Familial aggregation and prevalence of myopia in the Framingham Offspring Eye Study. *Arch Ophthalmol*. 1996;114:326–332.
- [11] Wang Q, Klein BE, Klein R, Moss SE. Refractive status in the Beaver Dam Eye Study. *Invest Ophthalmol Vis Sci.* 1994;35:4344–4347.
- [12] Midelfart A, Aamo B, Sjohaug KA, Dysthe BE. Myopia among medical students in Norway. *Acta Ophthalmol*. 1992;70:317-322.
- [13] Dandona R, Dandona L, Naduvilath TJ, Srinivas M, McCarty CA, Rao GN. Refractive errors in an urban population in Southern India: the Andhra Pradesh Eye Disease Study. *Invest Ophthalmol Vis Sci.* 1999;40:2810–2818.
- [14] Wu HM, Seet B, Yap EPH, Saw SM, Lim TH, Chia KS. Does education explain ethnic differences in myopia prevalence? A population-based study of young adult males in Singapore. *Optom Vis Sci.* 2001;78:234–239.
- [15] Luke LK L, Yung-Feng S, Chong-Bin T, Chien-Jen C, Loung-An L, Port-Tying H, Ping-Kang H. Epidemiologic study of ocular refraction among schoolchildren in Taiwan in 1995. *Optom Vis Sci.* 1999;76:275–281.
- [16] Jadoon MZ, Dineen B, Bourne RRA, Shah SP, Khan MA, Johnson GJ, Gilbert CE, Khan MD. Refractive Errors in the Adult Pakistani Population: The National Blindness and Visual Impairment Survey. *Ophthalmic Epidemiol* 2008;15:183-190.
- [17] Katz J, Tielsch JM, Sommer A. Prevalence and risk factors forrefractive error in an adult inner city population. *Invest Ophthalmol Vis Sci.* 1997;38:334–340.

- [18] Attebo K, Ivers RQ, Mitchell P. Refractive errors in an older population: The Blue Mountains eye study. *Ophthalmology*. 1999;106:1066–1072.
- [19] Wong TY, Foster PJ, Hee J, Pin Ng T, Tielsch JM, Chew SJ, Johnson GJ and Seah SKL. Prevalence and risk factors of refractive errors in adult Chinese in Singapore. *Invest Ophthalmol Vis Sci.* 2000;41:2486–2494.
- [20] Wu SY, Nemesure B, Leske MC. Refractive errors in a black adult population: The Barbados Eye Study. *Invest Ophthalmol Vis Sci.* 1999;40:2179–2184.
- [21] Gwiazda J, Scheiman M, Mohindra I, Held R. Astigmatism in children: changes in axis and amount from birth to six years. *Invest Ophthalmol Vis Sci.* 1984;25:88–92.
- [22] Saw SM. A synopsis of the prevalence rates and environmental risk factors for myopia. *Clin Exp Optom.* 2003;86:289-294.
- [23] Guggenheim JA, Hill C, Yam TF. Myopia, genetics and ambient lighting at night in a UK sample. *Br J Ophthalmol*. 2003;87:580-582.
- [24] Dong X, Dong, Ayala M, Lofgren S, Soderberg PG. Ultraviolet radiation induced catract: age and maximum acceptable dosage. *Invest Ophthalmol Vis Sci.* 2003;44:1150-1154.
- [25] Saw SM, Gazzard G,Koh D, Farook M, Widjaja D, Lee J, Donald T. H. Tan DTH. Prevalence rates of refractive errors in Sumarta, Indonesia. *Invest Ophthalmol Vis Sci.* 2002;43:3174-3180.
- [26] Gudmundsdottir E, Jonasson F, Jonsson V, Stefansson E, Sasaki H, Sasaki K. "With the rule" astigmatism is not the rule in the elderly. *Acta Ophthalmol.* 2000;78:642-646.
- [27] Pensyl CD, Harrson RA, Simpson P, Waterbor JW. Distribution of astigmatism among sioux Indians in South Dakota. *J Am Optom Assoc.* 1997;68:425-431.
- [28] Goss DA. Meridonial Analysis of with the rule astigmatism in Oklahoma *Indians*. *Optom Vis Sci.* 1989;66:281-287.
- [29] Goel.B.S, Amblyopia: modern concept and management. In current topics in ophthalmology. I Gupta, A.K (ED), B.I Churchil Livingstone New Dehli, 1993, p
- [30] Karki KJD Prevalence of Amblyopia in ametropia in clinical set up. *Katmandu University Medical Journal*. 2006;4;470-73.





Advances in Ophthalmology

Edited by Dr Shimon Rumelt

ISBN 978-953-51-0248-9
Hard cover, 568 pages
Publisher InTech
Published online 07, March, 2012
Published in print edition March, 2012

This book focuses on the different aspects of ophthalmology - the medical science of diagnosis and treatment of eye disorders. Ophthalmology is divided into various clinical subspecialties, such as cornea, cataract, glaucoma, uveitis, retina, neuro-ophthalmology, pediatric ophthalmology, oncology, pathology, and oculoplastics. This book incorporates new developments as well as future perspectives in ophthalmology and is a balanced product between covering a wide range of diseases and expedited publication. It is intended to be the appetizer for other books to follow. Ophthalmologists, researchers, specialists, trainees, and general practitioners with an interest in ophthalmology will find this book interesting and useful.

How to reference

In order to correctly reference this scholarly work, feel free to copy and paste the following:

Ashok Kumar Narsani, Shafi Muhammad Jatoi, Mohan Perkash Maheshwari and Khairuddin Shah (2012). Incidence of Refractive Error and Amblyopia Among Young Adults - A Hospital Based Study, Advances in Ophthalmology, Dr Shimon Rumelt (Ed.), ISBN: 978-953-51-0248-9, InTech, Available from: http://www.intechopen.com/books/advances-in-ophthalmology/incidence-of-refractive-error-and-amblyopia-among-young-adult-a-hospital-base-study-



InTech Europe

University Campus STeP Ri Slavka Krautzeka 83/A 51000 Rijeka, Croatia Phone: +385 (51) 770 447

Fax: +385 (51) 686 166 www.intechopen.com

InTech China

Unit 405, Office Block, Hotel Equatorial Shanghai No.65, Yan An Road (West), Shanghai, 200040, China 中国上海市延安西路65号上海国际贵都大饭店办公楼405单元

Phone: +86-21-62489820 Fax: +86-21-62489821 © 2012 The Author(s). Licensee IntechOpen. This is an open access article distributed under the terms of the <u>Creative Commons Attribution 3.0</u> <u>License</u>, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.



