

FREQUENCY OF RETINOPATHY AMONG DIABETICS ADMITTED IN A TEACHING HOSPITAL OF LAHORE

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Background: Diabetic retinopathy is a complication of diabetes that affects the blood vessels of the retina and leads to blindness. Although 4 – 8 million diabetics exist in Pakistan, very little work has been done on this complication of diabetes. The present study was undertaken to estimate the frequency of retinopathy among diabetics admitted in a teaching hospital of Lahore. **Methods:** Every patient of diabetes mellitus, admitted in departments of Medicine, Surgery, Ophthalmology and Obstetrics & Gynecology at Sir Ganga Ram Hospital, Lahore between June 2001 and September 2001 was included in the study. After adequate mydriasis, detailed fundus examination using indirect ophthalmoscope was carried out to determine the presence of diabetic retinopathy and its type. **Results:** Out of 4414 admissions, 540 patients – 340 females and 196 males were diabetics. Among 540 diabetics, 132 had Type-1 diabetes (24.4%) while 408 had Type-2 diabetes (75.6%). The duration of diabetes ranged from 10 to 12 years. The prevalence of diabetes among admitted patients was 12.2%. Among these 540 diabetics, 180 had diabetic retinopathy showing a prevalence of 33.3%. Non-proliferative retinopathy was present in 21.5% diabetics and proliferative retinopathy among 11.8% diabetics. The prevalence of retinopathy was significantly higher ($P < 0.001$) among males (42.8%) as compared to females (27.9%). The prevalence was similar (33.3%) among both Type-1 as well as Type-2 diabetes as well as similar to that reported from other countries. **Conclusion:** Since 33.3% of our diabetic population is suffering from retinopathy – a condition amenable to timely and cost-effective treatment, every diabetic should be made aware of the importance of regular ophthalmologic examination.

INTRODUCTION

Diabetic retinopathy is a complication of diabetes that affects the blood vessels of the retina and leads to blindness. The progression of retinopathy is orderly, advancing from mild abnormalities, characterized by increased vascular permeability, to moderate and severe non-proliferative diabetic retinopathy (NPDR), characterized by the growth of new blood vessels on the retina and posterior surface of the vitreous¹.

Retinopathy is the commonest complication of diabetes. Surveys from developed countries show that at any time, up to 10% of people with diabetes will have retinopathy². The annual incidence of retinopathy requiring ophthalmological follow up or treatment has been reported to average 1.5% after one year³. Untreated, between 6-9% of the people with proliferative retinopathy or severe non-proliferative disease would become blind each year. Findings, consistent from study to study, make it possible to suggest that, after 15 years of diabetes, approximately 2% of people become blind, while about 10% develop severe visual handicap³.

The knowledge of the retinopathy status of an individual is one part of the whole process of care in diabetes. The need to screen for diabetic retinopathy is uncontroversial. Early detection of sight threatening retinopathy and treatment by laser therapy has been shown to be effective in preventing the onset of visual impairment.

According to National Health Survey of Pakistan⁴, the prevalence of diabetes among population aged ≥ 25 year is 4.2%, while Shera et al⁵⁻⁶ have estimated a prevalence of 9.1 to 13.7%. These figures translate into 4-8 million diabetics in the country. In spite of this high number of diabetics very little work⁷⁻⁸ has been done on macrovascular and microvascular complications of diabetes. The present study was undertaken to estimate the frequency of retinopathy in a group of diabetics attending a teaching hospital in Lahore.

MATERIAL AND METHODS

The study was conducted between June 2001 and September 2001 at Sir Ganga Ram Hospital, attached to Fatima Jinnah Medical College, Lahore. Every patient of diabetes mellitus admitted in departments of Medicine, Surgery, Ophthalmology and Obstetrics & Gynecology was included in the study.

After obtaining informed consent, the information was obtained regarding the type, onset, duration, treatment, nature of diabetic control, associated systemic diseases and ocular complications. Each patient underwent a detailed ophthalmological examination by the principal author.

