

Progression to blindness in 20 years among patients with glaucomatous visual field loss in a tertiary hospital in the Philippines

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ABSTRACT

Objective To determine visual field (VF) rates of change among patients with glaucomatous VF loss and proportion of those becoming blind based on residual life expectancy and factors associated with fast progression.

Methods and analysis This was a retrospective study of the VFs of patients with glaucomatous VF defects in at least one eye. Baseline and final VFs were reviewed. Rates of VF change (decibels (dB)/year) for each eye, together with the residual life expectancy based on age and sex, were used to predict mean deviation/defect (MD) at the end of expected lifetime. Blindness was defined if computed MD was 22 dB (Octopus) or -22 dB (Humphrey) or worse in the better eye. Factors associated with fast progression (>1 dB/year for Octopus or <-1 dB/year for Humphrey) and blindness were determined.

Results There were 1016 eyes of 583 patients eligible. There was decline in VF MD/year in 613/1016 (60.3%), 95% CI (57.3% to 63.3%) of eyes; however, only 98/1016 (9.7%), 95% CI (7.9% to 11.5%) of eyes showed fast progression. Among patients with bilateral VFs, 43/433 (9.9%), 95% CI (7.1 to 12.8) of eyes were predicted to progress to blindness. In multivariate analysis, factors associated with fast progression were baseline MD ($p<0.001$) and male sex ($p=0.041$). Factors associated with blindness were age <60 years ($p=0.003$), baseline MD ($p=0.022$), bilateral glaucomatous VF defects ($p<0.001$) and fast progression ($p<0.001$).

Conclusion Patients reaching blindness in a routine clinical setting was 10%. Because of association of age and baseline MD on blindness, early disease detection is important. VF progression rates and residual life expectancy must be incorporated in glaucoma care.

Key messages

What is already known about this subject?

- ▶ A UK study revealed that the development of blindness using visual fields in a clinic setting is about 5%.

What are the new findings?

- ▶ Our study in the Philippines showed it is 10%.

How might these results change the focus of research or clinical practice?

- ▶ The rates of visual field progression and the expected life expectancy are important considerations in glaucoma care.

(LV) or pattern standard deviation (PSD). MD represents the difference in the overall sensitivity of the VF compared with the age-matched normal sensitivity values.² LV or PSD, which is more specific for localised VF loss, has been shown to underestimate true change as some diffuse loss also occurs in glaucoma.³ In the Early Management Glaucoma Trial and Collaborative Normal Tension Glaucoma Study, the mean rate of progression was -1.08 decibels (dB)/year and -0.2 to -2 dB/year, respectively.^{4,5} In a study by Heijl *et al*, among patients with glaucoma under clinical care, the mean rate of progression was -0.80 dB/year. However, fast progressors were also common.⁶ Determining rate of change would entail frequent VF testing to establish a trend. Once detected, the individual's residual life expectancy or life expectancy adjusted for age must be taken into consideration.⁷ Using these data, individualised care and avoidance of excess treatment and monitoring can be attained.

VF rates of change and the risk factors for progression among patients with glaucoma have been investigated in numerous studies.⁸ However, most of these studies are limited to the VF result only and did not consider estimates of blindness based on a

Glaucoma is a progressive disease that causes optic nerve head and visual field (VF) changes. Intraocular pressure (IOP) is considered the only modifiable risk factor in glaucoma and it is often given a significant role during treatment decisions.¹ However, because functional vision is a major determinant of quality of life, the rate of VF progression should be included in clinical management. The most common indices (depending on the instrument) to monitor for VF change are the mean defect or deviation (MD) and the loss variance



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patient's residual life expectancy. Furthermore, these types of studies are currently lacking in the Philippines. The significance of providing local data on the status of glaucomatous VF loss and the number of patients at risk of becoming blind within their lifetime will help determine whether efforts in the management of those with glaucomatous VF damage are adequate and properly administered. This study aims to help direct future efforts and resources in the care of patients with glaucoma.

The goal of this study is to provide information on the current trend of field progression among patients with glaucomatous field loss using data collected from a Philippine tertiary hospital. Specifically, this study aims to determine the rates of field change based on MD, the proportion of patients that will become blind during their lifetime and the factors associated with a fast rate of field loss and progression to blindness at death.

