Clinical Research

Prognosis value of Chinese Ocular Fundus Diseases Society classification for proliferative diabetic retinopathy on postoperative visual acuity after pars plana vitrectomy in type 2 diabetes

Tie-Zhu Lin^{1,2}, Yan Kong², Cheng Shi², Emmanuel Eric Pazo², Guang-Zheng Dai², Xian-Wei Wu², Ling Xu², Li-Jun Shen^{1,3}

¹Eye Hospital and School of Ophthalmology and Optometry, Wenzhou Medical University, Wenzhou 325000, Zhejiang Province, China

²He Eye Specialist Hospital, Shenyang 110034, Liaoning Province, China

³Ophthalmology Department, Zhejiang Provincial People's Hospital, Hangzhou 310000, Zhejiang Province, China

Correspondence to: Li-Jun Shen. Eye Hospital and School of Ophthalmology and Optometry, Wenzhou Medical University, Wenzhou 325000, Zhejiang Province, China. slj@mail.eye.ac.cn Received: 2021-10-22 Accepted: 2022-07-05

Abstract

• **AIM**: To compare the postoperative visual acuity among eyes with proliferative diabetic retinopathy (PDR) of different stages after pars plana vitrectomy (PPV) in type 2 diabetic patients.

• **METHODS:** A retrospective study was conducted for PDR eyes undergoing PPV in type 2 diabetic patients. All patients were divided into three groups based on Chinese Ocular Fundus Diseases Society (COFDS) classification for PDR: Group A (primary vitreous hemorrhage), Group B (primary fibrovascular proliferation) and Group C (primary vitreous hemorrhage and/or fibrovascular proliferative combined with retinal detachment). The postoperative visual acuity and the change between postoperative and preoperative visual acuity were compared among three groups. The associated risk factors for postoperative visual acuity were analyzed in the univariate and multiple linear aggression.

• **RESULTS:** In total, 195 eyes of 195 patients were collected in this study, including 71 eyes of 71 patients in Group A, 75 eyes of 75 patients in Group B and 49 eyes of 49 patients in Group C. The eyes in Group A got better postoperative best-corrected visual acuity (BCVA) compared to the eyes in Group B and C (0.48 ± 0.48 vs 0.89 ± 0.63 , *P*<0.001; 0.48 ± 0.48 vs 1.04 ± 0.67 , *P*<0.001; respectively). The eyes in Group A got more improvement of

BCVA compared to the eyes in Group B and C (1.07 ± 0.70 vs 0.73 ± 0.68 , P=0.004; 1.07 ± 0.70 vs 0.77 ± 0.78 , P=0.024; respectively). In the multiple linear regression analysis, primary fibro-proliferative type ($\beta=0.194$, 95%CI=0.060-0.447, P=0.01), retinal detachment type ($\beta=0.244$, 95%CI=0.132-0.579, P=0.02), baseline logMAR BCVA ($\beta=0.192$, 95%CI=0.068-0.345, P=0.004), silicone oil tamponade ($\beta=0.272$, 95%CI=0.173-0.528, P<0.001) was positively correlated with postoperative logMAR BCVA. Eyes undergoing phacovitrectomy had better postoperative BCVA ($\beta=-0.144$, 95%CI=-0.389 to -0.027, P=0.025).

• **CONCLUSION:** PDR eyes of primary vitreous hemorrhage type usually have better visual acuity prognosis compared to primary fibrovascular proliferation type and retinal detachment type. COFDS classification for PDR may have a high prognostic value for postoperative visual outcome and surgical management indications.

