Pattern and Presentation of Vitreo-Retinal Diseases: An Analysis of Retrospective Data at a Tertiary Eye Care Center in Nepal

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Purpose: We examined patients presenting in a tertiary eye hospital in Nepal, focusing on information relevant to screening and management programs for vitreo-retinal (VR) disease.

Design: Retrospective, cross-sectional study.

Methods: We reviewed all patients presenting for the first time to the VR-clinic over 1 year. We quantified patient demography, symptoms and duration, systemic diseases, ophthalmological examinations, diagnostic investigations, and final diagnoses.

Results: Of the 1905 cases, 1148 were male (60.3%). The 25th percentile of ages was 29 and 38 years for male and female, respectively; thus, female presented later (P < 0.0001). Hypertension was the commonest systemic disease (40.8%), followed by diabetes (32.5%). Age-related macular degeneration (AMD) and diabetic retinopathy (DR) affected 447 eyes (11.8%) and 46 eyes (10.9%), respectively. Male and female AMD and DR patients did not differ in age or disease duration. Similarly, age or disease duration for DR did not correlate with severity. Asymmetry of disease severity between eyes with AMD and DR was largest in patients with 1 normal eye. Presenting acuity was asymmetric between eyes (P < 0.0001) with people more often reporting once their right eyes had acuity of 6/18 or worse.

Conclusions: The screening of blood pressure and glucose levels combined with fundus photography could prevent many from progressing to life-changing visual impairment and blindness. Later reporting by females began at childbearing age; therefore, education and ocular screening could be useful in reproductive health programs. Clubbing VR disease screening with other established health programs like diabetes control program, hypertension clinics, school health program, and so on, would provide economical and sustainable approach.

Key Words: AMD, diabetic retinopathy, retinal diseases, laterality, sex differences

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METHODS

Setting

This study was conducted at the Tilganga Institute of Ophthalmology (TIO), Kathmandu, Nepal. The TIO is a tertiary eye care center in Nepal, focusing on information relevant to screening and management programs for vitreo-retinal (VR) disease.
eye care center with subspeciality clinics providing services for ophthalmological patients from all over Nepal, India, and Bhutan. All subspeciality clinics in the TIO, including the VR clinic, run from morning till afternoon, and an extended paying clinic operates in the evening. Despite this availability of extended facilities, 1843 cases (96.7%) presented to the routine public VR clinic. Only 36 cases (1.9%) presented to the paying clinic, whereas 26 cases (1.4%) reported to the emergency department.

Study Design and Ethics
The study has been approved by the TIO-Institutional Review Committee (TIO-IRC) vide letter number: Ref: 10/2018. The need for consent was waived by TIO-IRC because the retrograde study collected de-identified data from patient files.

Study Population
This study covers all cases presenting to the VR clinic for the first time over 1 year. Patients who presented for repeat or follow-up visits were not included. The study ran from January 1, 2010 to December 31, 2010. A key objective of this study is to provide a baseline reference for planned follow-up studies that will use similar analyses. A further benefit of recording this hospital-based study is that there are population-based studies of disease prevalence from both Nepal4,20 and Bhutan5 from 2009/2010 with which this study can be compared, adding extra value for both the region and similar countries.

Data Collection and Analysis
Detailed demographic information, presenting complaints, treatment history, and associated systemic diseases were noted. Best corrected visual acuity (BCVA), refractive status, Goldmann applanation tonometry, findings of slit-lamp examination, 90D biomicroscopy, fundoscopy, and diagnostic investigations performed were recorded. Diagnoses were noted separately for right and left eyes. Data collection was supervised by the head of the VR unit to ensure high quality and uniform practice. Hypertensive retinopathy was graded according to Keith-Wagener-Barker classification system,21 and DR was classified as per the modified Airlie House and Abbreviated Early Treatment Diabetic Retinopathy Study (ETDRS) classification of DR.22

The data were analyzed using SPSS (version 20.0, IBM, New York, NY) and MATLAB (2016b, The MathWorks, Natick, MA). Comparisons of the expected and observed frequency of sex or eye-wise effects were done using Chi-squared tests. To compare the 10th, 25th, 50th, 75th, and 90th percentiles of the distributions of Figure 1B for male and female, we applied a sampling-with-replacement-based bootstrap analysis23 to estimate the population means and the standard errors (SE) of the means of those percentiles. We employed 10,000 bootstrap cycles to insure the estimated means and SE converged to within 2 decimal places on 5 independent cross-validations. We then applied t tests employing the estimated means and SE.

