Clinical profile of adults with permanent visual impairment presenting to the Government tertiary care hospital In India

Praveen Venkatesha Sastry1, Pradeep Addagadde Venkataramana2, Srinivas Siddegowda3,

Affliations-

1Assistant Professor, Department of Ophthalmology, Mandya Institute of Medical Sciences, Mandya, Karnataka, India.

2Professor and HOD, Department of Ophthalmology, Mandya Institute of Medical Sciences, Mandya, Karnataka, India.

3Associate Professor, Department of Ophthalmology, Mandya Institute of Medical Sciences, Mandya, Karnataka, India.

Corresponding Author:

Praveen Venkatesha Sastry, Email:drvpsastry@gmail.com

Abstract

Purpose: To evaluate the causes of visual impairment in adults attending our Government tertiary care hospital, in India for the purpose of blindness disability certification to aid in better planning and prevention of blindness activities in India.

Methods: The study was a non-interventional cross-sectional record-based analysis on 411 adults with permanent visual impairment attending Ophthalmologyout patient department of Mandya Institute of Medical Sciences, Mandya, India for blindness disability certification during July 2019 to Dec 2020. Patients demographic data, educational background, need for disability certification and area of residence all noted after complete ocular examination and categorised depending on the etiology and percentage of visual impairment. Groups analyses and P<0.05 was taken as level of statistical significance.

Results: Of the 411 subjects, 65.2% were males and the majority >53% were in the 18-39 years age group. Overall, most frequent cause of permanent visual impairment was congenital anomalies (17.7%) and retinitis pigmentosa (17.03%). However, age group based data showed a trauma (59.09%) was the most common cause for 30% visual impairment (one eyed). Most common causes for 'low vision'

and 'blindness' in the study group was amblyopia (19.01%) and Congenital

anomalies (29.9%) respectively.

Conclusion: Congenital anomalies and retinitis pigmentosa were common cause of

visual impairment in our study, hence proper genetic counselling to expectant

mothers can mitigate these. Early & timely regular school screening activities by

authorities can help control the incidence of amblyopia and high refractive errors.

Policymakers should at regular intervals form consensus based on disability data to

better manage and empower the blind and mitigate the prevalence of ocular

morbidity.

Keywords: Blindness, disability, congenital anomalies, retinitis pigmentosa

INTRODUCTION

Visual deficits due to any cause can hamper social, psychological and emotional

wellbeing of an

individual. According to the National Sample

Organization(NSSO) approximately 12 million people in India, currently have some

form of blindness. Census data of India 2001, shows the prevalence of totally blind

due to permanent blindness in India to be 156 per lakh population, and those

suffering from low vision to be 61 per lakh population.²

Visual morbidity has tremendous consequences on the individual and the society in

general, especially in a developing economy like ours. 1,3 In-order to mitigate this.

visual disability certification is an important social and economic insurance measure

taken up by the Ministry of Social Justice and Empowerment, to provide

rehabilitation and vocational opportunities for the visually impaired in India. Visual

disability certificates are issued to only those who have a disability percentage of

≥40% as defined by the Gazette of India.4

461

In India, identification of the causes of visual morbidity is of paramount importance to assist in planning of national and social blindness prevention schemes, however most of the data accrued is from the voluntary registration of the blind and blindness registers of various institutes which issue blindness disability certificates for the purposes of employment, education and economic support. Due to the voluntary registration process of the blind, there is under-reporting especially in rural parts of India leading to missed cases and also data regarding the cause of visual impairment in patients with less than 40% visual handicap is missed out from our blindness registers which impacts the planning and prevention of blindness activities in the state and the country. A,5

Studies evaluating the clinical profile of patients suffering from permanent visual impairment are far and few in the state of Karnataka, India. Hence in our current study we aim to assess and evaluate the causes of visual impairment in patients attending our institution for the purpose of blindness disability certification to aid in better planning and prevention of blindness activities in India.