• **KEYWORDS:** proliferative diabetic retinopathy; pars plana vitrectomy; postoperative visual acuity; best-corrected visual acuity; Chinese Ocular Fundus Diseases Society

DOI:10.18240/ijo.2022.10.10

Citation: Lin TZ, Kong Y, Shi C, Eric Pazo E, Dai GZ, Wu XW, Xu L, Shen LJ. Prognosis value of Chinese Ocular Fundus Diseases Society classification for proliferative diabetic retinopathy on postoperative visual acuity after pars plana vitrectomy in type 2 diabetes. *Int J Ophthalmol* 2022;15(10):1627-1633

INTRODUCTION

D iabetic retinopathy (DR) is a common cause of vision loss and blindness. The global prevalence of retinopathy is estimated to be 22.27% among all diabetics^[1]. The advanced form of DR, proliferative diabetic retinopathy (PDR), is a sight-threatening illness^[2]. The global prevalence of PDR is estimated to be approximately 1.4% among all individuals with diabetes^[3]. In a Meta-analysis, the prevalence of PDR was 6% in Asian type 2 diabetic patients, accounting for 17%

DR patients^[4]. Vitreous hemorrhage, severe fibrovascular proliferation, combined tractional retinal detachment (TRD), and/or rhegmatogenous retinal detachment (RRD) are the primary signs of PDR, which are also considered as indication criteria for pars plana vitrectomy (PPV). The surgical goals include reducing bleeding-related vitreous opacities, alleviating vitreoretinal tension by removing the posterior hyaloid and epiretinal membranes, and using endo-photocoagulation to achieve proliferative tissue regression.

For the first cases of PDR, the Early Treatment Diabetic Retinopathy Study (ETDRS) severity scale classified the illness as mild, moderate, or high-risk PDR. Advanced PDR was defined as the presence of traction, retinal detachment (RD), rubeosis iridis, or fundus obscuration^[5]. Because of its complexity, most retinal surgeons managing diabetic patients do not use the whole ETDRS severity scale, according to several surveys^[6].

The number of diabetics in China is by far the highest. A common use classification for PDR in China based on retinal pathology and morphology was made by Chinese Ocular Fundus Diseases Society (COFDS) in 1985, and was kept in the modified version in 2014, including the early PDR stage (neovascular elsewhere or of the disk, with or without vitreous or preretinal hemorrhage), the fibrous proliferation stage, and the advanced PDR stage (TRD)^[7]. Though this classification has fewer details compared to ETDRS grading, it is more convenient for retinal surgeons in clinical practice. Based on this classification, we would like to know the prognosis of PDR after PPV surgery. In this study, we compared the postoperative visual acuity among PDR eyes of different stages in type 2 diabetic patients.

SUBJECTS AND METHODS

Ethical Approval The study design followed the tenets of the Declaration of Helsinki and was approved by the Institutional Review Board at He Eye Specialist Hospital [IRB (2020) K006.01]. Written informed consent was waived due to retrospective design.

Study Design and Participants This retrospective study was performed for PDR eyes undergoing PPV in patients with type 2 diabetes in He Eye Specialist Hospital between January 2017 and June 2020.

The criteria of inclusion were: 1) PDR eyes undergoing 23G PPV or phacovitrectomy treatment; 2) patients with type 2 diabetes mellitus; 3) more than 18 years old; 4) Snellen best-corrected visual acuity (BCVA) at baseline over light perception; and 5) the minimal follow-up time was 6mo. Exclusion as follows: 1) ocular pathology other than cataract and PDR that may affect visual acuity, such as pterygium, corneal opacity, glaucoma, age related macular degeneration, amblyopia; 2) history of ocular surgery; 3) cataract of grade 3 or more dense^[8]; 4) postoperative anatomical success was not

1628

achieved; and 5) other systemic diseases that may influence vision acuity, such as cerebral thrombosis, hematencephalon, intracranial tumors.