RESULTS
We will first give an overview of the general presentation of the 1905 cases. We will then present the ocular disease data as: the nonretinal diseases involving posterior segment other than retina, and the main analysis of the retinal diseases. Background data are presented in 5 Supplementary tables, http://links.lww.com/APJO/A56.
General Presentation

The 1905 new cases presenting to the VR clinic were split as: 1148 male (60.3%) and 757 female (39.7%). Demographic data are shown in Table 1. Figure 1A gives a breakdown of the presenting ages in 20-year cohorts for each sex. Figure 1B shows that, for the whole cohort. The 25th percentile was 29 years for male and 38 years for female. A bootstrap analysis revealed that the 10th, 25th, 50th and 75th percentiles were all significantly older for female than male by 3.95, 9.28, 7.25, and 3.08 years (ie, median is 5.6 years). The 10th percentile difference was significant at $P < 0.003$, and the others at $P = 0.0001$. It is noted that the 25th percentiles for the males were lower in the 20-year cohorts of Figure 1A as well.

A total of 802 cases (42.1%) were from the Kathmandu valley, 1038 cases (54.5%) were from Nepal outside the Kathmandu valley, and 65 cases (3.4%) were from other countries. A total of 1574 cases (82.6%) reported to the hospital as primary cases, whereas 331 cases (17.4%) were referred from other hospitals or health care centers.

The commonest presenting complaint was poor vision accounting for 1142 cases (59.9%), followed by trauma in 227 (11.9%), loss of vision in 155 (8.1%), floaters in 91 (4.8%), and vision not improving after cataract surgery in 44 (2.3%). All presenting complaints are summarized in Table 2. A total of 565 cases (30.5%) presented within a month of the onset of their symptoms, whereas 500 (27.0%) presented only after 12 months. In 53 cases, we could not affirm the time (Supplemental Digital Content 1, Table S1, http://links.lww.com/APJO/A56).

Hypertension was the commonest systemic disease associated in 250 cases (40.8%), followed by diabetes in 199 cases (32.5%) and combined diabetes and hypertension in 124 cases (20.2%) (Fig. 2). The other systemic diseases found in 40 cases (6.5%) are summarized in Supplemental Digital Content 1, http://links.lww.com/APJO/A56. The duration of systemic disease association was found to be 10 years in 24.0%.

We quantified interventions done at other centers before patients reported to our clinic, including surgical, laser, or ophthalmic injections. These cases included cataract surgery in 55 (2.9%), VR surgery in 27 (1.5%), retinal laser in 27 (1.5%), and primary repair of corneoscleral tear or eyelid lacerations in 20 (2.9%). All previous interventions are summarized in Table 2. A total of 584 cases (23.1%) had undergone any intervention.

Hematological testing was the most commonly diagnostic test performed in 584 cases (23.1%), followed by optical coherence tomography in 516 (20.4%), A-scan in 436 (17.2%), fundus fluorescence angiography in 89 (3.5%), and visual fields in 39 (1.5%). Diagnostic investigation was not indicated or not done in 687 cases (27.1%). Supplemental Digital Content 1, Table S4, http://links.lww.com/APJO/A56 summarizes the diagnostic procedures.

### Table 2. Presenting Symptoms

| Presenting Complaints                | Frequency | %  |
|--------------------------------------|-----------|----|
| Poor vision                          | 1142      | 59.9 |
| Trauma                               | 227       | 11.9 |
| Loss of vision                       | 155       | 8.1  |
| Floaters                             | 91        | 4.8  |
| Vision not improving after cataract surgery | 44    | 2.3  |
| Sudden loss of vision                | 43        | 2.3  |
| Routine eye check-up                 | 41        | 2.2  |
| Night blindness                      | 34        | 1.8  |
| Foreign body sensation               | 20        | 1.0  |
| Pain                                 | 18        | 0.9  |
| Flashes of light                     | 8         | 0.4  |
| Black spot in visual field           | 8         | 0.4  |
| White eye                            | 5         | 0.3  |
| Others                               | 69        | 3.6  |
| Total                                | 1905      | 100  |

BCVA was 6/18 or better in the right eye in 660 cases (34.8%), and in left eyes in 1025 cases (54.1%). In 751 cases (39.4%) right eyes, and 355 cases (18.6%) left eyes, the BCVA was 6/18 to 6/60. Figure 3 illustrates how surprisingly asymmetric these values were ($P < 0.0001$, see legend). These findings prompted us to explore our data to see whether any other such asymmetries occurred. Cases with BCVA <6/60 or no perception of light (NPL) were more common on the left, but not significantly so. NPL was found in 37 (1.8%) right eyes and 44 (2.3%) left eyes.