METHODS

The current analysis was a non-interventional, cross-sectional recordbasedstudy on 411 patients who attended Ophthalmology out-patient department (OPD) of Government tertiary care institute-Mandya Institute of Medical Sciences, Mandya, India, for the purpose of Blindness Disability Certification from July 2019 to Dec 2020. The study was conducted after due clearance from the institutional ethical committee clearance. The study conformed to all the tenets of the declaration of Helsinki.Purposive sampling method was used and 411 subjects were included as study subjects. All adults/subjects aged above 18 years, having permanent visual impairment, visiting the OPD of our institute for the purpose of disability certification

were included in the study. However, those with avoidable/treatable causes of decreased vision at presentation such as cataract, correctable refractive errors and records with incomplete data were excluded from the study.

A complete history, demographic data, educational status, residenceand cause for visual impairment was noted. Patientsdistant visual acuity after best possible correction in eye/eyes recorded. Complete ocular examination by slit lamp biomicroscopy, fundus examination by +90D and indirect ophthalmoscopy was done if required and observations noted. Findings of special investigations like B scan, Visual Evoked Potential, Optical Coherence Tomography, Visual fields if needed were also noted. Definitions of Visual impairment, Low vision and Blindness were set according to set criteria of Gazette of India as described below.^{4,7}

Visual impairment was defined according to the criteria set by the Gazette of India, as distance visual acuity less than 6/18 or visual field loss by predefined standardized conservative criteria in the better eye. "Low-vision" defined as visual acuity not exceeding 6/18 or upto 3/60 or in the better eye with best possible corrections or limitation of the field of vision subtending an angle of less than 40 degree up to 10 degree. "Blindness" means a condition where a person has visual acuity less than 3/60 in the better eye with best possible correction or limitation of the field of vision subtending an angle of less than 10 degree. ^{4,7}

Percentage of visual impairment was calculated, based on current guidelines for the evaluation of various disabilities and the procedure for certificationas described in table 1. 4,7

Table 1. Current guidelines for the evaluation of visual disabilities and the procedure for certification

