Diabetic Retinopathy in Type II Diabetics Detected by Targeted Screening Versus Newly Diagnosed in General Practice

Swati Agarwal, ¹*Ms* (*Ophth*), Rajiv Raman, ¹*Ms* (*Ophth*), *DNB*, Rani Padmaja Kumari, ¹*Ms* (*Ophth*), *FNB* (*Retina*), Himanshu Deshmukh, ¹*Dip Ophth*, Pradeep G Paul, ²*BsMs*, *Msc* (*Epidemiol*), Perumal Gnanamoorthy, ²*Msc* (*Stat*), Govindasamy Kumaramanickavel, ³*MD*, Tarun Sharma, ¹*MD* (*Ophth*), *FRCS* (*Edin*)

Abstract

Introduction: The aim of this study was to compare the occurrence of diabetic retinopathy in targeted screening diabetic patients (Group I) with newly diagnosed diabetic patients in general practice (Group II). Materials and Methods: This was an observational cross-sectional study. Data were obtained from 25,313 subjects who participated in the diabetic screening camps, and 128 newly diagnosed diabetes who presented to the diabetic retinopathy screening camps in general practice in rural and urban south India. The study variables were collected from all patients who underwent eye examination from the target screening detected diabetics [(n = 173) Group I] and those newly diagnosed in general practice [(n = 128) Group II]. The variations in prevalence of diabetic retinopathy and sight-threatening diabetic retinopathy in Group I and Group II and the factors affecting it were identified. Results: The occurrence of diabetic retinopathy was 6.35% (95% CI, 2.5-9.5) in Group I and 11.71% (95% CI, 5.6-16.4) in Group II. No significant difference was observed on occurrence of diabetic retinopathy, including sightthreatening retinopathy, in rural versus urban population and in Group I versus Group II. Patients diagnosed in general practice (Group II) with systolic blood pressure (BP) >140 were more likely to have retinopathy (P = 0.02). <u>Conclusions</u>: Diabetic retinopathy including sightthreatening complications was found at the time of diagnosis of diabetes in the targeted screening group as well as in newly diagnosed diabetics in the general practice group.

Ann Acad Med Singapore 2006;35:531-5

sight-threatening complications develop.

practice (Group II).

Study Population

Patients and Methods

The purpose of this study was to compare the occurrence

of diabetic retinopathy in targeted screening diabetic patients

(Group I) with newly diagnosed diabetic patients in general

All patients were enrolled between June 2003 and

September 2004. For Group I (targeted screening), diabetic

screening camps were organised in the two rural districts -

Kanchipuram and Vellore (73 camps) Tamil Nadu, India -

and urban Chennai (4 camps), Tamil Nadu, India. Patients

above the age of 30 years were screened for diabetes.

Key words: Diabetes mellitus, Diabetic retinopathy, General practice, Screening

Introduction

The epidemic of type II diabetes mellitus is now recognised worldwide.¹In India, it has been estimated that the population with type II diabetes would increase by 150% in 2025.^{2,3} As the population with type II diabetes increases, so does the prevalence of diabetic retinopathy and other microvascular complications.^{3,4}

It has been observed that the actual onset of diabetes mellitus occurs much earlier (could be 9 to 12 years) before it is clinically presented and diagnosed.⁵ Since microvascular complications are directly related to the duration of diabetes mellitus, it is important to screen the general population in order to detect potential or pre-clinical diabetics.^{6,7} This will enable early detection of diabetic retinopathy before

¹ Shri Bhagvan Mahavir Vitreoretinal Services

² Department of Ophthalmic Epidemiology

³ Department of Genetics and Molecular Biology

Sankara Nethralaya, Chennai, India

Address for Reprints: Dr Tarun Sharma, Sankara Nethralaya, 18, College Road, Chennai, Tamilnadu, India 600006. Email: drtaruns@gmail.com

Random blood glucose was measured with a glucometer (Accutrend Alfa, Boehringer Mannheim, Germany) using the finger-prick capillary method. Target screening-detected diabetes was defined as one who had random blood glucose value of 200 mg/dL or more. Among 25,313 subjects who participated in the diabetic screening camps, 4345 were known diabetics and 1130 were screening-detected diabetics individuals. Of these 1130, 173 (15.3%) reported for eye examination in the diabetic retinopathy camp. Of these 173, 125 (72.3%) were from the rural camps and 48 (27.7%) from the urban camps. All screening detected diabetics in the camps were referred to the general physicians or diabetologists for further evaluation of their diabetes status and follow-up.