Data Collection and Grouping Patients' gender, age, HbAlc, duration of diabetes, vitreous tamponade, previous photocoagulation, the use of anti-vascular endothelial growth factor (VEGF) agents from the medical charts were recorded. The records of surgical procedures were also reviewed. All the patients underwent Snellen BCVA at distance, slitlamp examination, intraocular pressure measurement, pupil dilated fundus examination, ultra-field scanning laser ophthalmoscopy (SLO; Optos[®] 200Tx, Optos[®], Dunfermline, UK) and ultrasonographic B-scan preoperatively. All eyes were divided into three groups based on PDR classification of COFDS using preoperative SLO and ultrasonographic B-scan imaging^[5]. Group A (primary vitreous hemorrhage): massive hemorrhage was seen in the vitreous on SLO image without obvious proliferative membrane noted, no vitreous tractional lesion or TRD was seen on ultrasonographic B-scan. Group B (primary fibrovascular proliferation): obvious fibroproliferative membrane on the optic disc or posterior pole area along the vascular arch was seen on SLO image, or obvious fibroproliferative traction was noted on ultrasonographic B-scan, without TRD or RRD. Vitreous hemorrhage was permitted. Group C (TRD/RRD type): TRD or RRD noted on SLO and/or ultrasound B-scan (Figure 1). That grouping would be corrected if obvious fibrovascular membrane or TRD was discovered intraoperatively, because of the low sensitivity of SLO and ultrasound B-scan for fibrovascular membrane in the presence of dense vitreous hemorrhage. The consistency analysis would be done between the real retinal condition and the initial classification. The classification was performed by two masked, experienced retina specialists (Wu XW and Kong Y). A third masked retina specialist (Lin TZ) made the final decision in cases where no agreement between the graders was achieved. The visual acuity at 6mo follow up was assigned as postoperative visual acuity for statistical analysis.

Surgical Techniques Under retrobulbar anesthesia, all patients received a comprehensive PPV using 23-gauge instrumentation (OS4TM, Oertli, Switzerland) and the Resight 500 (Carl Zeiss Meditec, Inc., Dublin, California, USA) for visualization. Fibrovascular membrane dissection was done according to each subject's specific demands and requirements. During the procedure, an endolaser was used. Replacement of the vitreous with gas (C_2F_6), fluid, or silicone oil was left to the surgeon's discretion. Phacoemulsification was conducted systematically before the PPV, through a 2.8 mm clean corneal incision, with the implantation of an acrylic foldable IOL in the case of a combined surgery. If eyes received intravitreal injection of anti-VEGF agents [ranibizumab (Lucentis; Genentech,

South San Francisco, CA), aflibercept (Eylea; Regeneron, Tarrytown, NY), or conbercept (Kanghong Biotech Company, Chengdu, Sichuan Province, China)], that was administered intraoperatively or 3-5d before surgery^[9].

Statistical Analysis For data analysis, Snellen visual acuity was transformed to logMAR visual acuity. The logMAR result for count-fingers was 1.9, hand motion was 2.3, light perception was 2.7, and no light perception was $3.0^{[10]}$. The inter-rater agreement for classification of groups was checked using intraclass coefficient (ICC). The agreement between reviewers was high: ICC=0.965 (P<0.001). Mean±SD and percentage (%) were used among the descriptive statistics. Categorical data was presented as percentages and analyzed using the Chi-square test. ANOVA was used to test homogeneity of variance among groups, and based on the homogeneity of variance, an independent t test was used for multiple comparison. Multiple linear regression was used to identify the factors associated with postoperative BCVA. All statistical analyses were carried out using SPSS software version 26.0 (SPSS Inc, Chicago, USA), with an alpha level of 0.05 (type I error).

RESULTS

Patient Characteristics In total, 195 eyes of 195 patients were collected in this study. At first, Group A with 76 eyes, Group B with 72 eyes and Group C with 47 eyes were classified by only using SLO and ultrasound B-scan imaging. There were 71 eyes of 71 patients (35 females and 36 males with a mean age of 54.37±11.30y) in Group A, 75 eyes of 75 patients (37 females and 38 males with a mean age of 50.95±9.00y) in Group B and 49 eyes of 49 patients (27 females and 22 males with a mean age of 47.43±13.32y) in Group C after correction by reviewing surgical records. Four eyes with primary vitreous hemorrhage presented fibrovascular proliferation intraoperatively. RD was found during surgery in 1 eye with primary vitreous hemorrhage and 1 eye with fibrovascular proliferation. The agreement between preoperative and intraoperative DR classification was excellent: ICC=0.981 (P<0.001). The baseline BCVA was comparable among three groups (Table 1). Postoperative Visual Acuity There were significant improvements in mean postoperative visual acuity compared to baseline in all three groups $(0.48\pm0.48 \text{ vs } 1.55\pm0.61, P < 0.001;$ 0.89±0.63 vs 1.62±0.57, P<0.001; 1.04±0.67 vs 1.81±0.57, P < 0.001; respectively, Figure 2). In the comparison among three groups, the eyes in Group A got better mean postoperative visual acuity compared to the eyes in Groups B and C (0.48±0.48 vs 0.89±0.63, P<0.001; 0.48±0.48 vs 1.04±0.67, P < 0.001; respectively), meanwhile the eyes in Group A got more improvement of mean postoperative visual acuity compared to the eyes in Groups B and C (1.07±0.70 vs 0.73 ± 0.68 , P=0.004; 1.07±0.70 vs 0.77±0.78, P=0.024;