Corrected Best corrected Impairment	Better eye Best	Worse eye	Percent	Disability category
6/24 to 6/60	Corrected	Best corrected	Impairment	
Less than 6/60 to 3/60	6/6 to 6/18	6/6 to 6/18	0%	0
Less than 3/60 No Light Perception 6/24 to 6/60 Or 6/24 to 6/60 40% III a (low vision) Visual field less than 6/60 to 3/60 50% III b (low vision) Less than 3/60 to No 60% III c (low vision) Less than 3/60 to No 60% III c (low vision) Less than 6/60 to 3/60 70% III d (low vision) Less than 6/60 to 3/60 70% III d (low vision) Less than 6/60 to 3/60 70% III d (low vision) Less than 3/60 to No 80% III e (low vision) Less than 3/60 to No 80% III e (low vision) Less than 3/60 to No 80% III e (low vision) Less than 3/60 to No 80% III e (low vision) Less than 3/60 to No 90% IV a (Blindness) Only HMCF Only HMCF Only Light Perception No Light Only Light Perception No Light		6/24 to 6/60	10%	0
Perception Percep		Less than 6/60 to 3/60	20%	1
6/24 to 6/60 Or Visual field less than 40 up to 20 degree around centre of fixation or hemianopia involving macula Less than 6/60 to 3/60 Less than 3/60 to No Light Perception Less than 6/60 to 3/60 Less than 6/60 to 3/60 Less than 6/60 to No Light Perception Less than 6/60 to No Light Perception Less than 6/60 to No Light Perception Less than 3/60 to No Light Perception IV a (Blindness) Only HMCF Only Light Perception No Light Perception No Light Perception No Light Perception		Less than 3/60 No Light	30%	II (One eyed
Visual field less than 40 up to 20 degree around centre of fixation or hemianopia involving macula Less than 6/60 to 3/60 Less than 6/60 to 3/60 Less than 6/60 to No 60% Light Perception Less than 6/60 to 3/60 To% Ill c (low vision) Ill d (low vision) Less than 6/60 to 3/60 To% Ill d (low vision) Ill e (low vision) Ill e (low vision) Less than 20 up to 10 degree around centre of fixation Less than 3/60 to No 90% IV a (Blindness) IV b (Blindness) Only HMCF Only HMCF Only Light Perception No Light No Light		Perception		person)
than 40 up to 20 degree around centre of fixation or hemianopia involving macula Less than 6/60 to 3/60 70% III d (low vision) Jess than 6/60 to 3/60 70% III d (low vision) Less than 6/60 to 3/60 70% III d (low vision) Less than 20 up to 10 degree around centre of fixation Less than 3/60 to 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only Light Perception Only Light Only Light Perception IV b (Blindness) Only Light Perception No Light	6/24 to 6/60 Or	6/24 to 6/60	40%	III a (low vision)
degree around centre of fixation or hemianopia involving macula Less than 6/60 to 3/60 70% III d (low vision) 3/60 Or Visual filed less than 3/60 to No Light Perception Less than 3/60 to Less than 3/60 to No 1/60 Or Visual field less than 3/60 to 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only Light Perception Only Light Perception Light Perception IV a (Blindness) IV b (Blindness) Only Light Perception No Light Perception	Visual field less	Less than 6/60 to 3/60	50%	III b (low vision)
centre of fixation or hemianopia involving macula Less than 6/60 to Less than 6/60 to 3/60 70% III d (low vision) 3/60 Or Visual filed Less than 3/60 to No 80% III e (low vision) less than 20 up to 10 degree around centre of fixation Less than 3/60 to No 90% IV a (Blindness) 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only HMCF Only Light Perception Only Light Perception No Light Only Light Perception No Light	than 40 up to 20	Less than 3/60 to No	60%	III c (low vision)
or hemianopia involving macula Less than 6/60 to 3/60 70% III d (low vision) 3/60 Or Visual filed less than 20 up to 10 degree around centre of fixation Less than 3/60 to No 10	degree around	Light Perception		
Less than 6/60 to 3/60 70% III d (low vision) 3/60 Or Visual filed less than 3/60 to No less than 20 up to 10 degree around centre of fixation Less than 3/60 to No less than 10 degree around centre of fixation Only HMCF Only HMCF Only Light Perception Only Light Perception No Light No Light	centre of fixation			
Less than 6/60 to 3/60 T0% III d (low vision) 3/60 Or Visual filed less than 3/60 to No 10 B0% III e (low vision) Less than 20 up to 10 Light Perception Less than 3/60 to No 10 B0% IV a (Blindness) Less than 3/60 to No 10 B0% IV a (Blindness) Less than 10 degree around centre of fixation Only HMCF Only Light Perception Only Light Perception No Light No Light	or hemianopia			
3/60 Or Visual filed Less than 3/60 to No 80% Light Perception Less than 3/60 to No 90% Less than 3/60 to No 90% Less than 3/60 to No 90% Light Perception Less than 3/60 to No 90% Light Perception IV a (Blindness) Only HMCF Only HMCF Only Light Perception Only Light Perception No Light Perception No Light Perception	involving macula			
less than 20 up to 10 degree around centre of fixation Less than 3/60 to Less than 3/60 to No 90% IV a (Blindness) 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only HMCF Only Light Perception OnlyLight Perception No Light No Light	Less than 6/60 to	Less than 6/60 to 3/60	70%	III d (low vision)
10 degree around centre of fixation Less than 3/60 to Less than 3/60 to No 90% IV a (Blindness) 1/60 Or Visual field Light Perception less than 10 degree around centre of fixation Only HMCF Only HMCF 100% IV b (Blindness) OnlyLight Perception Perception No Light Perception No Light	3/60 Or Visual filed	Less than 3/60 to No	80%	III e (low vision)
Centre of fixation Less than 3/60 to Less than 3/60 to No 90% IV a (Blindness) 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only HMCF Only Light Perception OnlyLight Only Light Perception Perception No Light No Light	less than 20 up to	Light Perception		
Less than 3/60 to Less than 3/60 to No 1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only Light Perception OnlyLight Only Light Perception Perception No Light No Light IV a (Blindness) IV b (Blindness)	10 degree around			
1/60 Or Visual field less than 10 degree around centre of fixation Only HMCF Only HMCF 100% IV b (Blindness) OnlyLight Only Light Perception Perception No Light	centre of fixation			
less than 10 degree around centre of fixation Only HMCF Only HMCF 100% IV b (Blindness) OnlyLight Only Light Perception Perception No Light No Light	Less than 3/60 to	Less than 3/60 to No	90%	IV a (Blindness)
degree around centre of fixation Only HMCF Only HMCF 100% IV b (Blindness) OnlyLight Only Light Perception Perception No Light No Light	1/60 Or Visual field	Light Perception		
Centre of fixation Only HMCF Only HMCF Only Light Perception Perception No Light No Light Only Light Perception No Light No Light	less than 10			
Only HMCF Only HMCF 100% IV b (Blindness) OnlyLight Only Light Perception Perception No Light No Light	degree around			
OnlyLight Only Light Perception Perception No Light No Light	centre of fixation			
Perception No Light Perception No Light	Only HMCF	Only HMCF	100%	IV b (Blindness)
No Light	OnlyLight	Only Light Perception		
	Perception	No Light Perception		
Perception	No Light			
	Perception			