### Table 1. Demographic Characteristics of Patients

| Parameters                | Female Patients | Male Patients |
|---------------------------|-----------------|--------------|
| No. patients              | 757             | 1148         |
| Mean age, y               | 52.3            | 47.1         |
| Median age, y             | 57              | 50           |
| Standard deviation        | 20.5            | 22.2         |
| Minimum age, y            | 2.5             | 0.17         |
| Maximum age, y            | 90              | 116          |

Nonretinal Diseases

Before detailing the VR diseases, we quickly summarize the other issues discovered. Posterior vitreous detachment was the commonest nonretinal disease with 79 cases (35 OS). There were 51 cases of endophthalmitis (24 OS): 33 (64.7%) were post-traumatic, 13 (25.5%) postoperative, and 5 (9.8%) endogenous in origin. The other diseases involving posterior segment other than the retina are enumerated in Supplemental Digital Content 1, Table S5, http://links.lww.com/APJO/A56.

**Figure 2.** Systemic disease prevalence. The breakdown of the commonest systemic diseases found in 613 (32.2%) of the 1905 patients. The values at the top of the columns give the percentage of the systemic disease patients (left), and the percentage of all patients (right). DM indicates diabetes mellitus; HT, hypertension.
Most of the trauma-related conditions involved the left eyes, but this bias did not reach statistical significance. When pooled together, atrophic bulbi, phthisis, and absolute eyes were more common in left eyes: 16 (0.8%) compared with only 8 right eyes (0.4%, \( P = 0.05 \), chi-square). Dislocated cataract was similar in left eyes: 14 (0.8%) compared with 10 right eyes (0.5%, \( P = 0.25 \), chi-square). Dislocated intraocular lens into the vitreous cavity was also seen in 7 left eyes (0.4%) and 3 right eyes (0.2%, \( P = 0.10 \), chi-square). Retained intraocular foreign body involved 7 left and 4 right eyes (\( P = 0.15 \), chi-square). Open- and close-globe injuries affected both eyes equally. The fundus was not visible in 6 left eyes (0.3%), and 7 right eyes (0.4%).

**Retinal Diseases**

The commonest retinal disease was AMD, affecting 223 right eyes (11.7%) and 224 left eyes (11.8%), followed by DR involving 208 left, and 208 right eyes (10.9%). RD was the third commonest, affecting 112 right eyes (5.9%) and 126 left eyes (6.6%), of which 71% were rhegmatogenous, 17.2% exudative, and 11.8% tractional RD. There were 150 cases of branch RVO (BRVO), 80 in right and 70 in left eyes. Myopic degeneration affected 63 right and 54 left eyes, and retinitis pigmentosa (RP) was seen in 56 right and 57 left eyes. Other retinal diseases diagnosed were hypertensive retinopathy in total of 87 eyes, central serous chorioretinopathy in 77 eyes, central RVO in 65 eyes, macular scar in 53 eyes, macular hole in 55 eyes, retinal vasculitis in 46 eyes, and others detailed in Table 3. 94.8% of full-thickness macular hole and 84.6% lamellar macular hole were unilateral. The fundus was found to be normal in 623 left eyes (32.7%) and 593 right eyes (31.1%). The types of AMD and the severity of DR are shown in Table 4.

Among the AMD patients, dry AMD was the commonest type (79.2%), followed by wet AMD (15.9%), retinal pigment epithelium (RPE) change (4%), and familial drusen (0.9%). In contrast to the results of all patients (Fig. 1), there was no statistical difference in the number of male (122) and female (100) AMD patients. The male and female AMD patients also did not differ significantly in age: male 71.0 ± 9.48 years, female 68.8 ± 10.3 years (mean ± SD years). Among these patients, 81.7% of the AMD was bilateral and 18.3% unilateral. As might be expected, patients with unilateral disease were significantly younger than those with bilateral disease: 67.0 ± 13.1 cf. 70.7 ± 8.93 years (\( P = 0.031 \), t test).