 Int J Ophthalmol,
 Vol. 15,
 No. 10,
 Oct.18,
 2022
 www.ijo.cn

 Tel:
 8629-82245172
 8629-82210956
 Email:
 ijopress@163.com



Figure 1 Ultra-field SLO and B-scan images of 3 stages of PDR Primary vitreous hemorrhage type: massive hemorrhage noted in the vitreous on SLO image (A), no vitreous tractional lesion or traction retinal detachment on ultrasound B-scan (B). Primary fibrovascular proliferation type: obvious fibroproliferative membrane on posterior pole area along the vascular arch noted on SLO image (C), without retinal detachment on ultrasound B-scan (D). Retinal detachment type: tractional retinal detachment noted on SLO image (E) and ultrasound B-scan (F). SLO: Scanning laser ophthalmoscopy; PDR: Proliferative diabetic retinopathy.



Figure 2 The preoperative and postoperative BCVA in three groups BCVA: Best-corrected visual acuity.

respectively). The mean improvement of postoperative visual acuity is comparable between Groups B and C ($0.73\pm0.68 vs$ 0.77 ± 0.78 , *P*=0.751). Thirty-seven (52.11%) eyes in Group A, 17 (22.67%) eyes in Group B and 8 (16.33%) eyes in Group C got visual acuity of \geq 20/40 after PPV therapy (*P*<0.001). The

COFDS classification for PDR

Table 1	Demographic	data and surg	gical methods	in	three	groups
---------	-------------	---------------	---------------	----	-------	--------

Items	Group A (<i>n</i> =71)	Group B (<i>n</i> =75)	Group C (<i>n</i> =49)	Р
Glycemic index (mmol/L)	7.39±2.70	7.37±2.67	6.70±2.38	0.294
HbAlc (%)	7.63±1.80	7.63±1.55	6.96±1.49	0.783
Creatinine (µmol/L)	111.71±177.73	124.67±196.12	79.64±39.03	0.582
Age (y)	54.37±11.30	50.95±9.00	47.43±13.32	0.768
Sex, male (%)	36 (50.7)	38 (50.7)	22 (44.9)	0.484
Duration of diabetes (y)	12.64±7.65	10.38±6.73	9.13±6.72	0.073
Baseline BCVA (logMAR)	1.55±7.65	1.62 ± 0.57	1.81±0.57	0.06
Vitreous tamponade				< 0.001
None	17	11	2	
Gas	42	25	15	
Silicone oil	12	39	32	
Previous photocoagulation, n (%)	36 (50.7)	36 (48)	18 (36.7)	0.004
Intravitreal anti-VEGF agents, n (%)				0.001
None	34 (47.9)	26 (34.7)	13 (26.5)	
Preoperatively	19 (26.8)	44 (58.6)	33 (67.4)	
Intraoperatively	18 (25.3)	5 (6.7)	3 (6.1)	
Phacovitrectomy	51 (71.8)	59 (78.7)	35 (71.4)	0.011

BCVA: Best-corrected visual acuity; VEGF: Vascular endothelial growth factor.