CF- Finger counting, HMCF- Hand movement close to face

Statistical analysis

Analysis was done using descriptive statistics in the form of percentages, mean and standard deviation. Continuous variables were analysed using 'one sample goodness-of- fit test'. All the recorded data were statistically analysed by IBM SPSS Statistics version 21 (IBM Corp., Armonk, N.Y., USA). For statistical significance, *p* value <0.05 was considered.

RESULTS

Of the 411 subjects who were part of the study, 268 were males and 143 females. The age of the study population ranged from 18 to 70 years, with a mean of 36.31±15.86 years. Majority of the study population were in the age group of 18-39 years accounting for 53.28% of cases. More than 55.76% of the population who came to avail the services of disability certification were from the rural areas, and there was a direct statistical relevance to the educational status and the ocular disability. Approximately 60.32% of the subjects who came to avail the certification had not received any formal school education or dropped out during early childhood and belonged to a lower economic strata as classified under the Kuppuswamy classification.⁹

In our current study the purpose of availing disability certification was highest for reasons of financial benefits, followed by travel allowance, educational benefits and medical issues respectively. However, the most common age group who availed the services were among the 18-39 age group, followed by 40-59 years age group as depicted in figure 1.

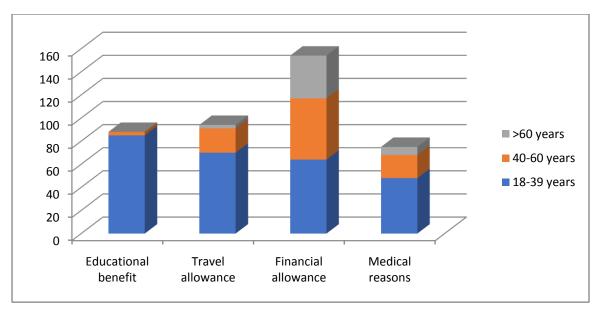


Figure 1: Reasons for availing disability certification in all age groups

Out of the 411 study subjects, 204 (49.63%) were categorized as 'blind' (with more than 80% visual impairment), 163(39.65%) were categorised as 'low vision' (40-80% visual impairment) and 44 (10.7%) were 'one eyed' (30% visual impairment). Overall, the most common cause for visual impairment noticed in our study was congenital anomalies (17.7%), followed by retinitis pigmentosa (17.03%). Retinitis pigmentosa was the most common cause of visual impairment in the age group of 18-39 years (p value <0.05), however in the 40 to 59 years age group, congenital anomalies were the most common. The values were statistically significant as observed in table 2 with p<0.05.