In part because of the higher rate of smoking by male in Nepal (32.8% among male cf. 15.8% among female), we decided to examine wet AMD eyes relative to fellow eyes. Two patients had familial bilateral drusen, and for this analysis they were classed as having dry AMD. Four patients had nonvisible fundus (mainly due to cataract) and were eliminated from the analysis. The age of male whose worst eye had wet AMD was not different from female: 66.0 ± 15.3 versus 65.8 ± 13.7 years. There were 40 males whose worst eye had wet AMD, versus 15 females, which was marginally significant (\( P = 0.063 \), t test, correcting for the relative abundance of males in the study population). Interestingly, patients whose worst eye had dry AMD were older than those with a worst eye having wet AMD [66.0 ± 15.3 versus 65.8 ± 13.7 years (\( P = 0.045 \)). Overall, there was a suggestion that males develop wet AMD earlier, and in relatively greater numbers compared with females. This might have been an effect of smoking, but further investigation is needed.

We also examined severity and laterality by scoring normal fundus to wet AMD on a scale of 1 to 3 (normal, dry, wet) and then...
examining the absolute value of the difference in scores between eyes. 6 patients had bilateral wet AMD OU. For the 186 patients whose best eye had dry AMD, only 30 had a worst eye with wet AMD. Patients whose best eye was normal were even more heterogeneous. Of those 36 patients, 17 had a fellow eye that was dry, and 19 had a fellow eye that was wet. Thus, the pattern of progression seemed to follow relative heterogeneity earlier, progressing to more bilateral disease later, rather than simple bilateral disease at each stage.

Among DR eyes, moderate nonproliferative DR (NPDR) was the commonest (34.6%), followed by severe NPDR (26.2%), proliferative diabetic retinopathy (PDR) (25.2%), and mild NPDR (13.9%). A total of 230 eyes had clinically significant macular edema (CSME): 50.9% involving left eyes and 49.1% right eyes. Like the AMD patients, the DR group contained more males (123) than females (91), but this was not significant. Their ages also did not differ at 58.7 ± 8.99 years for males, and 58.3 ± 11.21 years for females. Their durations of diabetes did not differ at 11.5 ± 6.79 and 10.4 ± 6.96 years, respectively. 9 patients had a nonvisible fundus in 1 eye and were removed from further analysis. We examined the laterality of DR in the remaining 205 patients.

We scored the 5 DR diagnostic-categories/severities from normal fundus to PDR as 1 to 5. As with AMD, the pattern of progression seemed to follow relative heterogeneity earlier, progressing to more bilateral disease later. As with AMD, we then binned subjects according to the diagnosis in their least affected eye. To quantify the degree of laterality, we took the absolute value of the difference of these severity steps in each pair of eyes. A box plot of the results is shown in Figure 4A. Basically, as with AMD, eyes tended to become more similar as severity of the best

### Table 3. Diseases Involving Retina

| Diagnosis                        | N     | % Across Retinal Diseases | % Across All Diseases |
|----------------------------------|-------|---------------------------|-----------------------|
| AMD                              | 223   | 13.5                      | 11.7                  |
| Diabetic retinopathy             | 208   | 12.6                      | 10.9                  |
| Retinal detachment               | 112   | 6.8                       | 5.9                   |
| BRVO                             | 80    | 4.8                       | 4.2                   |
| Myopic degeneration              | 63    | 3.8                       | 3.3                   |
| Retinitis pigmentosa             | 56    | 3.4                       | 2.9                   |
| Hypertensive retinopathy         | 45    | 2.7                       | 2.4                   |
| CSCR                             | 44    | 2.7                       | 2.3                   |
| CRVO                             | 33    | 2.0                       | 1.7                   |
| Macular scar                     | 31    | 1.9                       | 1.6                   |
| Macular hole                     | 30    | 1.8                       | 1.6                   |
| Retinal vasculitis               | 23    | 1.4                       | 1.2                   |
| Chorio-retinal scar              | 22    | 1.3                       | 1.2                   |
| Epiretinal membrane              | 20    | 1.2                       | 1.0                   |
| Macular dystrophy                | 16    | 1.0                       | 0.8                   |
| Irido-chorio-retinal coloboma    | 15    | 0.9                       | 0.8                   |
| Retinal break                    | 13    | 0.8                       | 0.7                   |
| Status post RD surgery           | 9     | 0.5                       | 0.5                   |
| Central retinal artery occlusion | 8     | 0.5                       | 0.4                   |
| Stargardt disease                | 5     | 0.3                       | 0.3                   |
| Retinal hemorrhage               | 4     | 0.2                       | 0.2                   |
| Cone dystrophy                   | 3     | 0.2                       | 0.2                   |
| Normal fundus                    | 593   | 35.8                      | 31.1                  |
| Total                            | 1656  | 100.1                     | 86.9                  |

AMD indicates macular degeneration; BRVO, branch retinal vein occlusion; CRVO, central retinal vein occlusion; CSCR, central serous chorioretinopathy; RD, retinal detachment.