METHODS

This was a retrospective cohort study using VF test data done at St. Luke's Medical Center, Quezon City, Metro Manila, Philippines of patients with glaucomatous VF defects. There was no direct patient and public involvement in this study as the anonymised VF data were collected and analysed without compromising data privacy of the subjects.

Patients with glaucomatous VF defects with at least five reliable VF tests using the same test programme for at least 3 years from July 1998 to April 2018 were included in the study. For the Octopus VF, a glaucomatous VF defect was defined as (1) MD >2dB and/or LV >6dB; and (2) presence of at least three contiguous abnormal points with p values less than 5% in the arcuate area in the pattern deviation plot on two consecutive examinations.⁹ For the Humphrey VF, a glaucomatous VF defect was defined as (1) a glaucoma hemifield test outside normal limits on at least two fields; (2) a cluster of three or more non-edge points, two of which are depressed on the pattern deviation plot at a p value of less than 5% and one of which is depressed at a p value of less than 1% on two consecutive fields; or (3) a pattern standard deviation (PSD) that occurs in less than 5% of normal fields on two consecutive fields.¹⁰ Unreliable VFs, defined to have a false-positive score of 20% or more, false-negative score of 33% or more, or fixation loss score of 20% or more, and a reliability factor of >15% (Octopus) were excluded. Patients 18 years old and below, as well as those who are blind at the start of the study (an MD of ≥22dB for the Octopus or ≤-22dB for the Humphrey in the better eye)¹¹ based on the baseline MD, were not included in the study.

VF test results of patients with glaucomatous VF defects were reviewed. Only VFs recorded on the Octopus VF (Haag-Streit, Koeniz, Switzerland) using the tG2 programme (central 30 degrees, size III white stimulus and TOP strategy) or the dG2 programme (central 30 degrees, size III white stimulus and Dynamic strategy) or on the Humphrey VF (Carl Zeiss Meditec, USA) using the

Table 1 Age-adjusted life expectancy based on the WHO Global Health Observatory data repository on the Philippines for the year 2015¹²

		Male	Female
Expectation of life at age x	<1 year	65.3	72
	1–4 years	66	72.4
	5–9 years	62.4	68.8
	10–14 years	57.5	63.9
	15–19 years	52.7	59.1
	20–24 years	48	54.3
	25–29 years	43.5	49.5
	30–34 years	39.1	44.8
	35–39 years	34.7	40.1
	40–44 years	30.4	35.5
	45–49 years	26.2	31
	50–54 years	22.3	26.7
	55–59 years	18.7	22.5
	60–64 years	15.5	18.6
	65–69 years	12.7	14.9
	70–74 years	10	11.5
	75–79 years	7.5	8.5
	80–84 years	5.5	5.9
	85+ years	4	4

WHO, World Health Organization.

30-2 programme (central 30 degrees, size III white stimulus and SITA-Fast or Standard strategy) were included in the study. The presence of a glaucomatous pattern of VF defects was determined based on the criteria stated previously. To eliminate learning effects, the first VF examination was disregarded.

The following data from the VF printout were obtained, tabulated and organised using Microsoft Excel 2011: birthdate, ages on the first and last VF test, sex, type of VF programme used, duration of follow-up, number of VF tests performed, baseline and final MD. The first and last recorded MD were referred to as the baseline and final MD, respectively. Rates of MD change were calculated in dB/year using ordinary least squares regression. MD at the expected death was determined by multiplying the rate of change and the patient's residual life expectancy. Residual life expectancies, based on age and sex, were collected from the World Health Organization Global Health Observatory data repository on the Philippines for the year 2015 (table 1).¹²

An MD of 22 dB (Octopus) or -22 dB (Humphrey) in the better eye is the value that corresponds to the US Social Security Act definition of 'statutory blindness'.¹¹ When determining blindness, both eyes had reached the necessary level of VF loss. If patients had fewer than five VFs performed in one of their eyes, then their baseline MDs were recorded and the eyes were either deemed to be stable (0 dB/year) or progressing (1.0 dB/year for