After adequate mydriasis, detailed fundus examination using indirect ophthalmoscope was carried out to determine the presence of diabetic retinopathy and its type. Diabetic retinopathy was classified as non-proliferative, when microaneurysms, haemorrhages (dot, blot or flame shaped) or hard exudates were seen in any quadrant of retina. Proliferative diabetic retinopathy was diagnosed, if neovascularisation of the retina or iris or angle, pre retinal or vitreous haemorrhage, and or tractional retinal detachment was present. SPSS version 10 was used for data analysis.

RESULTS

Total number of admissions during the study period was 4414. Out of these admissions, 540 patients were diabetics. Three hundred and forty four diabetics (63.7%) were female and 196 were males (36.3%). Table-1 shows the distribution of cases according to various characteristics.

The frequency of diabetes among admitted patients was 12.2%. The age of the diabetics ranged between 40 and 60 years except those from Department of Obstetrics & Gynecology, majority of whom were less than 45 years of age. Out of these 540 diabetics, 132 had Type-1 diabetes (24.4%) while 408 had Type-2 diabetes (75.6%). The duration of diabetes ranged from 10 to 12 years.

Among 540 diabetics, 180 had diabetic retinopathy showing a prevalence of 33.3% (Table-2). Non-proliferative retinopathy was present in 21.5% diabetics and proliferative retinopathy among 11.8% diabetics. The prevalence of retinopathy was significantly higher ($P < 0.001$) among males (42.8%) as compared to females (27.9%). Proliferative retinopathy was significantly higher ($P < 0.01$) among males (18.1%) as compared to females (8.1%). However, the difference in the prevalence of non-proliferative retinopathy among males (24.5%) and females (19.8%) was not significant.

The prevalence of retinopathy was similar (33.3%) among both Type-1 as well as Type-2 diabetes. Non-proliferative retinopathy was more common (22.5%) in Type-2 diabetes as compared to Type-1 diabetes (18.2%), whereas proliferative retinopathy was more common in Type-1 (15.1%) as compared to Type-2 diabetes (10.8%). However, the differences were not statistically significant ($P > 0.05$).

DISCUSSION

Knowledge of the retinopathy status of an individual is one part of the whole process of care in diabetes. The implicit “gold standard” for identifying and grading retinopathy is a retinal examination using indirect biomicroscopy by a senior ophthalmologist or seven field stereoscopic photographs of each eye interpreted by experienced readers.

Several studies⁹⁻¹⁰ have reported the cost effectiveness of screening for retinopathy. They have established that screening for diabetic retinopathy saves vision at a relatively low cost and this cost is many times less than the disability payments provided to people who go blind in the absence of a screening programme. In 1983, the annual cost of treating a diabetic at risk of blindness was estimated to be GBP 387/- compared with welfare benefits paid to a blind person of GBP 3575/- per annum. Similar results were reported in more recent American and European studies¹¹⁻¹².

Table – 1: Characteristics of Study Population

Characteristic	Number	Percent
Sex		
Male	196	37.3
Female	344	63.7
Total	540	100.0
Department of Admission		
Medicine	304	56.2
Surgery	52	9.7
Obstetrics & Gynaecology	52	9.7
Ophthalmology	132	24.4
Total	540	100.0

Type of Diabetics		
Type 1	132	24.4
Type 2	408	75.6
Total	540	100.0

Table – 2: Frequency of diabetic retinopathy among 540 diabetics according to sex and type of retinopathy

Characteristic	Number Examined	Positive for Retinopathy	Frequency (%)
Sex			
Male	196	84	42.8
Female	344	96	27.9
Total	540	180	33.3
Type of Retinopathy			
Non Proliferative	540	116	21.5
Proliferative			
Total	540	64	11.8
	540	180	33.3

Table 3 summarizes the results of various studies on diabetic retinopathy in developing countries. Ghana¹³ and Spain¹⁴ showed a significantly lower prevalence of 22.4% and 20.9% respectively ($P < 0.01$). Saudi Arabia¹⁵, Sri Lanka¹⁶ and Brazil¹⁷ reported a prevalence of 31.3%, which is equal to our figure of 33.3%. The prevalences reported from Egypt¹⁸ (42%) and India¹⁹ (48.1%) are significantly higher ($P < 0.05$) from our findings, while the prevalence noticed among South African²⁰ diabetics (40.3%) is not significantly different from the prevalence noticed in our study ($P > 0.05$). The reported prevalence among 3000 diabetics from Karachi⁷ (26.1%) is significantly lower than our findings. This could be due to the younger age of that population and the shorter duration of disease. In that study, the duration of diabetics in 52.2% of those suffering from retinopathy was 10 years or less, whereas in our study the duration of diabetes was 10 year or more.