Fable 2 Comparison o	f postoperative BCVA	among three groups
-----------------------------	----------------------	--------------------

BCVA	Group A	Group B	Group C	ЪР
Postoperative logMAR BCVA	0.48 ± 0.48	0.89±0.63ª	$1.04{\pm}0.67^{a}$	< 0.001
The change of postoperative logMAR BCVA	-1.07 ± 0.70	-0.73±0.68 ^a	-0.77 ± 0.78^{a}	0.009
Postoperative BCVA 20/40	37/71	17/75	8/49	< 0.001
Postoperative increase of BCVA of \geq 3 lines	55/71	34/75	20/49	< 0.001

BCVA: Best-corrected visual acuity. ${}^{a}P < 0.05 vs$ group A; ${}^{b}P$: ANOVA among three groups.

percentage of eyes with postoperative increase of visual acuity of \geq 3 lines was 77.46%, 45.33% and 40.82% in Groups A, B and C respectively (P<0.001; Table 2).

Postoperative Visual Acuity Associated Factors In the univariate analysis, the duration of diabetes (P=0.073), baseline BCVA (P < 0.001), previous photocoagulation (P = 0.004), classification of PDR (P<0.001), vitreous tamponade (P < 0.001), intravitreal anti-VEGF agents (P < 0.001), phacovitrectomy (P=0.011), and duration of operation (P=0.001) were significantly correlated with postoperative BCVA (Table 3). The variates with P < 0.2 were included in the multiple linear regression analysis, PDR of primary fibrovascular proliferation type (β =0.194, 95%CI=0.060-0.447, P=0.01), RD type ($\beta=0.244$, 95%CI=0.132-0.579, P=0.02), baseline logMAR BCVA (β=0.192, 95%CI=0.068-0.345, P=0.004), silicone oil tamponade ($\beta=0.272$, 95%CI=0.173-0.528, P<0.001) were positively correlated with postoperative logMAR BCVA, while phacovitrectomy was negatively correlated with postoperative logMAR BCVA (β =-0.144, 95%CI=-0.389 to -0.027, P=0.025; Table 4).

DISCUSSION

While the proportion of people with diabetes developing PDR

and severe visual loss decreased between 1980 and 2008 in populations with improved diabetes control^[11], the crude prevalence of visual impairment and blindness caused by DR increased significantly between 1990 and 2015, according to the most recent report of the Vision Loss Expert Group of the Global Burden of Disease Study^[12], owing largely to the increase in diabetes prevalence. PDR is characterized by the hallmark feature of pathologic preretinal neovascularization^[13]. The newly formed vessels could locate in the retina, or extend into vitreous or subretinal space. Structural alterations at the vitreoretinal interface encourage vasogenic cells to migrate and proliferate in the vitreous, resulting in vitreous hemorrhage and macular edema. The vitreous collagen mixes with the developing vascular endothelium to give it contractile properties. Simultaneously, fibrous tissue forming along new veins aligns with the posterior hyaloid, giving the vitreous contractile properties. Contraction of the vitreous can pull developing retinal vessels into the subhyaloid region, causing bleeding. Blood can break into the vitreous tissue concurrently or later if the hemorrhage is severe or the vitreous is substantially liquefied. TRD, secondary RRD, chronic macular edema, premacular hemorrhage, and recurrent