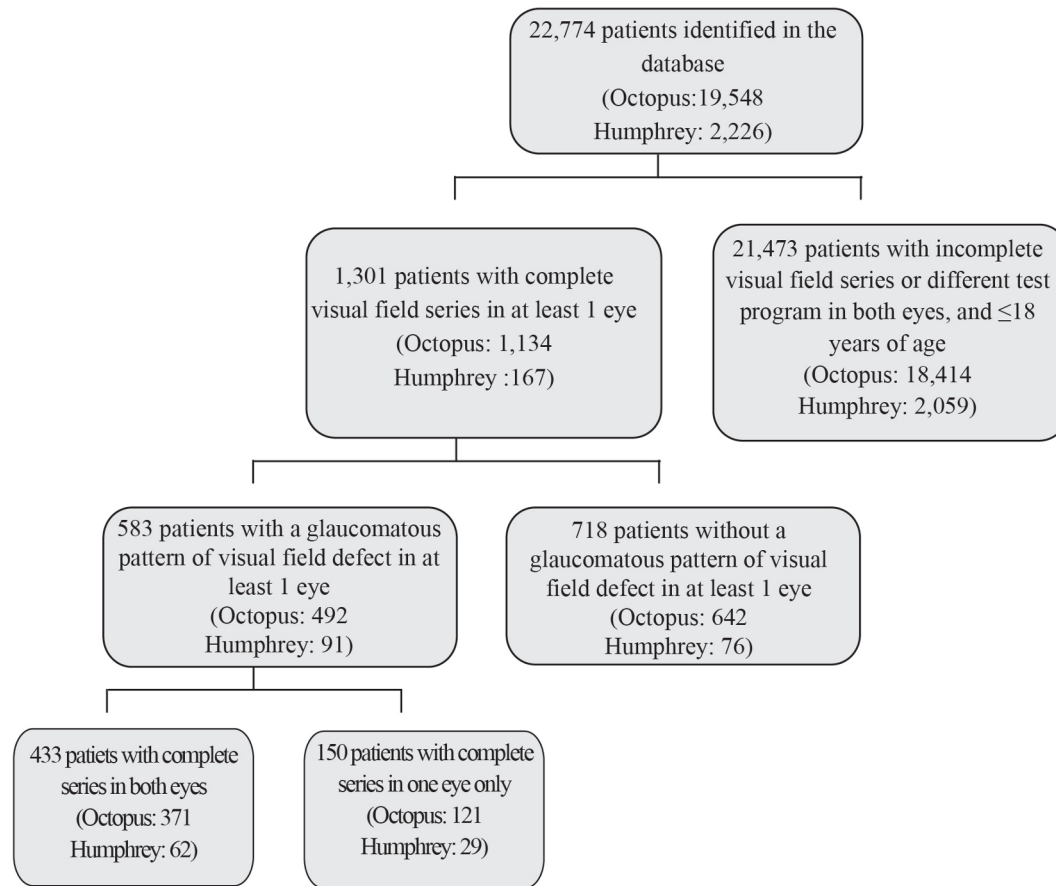


Figure 1 Flow chart of patient selection process.

Octopus or -1.0 dB/year for Humphrey) representing a 'best'-case and 'worst'-case scenario, respectively. If there was no recorded VF for the other eye, the eye was recorded as either healthy or blind representing a 'best'-case and 'worst'-case scenario, respectively. Fast rate of progression was defined as VF worsening of >1 dB/year (Octopus) or <-1 dB/year (Humphrey).⁶

RESULTS

Out of the 22,774 patients identified in the Octopus (19,548) and Humphrey (2,226) VF databases, only 583 patients met the inclusion criteria (figure 1). Four hundred thirty-three patients had VF series in both eyes, while 150 patients had series in only one eye. The characteristics of the patients in the study are given in table 2. Overall, majority of patients were female (62.4%) with a mean baseline age of 61.1 ± 12.6 years and a mean final age of 66.2 ± 12.5 . The mean residual life expectancy based on the final age was 18.8 ± 9.6 years. The most common VF programme used was the Octopus tG2 (50.9%) with a mean follow-up period of 5.1 ± 2.3 years and 6.6 ± 2.0 examinations.

Compared with the mean baseline MD, a significant ($p < 0.001$) decline in the mean final MD was only observed for the better eye using the Octopus VF. The other MD changes were non-significant (table 2, figure 2A,B).