For Group II (newly diagnosed in general practice), recently diagnosed diabetes individuals (during the last one month) were enrolled. There were 128 patients. Of these, 80 (62.5%) were referred from the diabetic clinics (urban) and 48 (37.5%) from the general physicians (rural). Forty-one (32%) were on oral hypoglycaemic agents, 9 (7%) were on insulin and the rest [78 (60.9%)] were on diet control and exercise.

Study Data

The study variables were collected from all patients who underwent eye examination. These variables included demographic data, height and weight for calculating body mass index (BMI) (normal range, up to 22.9; overweight, more than 23), blood pressure, physical activity (sedentary - rarely participate in any physical activity), smoking and alcohol status. Eye examination included assessment of Snellen's' visual acuity, slit-lamp evaluation (using handheld Heines HSL 100 CE), measurement of intraocular pressure and dilated fundus examination using binocular indirect ophthalmoscopy (Keeler Instruments Inc. PA, USA) and +20 D lens (Nikon). Fundus examination of all the patients was done by an experienced retinal specialist. Diabetic retinopathy was clinically graded as per the guidelines of Early Treatment Diabetic Retinopathy Study classification.8

Statistical Analysis

The SPSS (version 9.0) programme was used for statistical analysis. The prevalence was expressed as percentages with 95% confidence interval. Significance tests such as chi-square, *t*-test and Z test were applied at appropriate situation. *P* value <0.05 was considered statistically significant. The 2 groups were compared using Student's *t*-test for continuous variable and χ^2 test for dichotomous variables.

Results

Table 1 shows the baseline characteristics of Group I

(n = 173) and Group II diabetics (n = 128). The 2 groups were similar in terms of mean age, BMI, use of alcohol, smoking habits, physical activity and blood pressure. There were more males in Group I (P = 0.0005).

Table 2 shows the occurrence of diabetic retinopathy at the time of diagnosis of diabetes in Groups I and II. The diagnosis of diabetic retinopathy was made in 6.4% (95% CI, 2.5-9.5) and 11.7% (95% CI, 5.6-16.4) of individuals in Camps I and II, respectively. Sight-threatening diabetic retinopathy was observed in 4.6% in Group I and 7.8% in Group II. No significant difference was observed on occurrence of diabetic retinopathy in rural versus urban population in both groups.

Table 3 shows the influence of various factors on occurrence of diabetic retinopathy. None of the variables

 Table 1. Characteristics of Diabetic Patients in Group I (Targeted Screening) and Group II (Newly Diagnosed in General Practice)

Variable (Group I (n = 173) n (%)	Group II (n = 128) n (%)	Р
Age (y) Mean ± SD	54 ± 11	52 ± 12	0.13
Gender Male	103 (60)	50 (39)	0.0005
Physical activity Sedentary	67 (39)	57 (44)	0.34
BMI (Mean ± SD)	24 ± 4	24 ± 4	0.99
Smoking status Non-smoker	151 (87)	118 (92)	0.19
Alcohol status Non-drinker	146 (84)	116 (91)	0.12
Systolic BP (Mean ± SI	D) 132 ± 19	133 ± 20	0.65
Diastolic BP (Mean ± S	D) 82 ± 10	83 ± 11	0.41

BMI: body mass index; BP: blood pressure; SD: standard deviation

Table 2. Occurrence of Diabetic Retinopathy at the Time of Diagnosis in Group I (Targeted Screening) and Group II (Newly Diagnosed in General Practice)

Variable	Group I n = 173 (%)	Group II n = 128 (%)	Р
DR	11/173 (6.35%) (95% CI, 2.5-9.5)	15/128 (11.71%) (95% CI, 5.6-16.4)	0.22
NSTDR	3/173 (1.73%) (95% CI, 0.21-3.67)	5/128 (3.90%) (95% CI, 0.54-7.25)	0.43
STDR	8/173 (4.62%) (95% CI, 1.49-7.74)	10/128 (7.81%) (95% CI, 3.16-12.45)	0.36

DR: diabetic retinopathy; NSTDR: non-sight threatening diabetic retinopathy includes mild and moderate non-proliferative diabetic retinopathy; STDR: sight threatening diabetic retinopathy includes severe non-proliferative diabetic retinopathy, proliferative diabetic retinopathy and clinically significant macular oedema