Items	<i>n</i> =195	Postoperative logMAR BCVA (Snellen)	Р
Age (y), mean±SD	51.31±11.31		0.768
Gender, <i>n</i> (%)			0.484
Male	96 (49.2)	0.74±0.63 (0.18)	
Female	99 (50.8)	0.80±0.64 (0.16)	
Duration of diabetes (y), mean±SD	10.90±7.19		0.073
Glycemic index (mmol/L), mean±SD	7.21±2.61		0.233
HbAlc (%), mean±SD	7.46±1.65		0.783
Creatinine (µmol/L), mean±SD	108.70±163.79		0.582
Baseline BCVA, mean±SD	1.64±0.59		< 0.001
Previous photocoagulation, n (%)			0.004
Yes	90 (46.2)	0.64±0.55 (0.23)	
No	105 (53.8)	0.90±0.68 (0.13)	
Classification of PDR, <i>n</i> (%)			< 0.001
Primary VH type	71 (36.4)	0.48±0.48 (0.33)	
Primary fibrovascular proliferation type	75 (38.5)	0.89±0.63 (0.13)	
RD type	49 (25.1)	1.04±0.63 (0.09)	
Vitreous tamponade, n (%)			< 0.001
Fluid	30 (15.4)	0.66±0.63 (0.22)	
Silicone oil	83 (42.6)	1.08±0.61 (0.08)	
C_2F_6	82 (42)	0.51±0.52 (0.31)	
Intravitreal anti-VEGF agents, n (%)			0.001
None	73 (37.4)	0.58±0.53 (0.26)	
Preoperative	96 (49.2)	0.94±0.64 (0.11)	
Intraoperative	26 (13.3)	0.74±0.73 (0.18)	
Phacovitrectomy, <i>n</i> (%)			0.011
Yes (phacovitrectomy+IOL implantation)	145 (75.5)	0.58±0.48 (0.26)	
No (lens saving)	50 (24.5)	0.85±0.67 (0.14)	
Duration of operation(min), mean±SD	61.60±20.68		0.001

BCVA: Best-corrected visual acuity; SD: Standard deviation; PDR: Proliferative diabetic retinopathy; VH: Vitreous hemorrhage; RD: Retinal detachment; VEGF: Vascular endothelial growth factor; IOL: Intraocular lens.

 Table 4 Multiple analysis of linear regression for postoperative

 BCVA

Table 3 Univariate analysis for postoperative BCVA

Items	β	95%CI	Р
Classification of PDR			< 0.001
Primary VH type	-	-	
Primary fibrovascular proliferation type	0.194	(0.060, 0.447)	0.010
RD type	0.244	(0.132, 0.579)	0.002
Baseline logMAR BCVA	0.192	(0.068, 0.345)	0.004
Vitreous tamponade			
Fluid	-	-	
C_2F_6	-0.116	-	
Silicone oil	0.272	(0.173, 0.528)	< 0.001
Phacovitrectomy	-0.144	(-0.389, -0.027)	0.025

BCVA: Best-corrected visual acuity; CI: Confidence interval; PDR: Proliferative diabetic retinopathy; VH: Voireous hemorrhage; RD: Retinal detachment.

vitreous hemorrhage can all be caused by continued vitreous contraction.

Kroll *et al*^[14] established a classification of PDR that clearly distinguished different stages of the disease. Based on the hypothetic pathogenesis of PDR and clinical observations with preoperative examination techniques as well as intraoperative observations, the stages were briefly differentiated as stage A (proliferative changes with retina attachment), stage B (circumscribed TRD with macula attachment) and stage C (TRD in volving the macula), which is primarily based on retinal proliferative changes and macular status. In clinical trials, the severity of a subject's retinopathy over time can be categorized into discrete steps on Diabetic Retinopathy Severity Scale (DRSS). The DRSS is based on grading of fundus stereo photographs of 7 fields, and classifies DR per eye into 13 complex levels (7 levels for PDR). This grade can be influenced by the presence of severe vitreous hemorrhage. The procedure of seven standard mydriatic and stereoscopic 30-degree fundus fields (seven-field ETDRS) is also timeconsuming. Ultra-field SLO is a time-saving technique that can

cover up to 200 degrees of the fundus without mydriasis, and ultrasound B-scan could give the sagittal view of vitreous and retina. Though the dense vitreous hemorrhage shields the retina and could cause some bias in classification, we corrected that grouping using surgical records, and the agreement between preoperative and intraoperative classification was excellent. Only a few eyes with flat fibrovascular membrane or mild traction retinal detachment were incorrectly assigned to group A or B.