Figure 3 demonstrates the distribution of MD rates of change in all eyes. It is evident that majority of eyes worsened at a rate between 0 and 0.5 dB/year for the Octopus (35.7%, 95% CI 32.5% to 38.7%) and improved at the same rate for the Humphrey VF (32.7%, CI 25.3% to 40.1%). Overall, 60.3% (CI 57.3% to 63.3%) showed a decline in MD per year; however, only 9.7% (CI 7.9% to 11.5%) showed fast progression. Noticeably, 39.3% (CI 36.3% to 42.3%) showed improvement.

Table 3 shows the univariate analysis of the factors associated with a fast rate of progression. Majority of the fast progressors were male, 60 years old and above, and had a glaucomatous pattern of VF defect on both eyes. Their mean baseline MD was 7.7 ± 3.9 for the Octopus and -9.5 ± 4.8 for the Humphrey.

Baseline MD was shown to be the only statistically significant factor associated with a fast rate of progression but after controlling for the baseline MD in the multivariate analysis (table 4), sex became statistically significant as well. Males have 1.56 times the odds of fast progression, holding the baseline MD constant while the odds of fast progression decreases by 6% for every 1 dB worsening in baseline MD, holding sex constant.

Only 9.9% (95% CI 7.11 to 12.75) of patients with VF series in both eyes progressed to blindness in their expected lifetime (table 5). The 'best-case scenario'

Table 2 Patient demographics

Measure	All patients, n=583	Patients with series in both eyes, n=433		Patients with series in one eye only, n=150
Patients with glaucomatous VF defects in both eyes, n (%)	251 (43.1)	251 (58.0)		NA
Sex, n (%)				
Male	219 (37.6)	137 (31.6)		82 (54.7)
Female	364 (62.4)	296 (68.4)		68 (45.3)
Baseline age				
Mean±SD	61.1±12.6	61.5±11.5		59.8±15.1
Range	20 to 90	24 to 86		20 to 90
Final age				
Mean±SD	66.2±12.5	66.8±11.5		64.5±15.0
Range	25 to 93	31 to 91		25 to 93
Residual life expectancy				
Mean±SD	18.8±9.6	18.5±8.7		19.6±11.7
Range	2.7 to 54.3	2.7 to 49.5		4 to 54.3
VF programme, n (%)				
Octopus dG2	195 (33.4)	143 (33.2)		52 (34.7)
Octopus tG2	297 (50.9)	228 (52.4)		69 (46.0)
Humphrey 30-2	91 (15.6)	62 (14.4)		29 (19.3)
Follow-up period (years)				
Mean±SD	5.1±2.3	5.2±2.3		4.6±2.1
Range	2 to 15	2 to 15		2 to 14
Number of VF tests				
Mean±SD	6.6±2.0	6.7±2.0		6.4±1.9
Range	5 to 20	5 to 20		5 to 15
Baseline MD		Better eye	Worse eye	
Octopus				
Mean±SD		5.1±4.5	10.3±5.5	11.8±5.8
Range		-2.2 to 21.3	1.1 to 22.4	1.4 to 21.6
Humphrey				
Mean±SD		-5.2±6.7	-12.6±7.1	-14.1±7.8
Range		-21.8 to 0.9	-22.4 to -2.1	-21.8 to -2.4
Final MD				
Octopus				
Mean±SD		6.3±5.2	11±5.8	13.2±5.8
Range		-2 to 21.3	1.9 to 22.9	2.7 to 22.5
Humphrey				
Mean±SD		-4.8±5.8	-13.3±7.0	-14.9±7.5
Range		-20.9 to 0.7	-23.1 to -2.6	-20.0 to -3.7
Rate of VF field loss (dB/year)				
Octopus				
Mean±SD		0.17±0.69	0.16±0.87	0.30±0.79
Range		-3.40 to 4.10	-2.90 to 3.53	-2.00 to 3.37
Humphrey				
Mean±SD		0.08±0.40	-0.15±0.68	-0.20±0.80
Range		-0.75 to 1.56	-1.97 to 1.68	-1.90 to 1.50

MD, mean deviation ; SD, standard deviation ; VF, visual field.