Variable	Group I			Group II		
	n	Prevalence of DR (%)	Р	n	Prevalence of DR (%)	Р
Age (y)						
<49	53	3/53 (5.7)	0.55	43	6/43 (14)	0.386
>50	120	8/120 (6.7)		85	9/85 (10.6)	
Gender						
Male	103	7/103 (6.8)	0.52	50	9/50 (18)	0.07
Female	70	4/70 (5.7)		78	6/78 (7.7)	
Physical activity						
Sedentary	67	2/67 (3)	0.129	57	10/57 (17.5)	0.06
Non-sedentary	106	9/106 (8.5)		71	5/71 (7)	
BMI						
Normal	76	6/76 (7.9)	0.335	57	8/57 (14)	0.323
Overweight	97	5/97 (5.2)		71	7/71 (9.9)	
Smoking						
Non- smoker	151	9/151 (6)	0.42	118	15/118 (12.7)	0.274
Alcohol status						
Non-drinker	146	8/146 (5.5)	0.236	116	15/116 (12.9)	2.208
Systolic BP						
≤139	112	8/112 (7.1)	0.414	79	5/79 (6.3)	0.018
≥140	61	3/61 (4.9)		49	10/49 (20.4)	
Diastolic BP						
≤89	104	7/104 (6.7)	0.536	80	10/80 (12.5)	0.479
≥90	69	4/69 (5.8)		48	5/48 (10.4)	

Table 3. Influence of Variables on Diabetic Retinopathy in Groups I and II

DR: diabetic retinopathy; BMI: body mass index; BP: blood pressure

showed any influence in Group I, but systolic blood pressure >140 mm Hg was associated with high occurrence of diabetic retinopathy in Group II (P = 0.02).

Discussion

This paper highlights that one of the microvascular complications - diabetic retinopathy - in individuals with type II diabetes appeared to be somewhat higher in newly diagnosed patients from general practice (11.7%) compared to those diagnosed in the targeted screening group (6.4%). However, this difference did not reach statistical significance. Table 4 shows the prevalence of diabetic retinopathy in published reports, both in the diabetic clinics and in the general population. As expected, the prevalence of diabetic retinopathy was higher in those who were examined in the diabetic clinics than in the populationbased screening. The prevalence of diabetic retinopathy ranged from 7.3% to 25.5% in those diabetics who were detected in the diabetic clinics and 1.9% to11.2% in diabetics in the population-based screening.9,11-23 The reason for this difference could be that those who attended the diabetic clinics could have been symptomatic and had diabetes for longer period than those who attended the targeted screening camps. However, surprisingly the difference in prevalence of diabetic retinopathy between the two groups was not observed (7.6% in targeted screening vs 1.9% in

general practice) in the Hoorn Screening study, possibly due to the small sample size in the general practice group.9

Among those detected to have diabetic retinopathy (in the present study), sight-threatening diabetic retinopathy was evident in 4.6% in the targeted screening group and 7.8% in the general practice group. This is important as these patients could be referred to our base hospital for further investigations such as fluorescein angiography and prompt laser photocoagulation. In doing so, we hoped to avoid potential complications of proliferative diabetic retinopathy and stabilise vision in eyes with diabetic maculopathy. However, it is worth mentioning that of the 1130 diabetic individuals detected in Group I, only 15.3% attended the diabetic retinopathy camp. Non-response of such a magnitude calls for creating greater awareness among the masses on diabetes and its microvascular complications.²⁶ Being a progressive disorder, a one-time screening effort would not suffice to alleviate the socioeconomic burden of diabetic retinopathy. These screening programmes need to be regularly repeated in order to reduce the visual morbidity of type II diabetics.

Our study did not elucidate any risk factors related to diabetic retinopathy in Group 1; only in Group II, one factor - systolic blood pressure - was found to be related to diabetic retinopathy. Likewise, in a study done by Rema

Year, Author	Country (n)	Age/Mean age (y)	Prevalence of DR (%)
Clinic-based Studies			
2005, Al Zuabi et al ¹¹	Kuwait (92)	-/-	7.6
2003, Spijkerman et al ⁹	Netherlands (60)	$>50/61.4 \pm 7.0$	1.9
2002, Talu et al ¹²	Romania (487)	>40/58.5	14.3
2002, Chowdhury et al ¹³	England (292)	<40/ South Asians 35.3 years Europeans 33.5 years	South Asians 17.5 Europeans 7.9
2002, Liu et al ¹⁴	China (773)	-/58	21
2001, Unuigbe et al ¹⁵	Nigeria (66)	-/-	23
2001, Tzeng et al ¹⁶	Taiwan (148)	-/-	25.5
2000, Rema et al ¹⁷	India (438)	-/48 ± 14	7.3
1998, Weerasuriya et al18	Sri Lanka (597)	$-/42.3 \pm 6.2$	15
1998, Wang et al ¹⁹	Hong Kong (474)	$-/53.6 \pm 0.6$	21.9
Present study	India (128)	$40/52 \pm 12$	11.7
Population-based Studies			
2005, Rema et al ¹⁰	India (354)	≥20/48 ± 12	5.1
2003, Spijkerman et al ⁹	Netherlands (195)	$>50/63.4 \pm 7.0$	7.6
2002, Van Leiden et al ²⁰	Netherlands (626)	$>50/65.7 \pm 6.6$	9
2001, Perusicova ²¹	Czech Republic (314)	20-65	3.5
2001, de Fine Olivarius et al ²²	Denmark (1251)	≥40/-	5
2001, West et al ²³	United States (4774)	≥40/-	9
1997, Klein et al ²⁴	United States (354)	≤30 type I/-	1.3
1997, Nagi et al ²⁵	United Kingdom (169)	≥15/-	11.2
Present study	India (173)	$40/54 \pm 11$	6.35