### Table 4. Types of AMD and Severity of DR

| Diagnosis     | Right Eye | Left Eye | Total |
|---------------|-----------|----------|-------|
| AMD           | N         | %        | N     | %     |
| Dry           | 175       | 78.5     | 179   | 79.9  | 354   | 79.2  |
| Wet           | 37        | 16.6     | 34    | 15.2  | 71    | 15.9  |
| RPE change    | 9         | 4.0      | 9     | 4.0   | 18    | 4.0   |
| Familial drusen | 2    | 0.9      | 2     | 0.9   | 4     | 0.9   |
| Total         | 223       | 100.0    | 224   | 100.0 | 447   | 100.0 |
| DR            |           |          |       |       |       |       |
| Mild NPDR     | 33        | 15.9     | 25    | 12.4  | 58    | 13.9  |
| Moderate NPDR | 68        | 32.7     | 76    | 36.5  | 144   | 34.6  |
| Severe NPDR   | 57        | 27.4     | 52    | 26.4  | 109   | 26.2  |
| PDR           | 50        | 24.0     | 55    | 26.4  | 105   | 25.2  |
| Total         | 208       | 100.0    | 208   | 100.0 | 416   | 100.0 |

AMD indicates macular degeneration; DR, diabetic retinopathy; NPDR, nonproliferative diabetic retinopathy; RPE, retinal pigment epithelium; PDR, proliferative diabetic retinopathy.
eye increased. As shown in Figure 4B, severity of DR was not correlated with duration of disease.

**DISCUSSION**

The visual handicap experienced by individuals suffering from unilateral eye disease, such as macular hole, is reported to be strongly influenced by ocular dominance. Ocular-dominance and handedness are associated, with about 65% of right-handers, and 43% of left-handers, being right eye-dominant. Some of our data (Fig. 3) suggested that eye dominance played a role in patients reporting to the hospital, with the predominantly right-handed subjects only reporting that their right eye had BCVA worse than 6/18 once. This type of asymmetry has been reported before. AMD and DR were very homogenous with respect to age and sex; however, when the best eye was normal, the range of severity in the fellow eye was surprisingly broad (eg, Fig. 4). Taking together the results might mean that if their nondominant eye is affected, then patients tended not to notice their sight threatening disease, indicating an excellent argument for more screening. Given the subjects only reporting (Fig. 3), we subsequently analyzed other data by eye (Tables 3 and 4, and Supplemental Digital Content 1, Table S4, http://links.lww.com/APJO/A56) to elucidate any eye-wise biases; however, few were found.
The preponderance of males in the total group was 60.3%, and for subgroup of AMD and DR patients combined it was 56.5%, marginally more than for females (P = 0.052). This agrees with the findings of some hospital-based studies on VR diseases, but differs from a Nepalese population-based study from the same era, in which only 45.5% were male, and 1 in India with only 45% male. Our results could be due to being reserved due to social norms and therefore not coming forward for medical check-ups. Here the 25th percentile of ages reporting was nearly 10 years higher for female (P < 0.0001, t test 9.27 years; 95% confidence level (CL) 6.05 and 12.5 years). The mean and median ages of male and female (49 years and 54 years, respectively) tally well with other studies of VR diseases. Reported ages ranged from 2 months to 116 years.

Only 41 cases (2.2%) presented for routine check-up. Chronic retinal diseases like AMD, DR, RD, RVO, RP, and so on, ranked high in the diagnostic list, but only 2.2% of cases were found to present for routine check-up, indicating that there is a need to emphasize patient education and counseling about the importance of regular check-ups and follow-ups. A total of 565 cases (30.5%) presented within a month, 507 cases (27.1%) between 1 to 6 months, 280 cases (15.1%) between 6 to 12 months, and 500 cases (27.0%) presented after 12 months from the onset of the symptoms. The delayed presentation time might be because of illiteracy and ignorance (the overall literacy rate of Nepal for population aged 5 years and above was 65.9% in 2011, 75.1% for male and 57.4% for female). Remote areas and associated poor accessibility to modern medical services encourage people initially to rely on traditional methods of healing.