Table 2: Age wise distribution of the aetiology of visually impairment in our study

Etiology	Age group (years)			Total n (%)
	18-39	40-59	≥60	
Congenital anomalies	37	28	8	73 (17.7%)
Retinitis pigmentosa	42	24	4	70 (17.03%)
Amblyopia	31	12	0	43 (10.4%)
Macular pathology	25	8	2	35 (8.5%)

Trauma	25	5	3	33 (8.02%)
Pathological myopia	20	12	0	32 (7.7%)
Glaucoma	2	20	8	30 (7.29%)
Optic atrophy	20	8	0	28 (6.81%)
Corneal scar	6	11	6	23 (5.59%)
Diabetic retinopathy	4	8	6	18 (4.37%)
Complicated	0	7	6	13 (3.16%)
pseudophakia				
Retinal detachment	6	4	0	10 (2.43%)
Chronic uveitis	1	2	0	3 (0.72%)
Total	219	149	43	411
	(53.28%)	(36.25%)	(10.46%)	
P value	0.00*	0.00*	0.50	0.00*

^{*} p<0.05 statistically significant

On further analysis of the aetiology of blindness among 'blind', 'low vision' and 'one eyed' in various age groups the most common cause for 30% visual impairment (one eyed) was trauma (59.09%)(p<0.05)* which was more commonly seen in the age group of 18-39 years (45.4%) and the majority of them were of male gender accounting for 14 patients. In the age group of 40-59 years and above 60 years, the common cause for being one eyed or 30% visual impairment was, complicated cataract surgery/pseudophakia and glaucoma accounting for 15.9% and 11.36% cases respectively as depicted in figure 2.

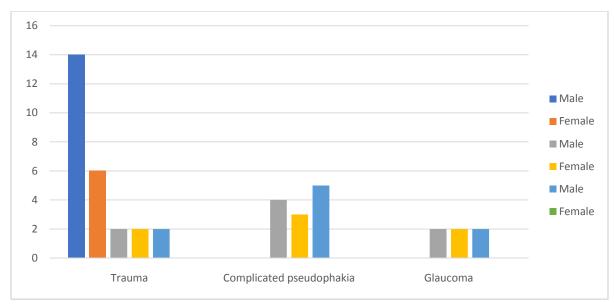


Figure 2: Age and sex distribution for causes of 'one eyed' (30% visual impairment)

Most common causes for 'low vision' (40% to 80% visual impairment) in the study group was amblyopia (19.01%)(p<0.05), followed by pathological myopia (17.17%) (p<0.05) and macular pathology (13.49%). Most of the study subjects having 'low vision' were in the age group of 18-39 years as observed in figure 3.

Out of the 204 participants who were categorized as 'blind' in our study, Congenital anomalies accounted for 29.9%(p<0.05) of patients, followed by retinitis pigmentosa observed among 25.49%(p<0.05) subjects across all the age groups. A majority of the patients classified as 'blind' or had visual impairment more than 80% were found in the 18-39 year age group as depicted in figure 4.

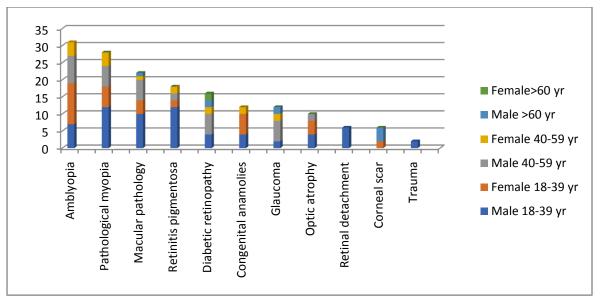


Figure 3: Age and sex distribution for causes of 'low vision' (40-80% visual impairment)

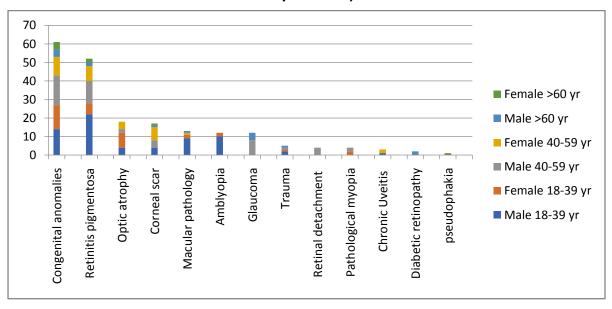


Figure 4: Age and sex distribution for causes of 'blind' (> 80% visual impairment)