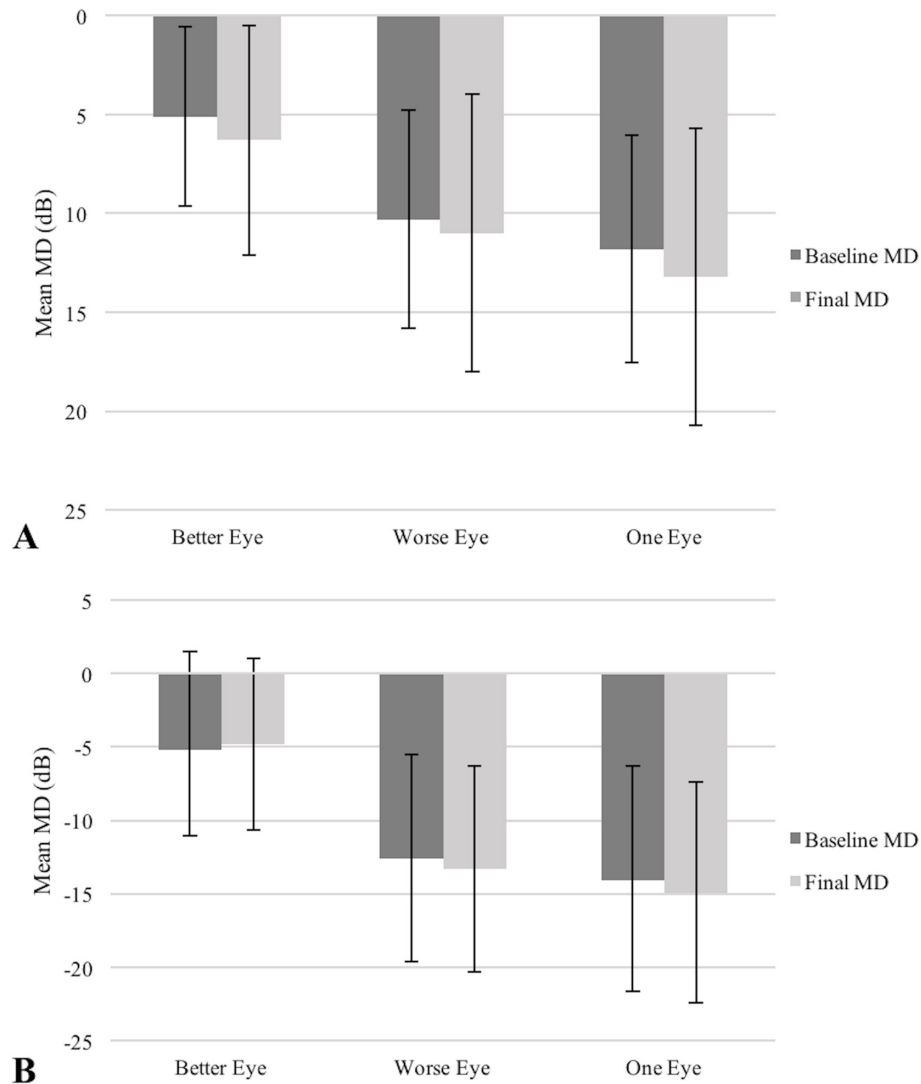


Figure 2 Mean baseline and final MD of eyes included in the study for the Octopus VF (A) and Humphrey VF (B).