Systemic disease association was not found in 1292 of the 1905 cases. As shown in Fig. 2 (and Supplemental Digital Content 1, Table S2, http://links.lww.com/APJO/A56), among the 613 remaining cases, the commonest associated diseases were hypertension (40.8%), diabetes (32.5%), and both (20.2%). This agrees with the Tehran eye study which reported hypertension (21.14%), followed by diabetes (15.99%). By contrast, a Nigerian study found diabetes to be more common (14.6%) than hypertension (13.2%). Other systemic diseases were found in 40 of our cases (6.5%, Supplemental Digital Content 1, Table S2, http://links.lww.com/APJO/A56). The duration of systemic disease association was <5 years in 210 cases (42.8%), 5 to 10 years in 163 cases (32.2%), and >10 years in 118 cases (24.0%).

The commonest VR disease was AMD affecting 11.8%, followed by DR at 10.9%; matching another Nepalese study which reported that AMD was the commonest VR disease at 28.3%, followed by DR at 17.9%. This does not match with a Nigerian hospital study, which reported DR (24.9%) as the commonest VR disease, followed by hypertensive retinopathy (13.3%) and AMD (10.7%). In our case, 208 of 324 (64.2%) confirmed diabetic cases had some form of DR, which did not tally with population-based studies (10.5%). In our study, 117 left eyes and 113 right eyes of 323 patients had CSME but a population-based study in Nepal found it in only 2 of 305 diabetic cases. This disparity could be explained by the early onset of macular edema causing a high percentage of CSME patients presenting to the TIO.

RD was the third commonest VR disease in our study, affecting 112 right eyes (5.9%) and 126 left eyes (6.6%), whereas a Nepalese population-based study reported population prevalence of 0.10%. A hospital-based study in Ethiopia reported RD as the second commonest VR disease at 24.5%. Of 65 cases of full-thickness macular hole in our study, 33 cases (50.8%) affected right eyes, whereas others have reported 48% of right eyes affected. In our study, 94.8% of the full-thickness macular hole and 84.6% of the lamellar macular hole were unilateral, which are similarly reported in other studies, and are basically the consequence of factors affecting the macula locally. When such unilateral macular diseases affect the dominant eyes of the individuals, they cause greater functional visual impairment.

In eastern Asia, pathological myopia is a major issue found in 80% to 90% of school-leavers, and 10% to 20% of those completing secondary school, and contributes to RD numbers. The prevalence of myopia varies from 0.8% to 53.4%, depending on geographical area, age, occupation, and ethnicity. In Nepal, Sherpa children had a prevalence of 2.9% as compared with 21.7% for Tibetan children. A myriad of myopic complications like atrophic retinal holes and RD, choroidal neovascular membranes, degeneration, cataract, and glaucoma, cause visual loss warranting attention. Thapa et al reported the population prevalence of macular hole as 0.20% in Nepal. Asymptomatic macular holes occur at a prevalence of 6.26% among high myopes with >−2 diopters.

**Limitations**

19 patients were excluded due to incomplete data. AMD was not classified as per the Age-Related Eye Disease Study (AREDS) classification system. Only the final diagnoses, but not ocular comorbidities, have been considered for analysis, which might have altered the reported disease patterns. The ocular dominance and handedness were not recorded in the case records of the study population, but based on the literature reviewed and cited above, we speculated that if only the nondominant eye was affected, this might cause the patients to present late. In future studies it would be valuable to record the dominant eye. Similarly, the reviewed medical records of the diabetic patients under study did not differentiate the type of diabetes, which we will consider in follow-up studies.

**CONCLUSIONS**

We found that without screening programs, patients often fail to notice developing visual impairment until the disease progresses to advanced stage, especially in their nondominant eye. In Nepal, and perhaps in similar countries, females report later for care than men. That was true from childbearing age, so education and screening could be usefully coupled for reproductive health programs. This study indicates that low-cost screening and management programs for retinal disease could be of immense value in developing countries. The high prevalence of hypertension and diabetes among retinal disease patients suggest that a simple screening of blood pressure and glucose levels combined with fundus photography could prevent many from progressing to life-changing visual impairment and blindness. We recommend that coordinating with other established health programs would provide economical and sustainable screening methods such as diabetes control program for screening DR, hypertension clinics for screening hypertensive retinopathy, reproductive health program for VR diseases affecting female of childbearing age, comprehensive school health program for pathological myopia and hereditary diseases among school children, and counseling.
and rehabilitation centers for counseling people affected with permanent visual impairment or blindness, and other hereditary diseases.

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