Table 4. Comparison with Published Reports

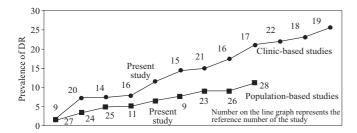


Fig. 1. Comparison with published reports.

et al,¹⁰ none of the clinical or biochemical variables were associated with diabetic retinopathy. In contrast, the Hoorn Screening study noted that blood pressure, lipid concentration and BMI are associated with microvascular complications.⁹

Figure 1 shows that the prevalence of diabetic retinopathy is always high in clinic-based studies. Estimation of diabetic retinopathy occurrence in the present study was somewhat in the mid-range in Group I, but at the junction of one-third and upper two-thirds in Group II

Though the gold standard technique for the detection of diabetic retinopathy is fundus photography, we used indirect

ophthalmoscopy as described by Dandona et al²⁷ in a population-based study. Moss et al²⁸ reported that sensitivity and specificity of indirect ophthalmoscopy for detecting any retinopathy was 82 % (95% CI, 80-84) and 95% (95% CI, 94-96), and for sight threatening retinopathy, 72% (95% CI, 73-86) and 100% (95% CI, 99-100), respectively. The British Diabetic Association recommended that for any screening tool to be able to detect diabetic retinopathy, the sensitivity should be around 80%, and specificity, 95%. Availability of HiMag attachment with Keeler indirect ophthalmoscope allowed us to view the macular area with high magnification (5 times); hence, this tool could be used as an initial screening procedure for mass community screening for diabetic retinopathy.

In conclusion, diabetic retinopathy including sightthreatening complications was found at the time of diagnosis of diabetes in targeted screening group as well as in newly diagnosed diabetics in general practice group.

Acknowledgements

We would like to acknowledge the Lions Club International Foundation and the RD Tata Trust for their financial support.

REFERENCES

- Currie CJ, Kraus D, Morgan CL, Gill L, Stott NC, Peters JR. NHS acute sector expenditure for diabetes: the present, future, and excess in-patient cost of care. Diabet Med 1997;14:686-92.
- Wild S, Roglic G, Green A, Sicree R, King H. Global prevalence of diabetes: estimates for the year 2000 and projections for 2030. Diabetes Care 2004;27:1047-53.
- Ramachandran A, Jali MV, Mohan V, Snehalatha C, Viswanathan M. High prevalence of diabetes in an urban population in South India. BMJ 1988;297:587-90.
- Namperumalsamy P, Nirmalan PK, Ramasamy K. Developing a screening program to detect sight-threatening diabetic retinopathy in South India. Diabetes Care 2003;26:1831-5.
- Harris MI, Klein R, Welborn TA, Knuiman MW. Onset of NIDDM occurs at least 4-7 yr before clinical diagnosis. Diabetes Care 1992;15:9.
- Thompson TJ, Engelgau MM, Hegazy M, Ali MA, Sous ES, Badran A, et al. The onset of NIDDM and its relationship to clinical diagnosis in Egyptian adults. Diabet Med 1996;13:337-40.
- American Diabetic Association. Screening for type 2 diabetes. Diabetes Care 2004;27Suppl1:S11-S14.
- Early Treatment Diabetic Retinopathy Study Research Group. (Early photocoagulation for diabetic retinopathy. ETDRS report No. 9). Ophthalmology 1991;98(5 Suppl):766-85.
- Spijkerman AM, Dekker JM, Nijpels G, Adriaanse MC, Kostense PJ, Ruwaard D, et al. Microvascular complications at time of diagnosis of type 2 diabetes are similar among diabetic patients detected by targeted screening and patients newly diagnosed in general practice: the hoorn screening study. Diabetes Care 2003;26:2604-8.
- Rema M, Premkumar S, Anitha B, Deepa R, Pradeepa R, Mohan V. Prevalence of diabetic retinopathy in urban India: the Chennai Urban Rural Epidemiology Study (CURES) eye study, I. Invest Ophthalmol Vis Sci 2005;46:2328-33.
- Al-Zuabi H, Al-Tammar Y, Al-Moataz R, Al-Sabti K, Wani VB, Hamama F, et al. Retinopathy in newly diagnosed type 2 diabetes mellitus. Med Princ Pract 2005;14:293-6.
- Talu S, Kaucsar E, Soreanu A. Diabetic retinopathy in newly diagnosed patients with type II diabetes mellitus [Romanian]. Ophthalmologia 2002;54:27-30.
- Chowdhury TA, Lasker SS. Complications and cardiovascular risk factors in South Asians and Europeans with early-onset type 2 diabetes. QJM 2002;95:241-6.
- 14. Liu DP, Molyneaux L, Chua E, Wang YZ, Wu CR, Jing H, et al. Retinopathy in a Chinese population with type 2 diabetes: factors affecting the presence of this complication at diagnosis of diabetes.