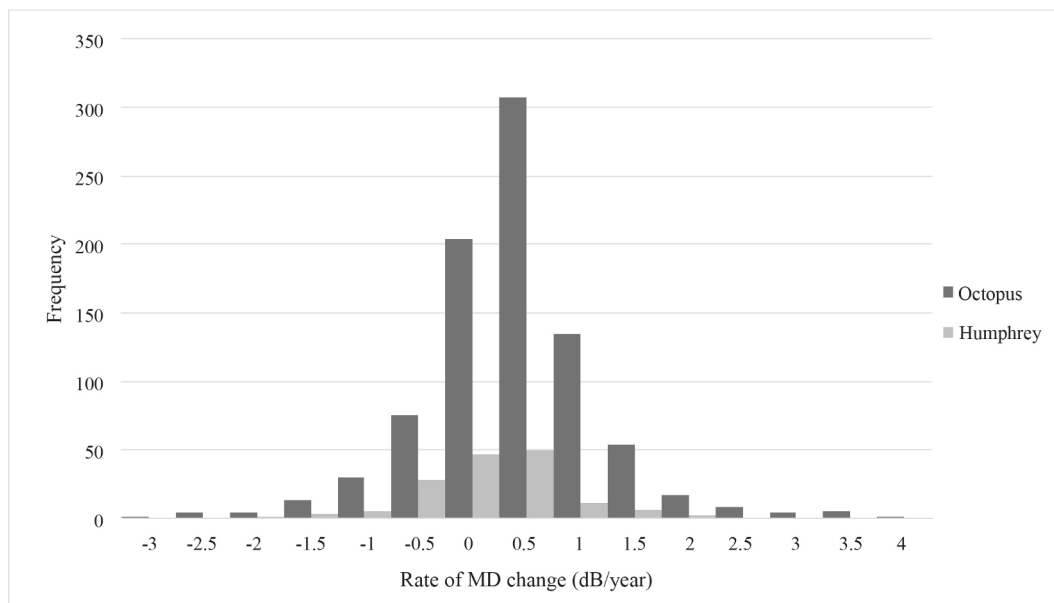


Figure 3 Distribution of the rates of progression of all eyes expressed in dB/year.



Table 3 Univariate analysis of factors associated with a fast rate of progression

Factor	Fast rate of progression MD, n (%)	P value	ORs (95% CI)
Sex			
Male	43 (13.7)	0.135	1.37 (0.91 to 2.08)
Female	60 (9.1)		
Age group			
Less than 60 years old	38 (9.3)	0.455	0.85 (0.56 to 1.30)
60 years old and above	65 (10.8)		
Baseline MD		0.001	0.91 (0.86 to 0.94)
Presence of glaucomatous visual field defect			
Bilateral	55 (11.4)	0.232	1.28 (0.85 to 1.93)
Unilateral	48 (9.1)		

MD, mean defect or deviation.

revealed a slight increase to 12.4% (95% CI 9.68 to 15.02) and a further increase to 15.8% (95% CI 12.82 to 18.74) for the 'worst-case scenario'.

The factors associated with blindness at expected death using univariate and multivariate analysis are displayed in tables 6 and 7, respectively.

Majority of patients who were predicted to reach blindness were male, less than 60 years of age, had a glaucomatous pattern of VF defect on both eyes, and had a progression rate of >1 dB/year for the Octopus and <-1 dB/year for the Humphrey VF. The mean initial MD among these patients was 8.2±4.2 (Octopus). Less than 60 years of age, baseline MD, presence glaucomatous VF defect pattern on both eyes and a fast rate of progression were all statistically significant factors associated with blindness at death. For every 1 dB worsening in baseline MD, the odds of becoming blind at death increased by about 10% holding age, presence glaucomatous VF defect pattern on both eyes and the MD rate of progression constant.

DISCUSSION

This study revealed an overall worsening of the MD in the final VF examination (except for the better eyes tested using the Humphrey VF). The difference in the mean final MD from the initial mean baseline MD, however, was not significant (except for the better eyes tested in the Octopus VF) and was not high (up to less than 1.5 dB). Given the mean follow-up period of around 5 years, this would indicate a progression rate of only about 0.3 dB/year (Octopus) or -0.3 dB/year (Humphrey). Using the individual progression rates and residual life

expectancy, almost 10% were predicted to become blind in their expected lifetime. Depending on the reasons for testing only one eye, this number can reach up to 15.8% ('worst-case scenario'). Although these results appear optimistic, they are less positive compared with the study of Saunders *et al* where in cohorts in UK clinics only 5.2% were shown to progress to blindness.¹³ Also, considering that the progression rate used in their study was -1.5 dB/year compared with -1.0 dB/year in our study for worst-case scenario, our results still showed a higher number of patients who will go blind. For better comparison, we patterned the criteria for blindness in our study with their study (MD worse than or equal to 22 dB). Patients included in their study, however, were being monitored at specialist glaucoma clinics, whereas the patients in our study were not treated exclusively by glaucoma specialists. Furthermore, blindness in our study could have been underestimated since the basis of the diagnosis of glaucoma was simply based on VF and several subjects may have not had glaucoma in the first place since structural correlation is also very important in making the diagnosis of glaucoma. This study was only based on VF follow-ups with no data on other clinical aspects limiting the veracity of the true clinical diagnosis of glaucoma. In addition, other factors (IOP control, type of glaucoma, compliance to medications) which were not included in the study may also have contributed to this difference. Our institute is a tertiary hospital with an eye institute comprising general ophthalmologists and ophthalmology subspecialists. It

Table 4 Multivariate analysis of factors associated with a fast rate of progression

	ORs	95% CI	P value
Sex (M:F)	1.56	1.1 to 2.41	0.041
Baseline MD	0.94	0.83 to 0.91	<0.001

F, female; M, male; MD, mean defect or deviation.