DISCUSSION

Blindness Disability certification is a social and economic insurance measure for the underprivileged visually impaired of our society. Blindness registers, National blindness surveys and voluntary registration of the blind, although are efficient measures to assess the patterns and aetiology of visual impairment in the society, a

lot of data is missed especially of those patients with visual impairment less that 40 percent, as per the criteria set by the Ministry of social justice and empowerment. Are According to the data accrued from our study, the majority of the subjects seeking blindness disability certification were males between the age group of 18 to 60 years for financial & employment allowance, travel allowance and for availing educational benefits. The study findings were in line with the analysis conducted in 2013 in Maharashtra by *Joshi et al* wherein the reasons for availing this social scheme was primarily seen in males for education, employment and financial reasons in the same order. Similarly, the study findings of a higher proportion of males seeking disability certification were observed in *Ghosh et al* and *Kareemsab et al* 2, probably due to the social obstacles for females and long held beliefs of males as bread winners and more accessibility of social services and employment opportunities in our society still prevalent in our country.

Overall, the most common cause for visual impairment in our analysis was congenital anomalies followed by retinitis pigmentosa. Our observations were consistent with the study findings of *Kareemsab et al*¹² wherein congenital anomalies (22.05%), refractive errors (19.85%), and retinitis pigmentosa (18.01%) were common similar to that of the *Titiyal et al* ¹³ study and *Ambastha et al*¹⁴ study who opined that congenital anomalies due to macular scar was the most prevalent cause of visual disability. In a similar study by *Ghosh et al* ¹¹, phthisis bulbi followed by congenital anomalies like microphthalmos was the most frequent occurrence. Our study findings contrasted to that of the western studies wherein ARMD (age related macular degeneration) was the commonest cause of visual disability as observed in the *Avisar et al* ¹⁵ and *Bunce et al* ¹⁶ study.

The most frequent cause of visual impairment less than 40 percent (one eyed) who were not eligible for disability certification was ocular trauma especially among young males 18-39 years, probably due to outdoor activities for the purpose of employment. However, our study findings were not in line with that of *Ambastha et al* ¹⁴ study in Bihar, who opined that the most common cause of visual impairment of less than 40 percent was amblyopia. The high ocular morbidity observed in our study due to trauma can be mitigated by imparting occupational safety and education standards to all workers at their work space.

In patients with 'low vision' (40% to 80% visual handicap), amblyopia, pathological myopia and macular pathology were the commonest causes in our study in the 18 to 39 year age group. Amblyopia and pathological myopia leading to visual handicap can be easily tackled by maintaining timely quality school screening activities and providing spectacle and low vision aid services in educational institutes similar to as suggested by the *Ambastha et*al ¹⁴ and Ghosh et al ¹¹study.

In those with > 80 percent visual disability or 'blind', who approached to avail disability certification was due to congenital anomalies and retinitis pigmentosa in the early age group, whereas in the beyond 60year age group complicated pseudophakia & glaucoma was the most common cause in our population, whereas Bunce et al ¹⁶ study opined that ARMD was the commonest cause of visual disability in the western population. However Indian studies like Ghosh et al ¹¹ and Titiyal et al ¹³back our study results.

LIMITATIONS OF THE STUDY: A larger sample size which included all the institutions in our region would provide a better representative sample than the one available in our study.

European Journal of Molecular & Clinical Medicine ISSN 2515-8260 Volume10, Issue 03, 2023

CONCLUSION

Congenital anomalies and retinitis pigmentosa were the most common cause of

permanent visual impairmentin our study, hence proper genetic counselling to

expectant mothers and education regarding eye care in the neonatal age group can

mitigate the incidence of increased ocular morbidity. Early & timely regular school

screening activities by authorities can help control the incidence of amblyopia and

high refractive errors causing low vision in the younger age group to improve their

social, economic and emotional wellbeing. Improve the standards of cataract

surgeries by providing good training facilities and infrastructure for ophthalmologists

to minimise the chances of complicated pseudophakia. Policymakers should set

occupational safety standards and create safe working conditions at all work

stations. Authorities need to relook at the causes of ocular morbidity at regular

intervals and form consensus to better manage and empower the blind and mitigate

the prevalence of ocular morbidity in the country.