Diabetes Res Clin Pract 2002;56:125-31.

- Unuigbe EI, Omeife H, Edema T, Ukoli FA. Microalbuminuria and associated factors in newly diagnosed diabetics. Niger Postgrad Med J 2001;8:187-92.
- Tzeng TF, Hsiao PJ, Hsieh MC, Shin SJ. Association of nephropathy and retinopathy, blood pressure, age in newly diagnosed type 2 diabetes mellitus. Kaohsiung J Med Sci 2001;17:294-301.
- Rema M, Deepa R, Mohan V. Prevalence of retinopathy at diagnosis among type 2 diabetic patients attending a diabetic centre in South India. Br J Ophthalmol 2000;84:1058-60.
- Weerasuriya N, Siribaddana S, Dissanayake A, Subasinghe Z, Wariyapola D, Fernando DJ. Long-term complications in newly diagnosed Sri Lankan patients with type 2 diabetes mellitus. QJM 1998;91:439-43.
- Wang WQ, Ip TP, Lam KS. Changing prevalence of retinopathy in newly diagnosed non-insulin dependent diabetes mellitus patients in Hong Kong. Diabetes Res Clin Pract 1998;39:185-91.
- 20. Van Leiden HA, Dekker JM, Moll AC, Nijpels G, Heine RJ, Bouter LM, et al. Blood pressure, lipids, and obesity are associated with retinopathy: the hoorn study. Diabetes Care 2002;25:1320-5.
- Perusicova J. Prevalence of dyslipidaemia, hypertension and vascular complications in newly diagnosed diabetics (prospective study: part 2) [Czech]. Vnitr Lek 2001;47:146-50.
- 22. de Fine Olivarius N, Nielsen NV, Andreasen AH. Diabetic retinopathy in newly diagnosed middle-aged and elderly diabetic patients. Prevalence and interrelationship with microalbuminuria and triglycerides. Graefes Arch Clin Exp Ophthalmol 2001;239:664-72.
- 23. West SK, Klein R, Rodriguez J, Munoz B, Broman AT, Sanchez R, et al. Diabetes and diabetic retinopathy in a Mexican-American population: Proyecto VER. Diabetes Care 2001;24:1204-9.
- Klein R, Palta M, Allen C, Shen G, Han DP, D'Alessio DJ. Incidence of retinopathy and associated risk factors from time of diagnosis of insulindependent diabetes. Arch Ophthalmol 1997;115:351-6.
- Nagi DK, Pettitt DJ, Bennett PH, Klein R, Knowler WC. Diabetic retinopathy assessed by fundus photography in Pima Indians with impaired glucose tolerance and NIDDM. Diabetic Med 1997;14:449-56.
- 26. Will JC, German RR, Schuman E, Michael S, Kurth DM, Deeb L. Patient adherence to guidelines for diabetic eye care: results from the diabetic eye disease follow-up study. Am J Public Health 1994;84:1669-71.
- Dandona L, Dandona R, Naduvilath TJ, Mc Carty CA, Rao GN. Population based assessment of diabetic retinopathy in an urban population in southern India. Br J Ophthalmol 1999;83:937-40.
- Moss SE, Klein R, Kessler SD, Rechie KA. Comparison between ophthalmoscopy and fundus photography in determining severity of diabetic retinopathy. Ophthalmology 1985;92:62-7.