Table 5 Proportion of patients likely to become blind in the course of their lifetime

Blindness at death	% Blind (95% CI)
Patients with series in both eyes, n=433	9.9 (7.11 to 12.75)
All patients best-case scenario, n=583	12.4 (9.68 to 15.02)
All patients worst-case scenario, n=583	15.8 (12.82 to 18.74)

Table 6 Univariate analysis of factors associated with blindness at expected death

Factor	Expected blind at death MD, n (%)	P value	ORs (95% CI)
Sex			
Male	17 (12.4)	0.25	1.46 (0.76 to 2.79)
Female	26 (8.8)		
Age group			
Less than 60 years old	24 (13.8)	0.030	2.00 (1.06 to 3.78)
60 years old and above	19 (7.4)		
Baseline MD		<0.001	1.14 (1.19 to 1.31)
Presence of glaucomatous visual field defect			
Bilateral	38 (16.5)	<0.001	7.76 (2.99 to 20.13)
Unilateral	5 (2.5)		
MD rate of progression			
Fast	21 (65.6)	<0.001	32.72 (14.03 to 76.29)
≤ 1 dB/year (Octopus), ≥ -1 dB/year (Humphrey)	22 (5.5)		

MD, mean defect or deviation.

also has a laser and diagnostic centre where different procedures are done and where ophthalmologists or glaucoma specialists from different hospitals can refer their private patients to. The source of our data was from the eye diagnostic centre where we did not have access to the clinical records of the patients who underwent the VF tests.

The rate of progression of the worse eye in this study with an average follow-up of 5 years using the Humphrey VF was -0.3 dB/year, slightly worse than the 5-year study of Verma *et al*, which was -0.12 ± -0.51 dB/year also using the Humphrey; however, they used a software (Progressor) converting Humphrey sensitivities to pointwise trend analysis.¹⁴ We used a global index specifically MD trend analysis. Also, their patients included only patients with primary angle closure glaucoma, while this study's patients had no specific glaucoma diagnosis and may not even have had any glaucoma at all so direct comparisons on rates of progression may not be possible.

Baseline MD was found to be positively associated with blindness at death. The mean baseline MD among patients who were predicted to reach blindness in this study was 8.2 ± 4.2 (Octopus). This is consistent with the study of Saunders *et al* which demonstrated that most patients at risk of blindness had greater VF defect (MD

worse than -6 dB in Humphrey) at baseline.¹³ A person with a worse MD will be at a higher risk for blindness since he or she will be much closer to the 'blindness threshold' compared with a person with a better MD provided that all other influencing factors are equal. This supports the importance of early disease detection.

Several treatment trials have demonstrated the relationship between initial MD and progression rates. In the Ocular Hypertension Treatment Study and Early Management Glaucoma Trial, a worse baseline VF MD led to increased progression rates compared with those with a better MD.^{4 15} In contrast, a study by Forchheimer *et al* found out that baseline VF MD had no effect on the rate of VF change.¹⁶ This study, on the other hand, indicated a negative association between baseline MD and fast rate of progression. A similar outcome was also shown in a study by Heijl *et al* which revealed that worse baseline VF status was associated with a slower rate of progression.⁶ One reason may be the 'floor' effect experienced by severely damaged eyes. That is, a VF with a severe defect cannot progress as much as a field with less damage.^{6 13}

Fast rate of progression (>1 dB/year for the Octopus and <-1 dB/year for the Humphrey VF), another significant factor for blindness, was seen in more than 60% of patients that were predicted to become blind at death. A similar rate of VF loss was observed in the study by Rossetti *et al*, where all eyes that developed blindness worsened at a rate of -1.1 dB/year (Humphrey).¹⁷ Thus, progression rates should be incorporated in glaucoma care to determine whether treatment should be adjusted in order to prevent blindness without overlooking the possibility of 'floor effect' among severely damaged eyes.