Diabetic retinopathy is a complication of both Type-1 and Type-2 diabetes mellitus.

Table – 3: Comparison with the prevalence of Retinopathy in Various Countries

Country	Author	Year	No of Subjects	Retinopathy (%)
Ghana	Ndiaye et al ¹³	1999	129	22.4
Spain	Lopez et al ¹⁴	2002	3544	20.9
Saudi Arabia	El. Asrar et al ¹⁵	1998	502	31.3
Sri Lanka	Fernando ¹⁶	1993	1003	31.3
Brazil	Gomes et al ¹⁷	2002	50	31.3
Egypt	Herman et al ¹⁸	1998	6052	42.0
India	Singh et al ¹⁹	2001	52	48.1
South Africa	Rotchford et al ²⁰	2002	203	40.3
Pakistan	Akhtar ⁷	1991	3000	26.1
Pakistan	Present Study	2003	540	33.3

Prevalence is related primarily to the duration of disease and secondarily to quality of blood sugar control. Aiello et al have shown that after 20 years, nearly all Type-1 diabetics and greater than 60% of Type-2 diabetics will have retinopathy regardless of diabetic control²¹. Vision-threatening retinopathy rarely occurs in Type-1 patients in the first 3-5 years after diagnosis or before puberty. Over the next 20 years, nearly all Type-1 patients will have some degree of retinopathy. Up to 21% of Type-2 patients have retinopathy at the time of initial diagnosis related to prolonged hyperglycemic (“borderline”) states. The majority (approximately > 60%) will develop retinopathy over the subsequent years²¹.

The Wisconsin Epidemiologic Study of Diabetic Retinopathy (WESDR) concluded that 3.6% of those diagnosed with Type-1 diabetes, and 1.6% of those diagnosed with Type-2 diabetes, were legally blind. For Type-1 diabetics, blindness was due to diabetic retinopathy in 86% of the cases. For Type-2 diabetics, blindness was related to retinopathy in 33% of the cases; the percentage was lower due to other ocular causes²².

The United Kingdom Prospective Diabetic Study (UKPDS) is the longest and largest study of Type-2 patients. The study revealed that improved control led to a reduction in retinopathy, a 25% reduction in overall microvascular complications and that one point decrease in HbA1c was associated with 35% reduction in risk for microvascular complications²³. UKPDS has further shown a slowed progression of retinopathy with improved control. The end result is preservation of sight, decreased morbidity and decreased need for more expensive intervention.

Screening saves vision at a relatively low cost, much lower than with later interventions like involving intra-ocular surgery. The personal and societal costs are reduced with increased productivity, decreased morbidity, and decreased disability. This is a valuable strategy to identify patients with asymptomatic macular edema and proliferative retinopathy. Timely intervention with laser photocoagulation, when appropriate, can prevent visual loss.

ACKNOWLEDGEMENT

The secretarial assistance of Mr. Mohammad Hanif and Mr. Liaquat Ali Butt is gratefully acknowledged.