Conflict of interest- Nil

Financial interest - Nil

REFERENCES

1. Demmin DL, Silverstein SM. Visual Impairment and Mental Health: Unmet

Needs and Treatment Options. Clin Ophthalmol. 2020;3;14:4229-4251.

2. National Sample Survey Organization, Ministry of Statistics and Programme

Implementation, Government of India, Round Number 37th in 1981, 47th in 1991 and

58th in 2002.

3. Marmamula S, Barrenakala NR, Challa R, Kumbham TR, Modepalli SB,

Yellapragada R, Bhakki M, Khanna RC, Friedman DS. Prevalence and risk factors

for visual impairment among elderly residents in 'homes for the aged'in India: the

472

Hyderabad Ocular Morbidity in Elderly Study (HOMES). BrJOphthalmol. 2021; 1;105(1):32-6.

- 4. Ministry of Social Justice and Empowerment. Guidelines for evaluation of various disabilities and procedure for certification. Notification dated 1st June, 2001. The Gazette of India extraordinary. Part 1. Section 1. No 154.
- 5. Bandyopadhyay S, Bandyopadhyay SK, Biswas J, Saha Dutta Chowdhury M, Dey AK, Chakrabarti A. Visual Impairment Registry of Patients from North Kolkata, Eastern India: A Hospital-based Study. J Ophthalmic Vis Res. 2018; 13(1):50-54.
- 6. Black K. Business statistics: for contemporary decision making. John Wiley & Sons; 2019;12.
- 7. Guidelines for the evaluation of various disabilities and procedures for certification. The Gazette of India extraordinary Part II; Section 3: sub section (ii) No 61.
- 8. Mehta CR, Patel NR. IBM SPSS exact tests. Armonk, NY: IBM Corporation. 2011;17:23-4.
- 9. Shaikh Z, Pathak R. Revised Kuppuswamy and BG Prasad socio-economic scales for 2016. Int J Community Med Public Health. 2017; 4(4):997-9.
- 10. Joshi RS. Causes of visual handicap amongst patients attending outpatient department of a medical college for visual handicap certification in central India. J Clin OphthalmolRes. 2013;1;1(1):17.
- 11. Ghosh S, Mukhopadhyay S, Sarkar K, Bandyopadhyay M, Maji D, Bhaduri G. Evaluation of registered visually disabled individuals in a district of West Bengal, India. Indian journal of community medicine: official publication of Indian Association of Preventive & Social Medicine. 2008;33(3):168.
- 12. Kareemsab D, Rachaiah NM. Balasubramanya. Prevalence of leading causes of certification for blindness and partial sight. J Clin Diagn. 2011;5(8):1624-6.
- 13. Titiyal JS, Pal N, Murthy GV, Gupta SK, Tandon R, Vajpayee RB, Gilbert CE. Causes and temporal trends of blindness and severe visual impairment in children in schools for the blind in North India. Br J Ophthalmol. 2003;1;87(8):941-5.

- 14. Ambastha A, Kusumesh R, Sinha S, Sinha BP, Bhasker G. Causes of visual impairment in applications for blindness certificates in a tertiary center of Bihar and its role in health planning. Indian J Ophthalmol. 2019;67(2):204.
- 15. Avisar R, Friling R, Snir M, Avisar I, Weinberger D. Estimation of prevalence and incidence rates and causes of blindness in Israel, 1998-2003. IMAJ-RAMAT GAN-. 2006;1;8(12):880.
- 16. Bunce C, Evans J, Fraser S, Wormald R. BD8 certification of visually impaired people. BrJ Ophthalmol. 1998;1;82(1):72-6.