Less than 60 years of age was shown to be a significant factor for blindness at death. Patients in this age group have greater residual life expectancy, therefore, are likely to become blind assuming that no treatment changes are

Table 7 Multivariate analysis of factors associated with blindness at expected death

	ORs	95% CI	P value
Age group	3.72	1.58 to 8.77	0.003
Bilaterality	9.83	2.77 to 34.82	<0.001
Baseline MD	1.10	1.14 to 1.34	0.022
Rate of progression	59.72	19.29 to 184.86	<0.001

MD, mean defect or deviation.

made. This finding is in contrast to the study of Rossetti *et al* which revealed that older age was a risk factor for blindness.¹⁷ A possible explanation for this difference is the fact that blindness was only a predicted outcome in our study and it is presumed that no escalation of treatment is made so that progression is allowed to continue at the same rate until the patient's residual life expectancy. Thus, assuming all other factors are held constant, the total amount of VF decay over a younger patient's lifetime will be greater than that of an older person.

Male sex was found to be a significant factor for a fast rate of progression after controlling for the baseline MD. Although studies on animal models have found evidence on the neuroprotective effects of female sex hormones, human studies have revealed inconsistent results.¹⁸ Other possible explanations could be a poorer IOP control and compliance, a more aggressive type of glaucoma or the presence of another ocular disease in men.

This study displayed visual loss per year, on average. However, the distribution of the rate of change varied and showed that most eyes only progressed at a rate of 0 to 0.5 dB/year. Interestingly, approximately 39.3% of eyes showed improvement in the VF. Saunders *et al* reported positive MD rates in 33% of eyes in their study; however, this could be attributed to VF measurement variability.¹³ A study by Russell *et al* showed that measurement variability increases as the sensitivity decreases to a level of -10 dB and peaks at around -20 dB.¹⁹ Improvement could also be due to learning effects. Our study tried to minimise learning effects by eliminating the first VF examination but the Collaborative Initial Glaucoma Treatment Study showed that improvement in the VF was still possible after the succeeding tests.²⁰ The effect or treatment of a concomitant ocular disease could also be another reason for improvement on the VF. This, however, was not determined in this study but may be helpful in future studies.

This study had certain limitations. Because of its retrospective design, some data were not available or reviewed. Some eyes with glaucomatous VF damage may not actually have had glaucoma, however, the glaucomatous VF criteria used in this study as well as regular VF monitoring for 3 years makes this unlikely. Procedures done within the study period may have influenced the VF, such as sensitivity improvement with cataract extraction. Other ocular disease/procedures which may have contributed to worsening of the VF were not taken into consideration. However, compared with prospective studies which may have an influence on a patient's adherence to follow-up and treatment, ours represents routine clinical setting which will be of more help to determine whether current efforts to slow down progression are adequate. Furthermore, while other studies have determined blindness based on the final MD, this study predicted blindness based on an individual's residual life expectancy instead. This provides a reflection of how a person's VF will progress if no change in treatment takes place and it underscores the importance on individualised treatment. Another recommendation would perhaps also include

visual impairment as an endpoint (-14dB cut-off) for better comparison with other studies.

In conclusion, this study revealed an overall worsening of the VF MD in the majority of eyes at 0–0.5 dB/year. This study also showed that most patients under a routine clinical setting will not reach blindness in their lifetime. Baseline MD, less than 60 years of age, presence of glaucomatous VF defect on both eyes and a fast rate of progression (>1 dB/year for Octopus and <1 dB/year for Humphrey) were shown to be significant factors for blindness. This study also demonstrated that the rates of progression of eyes had considerable variability, and while the male sex was a significant factor for fast progression, the initial MD exhibited an inverse relationship. Because of the association of age and baseline MD on blindness, the role of early diagnosis is critical in blindness prevention. Rate of progression should also be determined especially among those with less severe VF damage.

Contributors JAJT: responsible for overall content, concept, polishing. CCNC: conduct of the study, manuscript writing, statistics, polishing. JMDL: submission, polishing.

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Patient consent for publication Not required.

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Data availability statement All data relevant to the study are included in the article.

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