REFERENCES

1. American Diabetes Association. Diabetic Retinopathy. Diabetes Care 2002;25:S90-S93

2. Mcleod BK, Thompson JR, Rosenthal AR. The prevalence of retinopathy in the insulin-requiring diabetic patients of an English country town. *Eye* 1988;2:424-430.
3. Amos AF, McCarty DJ, Zimmet P. The rising global burden of diabetes and its complications : estimates and projections to the year 2010. *Diabetic Med* 1997; 14 (Suppl) : S7-S85.
4. Pakistan Medical Research Council. National Health Survey of Pakistan, Islamabad: Pakistan Medical Research Council; 1997:54.
5. Shera AS, Rafique G, Khwaja IA, Baqai S, King H. Pakistan National Diabetes Survey: prevalence of glucose intolerance and associated factors in Baluchistan province. *Diabetes Res Clin Pract* 1999; 44: 49-58.
6. Shera AS, Rafique G, Khwaja IA, Ara J, Baqai S, King H. Pakistan national diabetes survey: prevalence of glucose intolerance and associated factors in Shikarpur, Sindh Province. *Diabet Med* 1995;12:1116-21.
7. Khan AJ. Prevalence of diabetic retinopathy in Pakistan subjects. A pilot study. *J Pak Med Assoc* 1991;41:49-50.
8. Wajid SA, Khan MD. Causes of irreversible blindness. *J Coll Physicians Surg Pak* 2001;11:561-5.
9. Klein R, Klein B, and Moss S. The Wisconsin Epidemiological Study of Diabetic Retinopathy: a review. *Diabetes Metab Rev* 1989; 5: 59-70.
10. Lairson DR, Pugh JA, Kapadia AS, Lorimor RJ, Jacobson J, Velez R. Cost- effectiveness of alternative methods for diabetic retinopathy screening. *Diabetes Care* 1992;15:1369-77.
11. Matz H, Falk M, Gottinger W, Kieslborch G. Cost-benefit analysis of diabetic eye disease. *Ophthalmologica* 1996; 210: 348-53.
12. James M, Turner DA, Broadbent DM, Vora J, Harding SP. Cost effectiveness analysis of screening for sight threatening diabetic eye disease. *BMJ* 2000; 320:1627-31.
13. Nadiaye MR, Cisse A, De Medeiros M, Wane A, Kameni A, Ndiaye-Roth PA, Seye-Ndiaye C, Ba EH, Ndiaye PA, Wade A. Prevalence of diabetic retinopathy at the Dakar University Hospital Center. *Dakar Med J* 1999; 44:158-61.
14. Lopez IM, Diez A, Velilla S, Rueda A, Alvarez A, Pastor CJ. Prevalence of diabetic retinopathy and eye care in a rural area of Spain. *Ophthalmic Epidemiol* 2002;9:205-14.
15. El-Asrar AM, Al-Rubeaan KA, Al-Amro Sa, Kangave D, Moharram OA. Risk factors for diabetic retinopathy among Saudi Diabetics. *Int Ophthalmol* 1998-99; 22:155-61.
16. Fernando Dj, Siribaddana S, De Silva, Subasinge Z. Prevalence of retinopathy in a Sri Lankan diabetes clinic. *Ceylon Med J* 1993; 38:120-3.
17. Gomes MB, Dorigo E, da Silva Junior GR, Goncalves MR, Neves R. Prospective study of development of microalbuminuria and retinopathy in Brazilian IDDM patients. *Acta Diabetol* 2000;37:19-25.
18. Herman WH, Aubert RE, Engelgau MM, Thompson TJ, Ali MA, Sous ES, et al. Diabetes mellitus in Egypt: glycaemic control and microvascular and neuropathic complications. *Diabet Med* 1998; 15:1045-51.
19. Singh SK, Behre A, Sing MK. Diabetic retinopathy and microalbuminuria in lean type 2 diabetes mellitus. *J Assoc Physicians India* 2001; 49:439-41.
20. Rotchford AP, Rotchford KM. Diabetes in rural South Africa-an assessment of care and complications. *S Afr Med J* 2002; 92:536-41.
21. Aiello LP, Gardiner TW, King GL, Blankenship G, Cavallerano JD, Ferris FL, et als: Diabetic Retinopathy (Technical Review). *Diabetes Care* 1998; 21:143-156.

22. Klein R, Klein BEK, Moss SE, Davis MD, DeMets KL. The Wisconsin Epidemiologic Study of Diabetic Retinopathy. *Ophthalmology* 1987; 94:747-753.
23. Kohner EM, Stratton IM, Aldington SJ, Holman RR, Matthews DR. UK Prospective Diabetes Study (UKPDS) Group. Relationship between the severity of retinopathy and progression to photocoagulation in patients with Type 2 diabetes mellitus in the UKPDS (UKPDS 52). *Diabet Med* 2001; 18:178-84.

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