For about 40y, PPV has been utilized in PDR. Several papers describe the visual acuity outcomes in PDR eyes following vitrectomy surgery. Ratnarajan *et al*^[15] reported that the mean postoperative visual acuity of PDR eyes undergoing PPV was significantly improved during 44mo of follow up. Additionally, we observed a considerable improvement in postoperative visual acuity in PDR eyes at various stages following PPV surgery. A total of 62 (31.79%) eyes got postoperative visual acuity of $\geq 20/40$, and 109 (55.90%) eyes got over 3 lines improvement of postoperative visual acuity. We classified the eyes of PDR into three groups based on the PDR classification of COFDS: primary vitreous hemorrhage, primary fibrovascular proliferation and RD. We discovered that the eyes with primary vitreous hemorrhage had significantly better postoperative visual acuity than the other two groups. Postoperative visual acuity improvement was also greater in eyes with primary vitreous hemorrhage. Early vitrectomy, according to the Diabetic Retinopathy Study, increased the likelihood of restoring visual acuity greater than 20/40. In this study, more eyes with primary vitreous hemorrhage had postoperative visual acuity of 20/40 or greater than a three-line improvement in comparison to the other two PDR types.

Additionally, in univariate and multiple linear regression analyses, the type of primary vitreous hemorrhage was identified as a positive risk factor for postoperative BCVA. Nishi *et al*^[16] also discovered a correlation between vitreous hemorrhage and improved postoperative visual acuity in the absence of a fibrovascular membrane. Vitreous blood is typically caused by retinal neovascularization in PDR eyes. However, retinal function is not always compromised and is frequently preserved in the absence of RD. On the other hand, bleeding can enhance the vitreous contractile property by introducing vasogenic and fibrogenic materials, and chronic bleeding can result in the formation of fibrovascular tissue or even RD. This finding may suggest that undergoing surgery without reservation provides a greater therapeutic benefit.

Improved baseline BCVA predicts improved postoperative BCVA in PDR eyes undergoing PPV has been extensively reported. Ramezani *et al*^[17] reviewed 236 PDR eyes undergoing vitrectomy and silicone oil injection. The findings indicated that a higher baseline BCVA was associated with a greater risk of developing final visual acuity greater than 20/200. Shen and Yang^[18] also discovered that the only variable associated with final visual acuity was preoperative visual acuity. Though functional failure could be caused by retinal redetachment, we included in the current study only eyes with postoperative macular attachment. We hypothesize that macular ischemia may be a significant factor influencing postoperative visual acuity recovery in PDR eyes, but this hypothesis requires further investigation in the future.

Typically, retinal surgeons prefer to choose the tamponade materials intraoperatively. We frequently use silicone oil or gas to treat retinal anatomical instability, such as TRD or extensive fibrous proliferation. Rush *et al*^[19] demonstrated in a randomized clinical trial that vitreous substitution with C_3F_8 gas resulted in improved visual acuity at 6mo when compared to silicone oil tamponade in PDR eyes. That result corroborated ours. In comparison to eyes with fluid or gas tamponade, eyes with silicone oil tamponade were more likely to have poor postoperative visual acuity. Numerous studies hypothesized that silicone oil posed a threat to the retina's overall health^[20]. Christensen and la Cour^[21] reported that after silicone oil tamponade, the inner retinal layers of the macula become thinner. This may account for the poor postoperative visual acuity observed in eyes treated with silicone oil tamponade.

Cataract formation is a common complication of vitrectomy^[22]. which can impair postoperative visual acuity, which is why many retinal surgeons prefer phacovitrectomy for older patients, which can save money and avoid the need for additional cataract surgery. Additionally, retinal surgeons may benefit from enhanced visibility and the ability to perform more extensive vitrectomy while in an aphakic state during surgery. To minimize the impact of cataract on visual acuity and fundus evaluation at baseline, we excluded the severe cataract (grade 3 or more dense) patients. The results indicated that eyes that had undergone phacovitrectomy were more likely to have improved postoperative visual acuity in this study. We hypothesized that the poorer postoperative visual acuity in eyes with lens preservation was due to cataract progression following vitrectomy. On the basis of the development of surgical techniques and instruments for small-gauge PPV, we advocated for phacovitrectomy in older patients.

This study has some limitations. It is a retrospective case control study with a brief follow-up period; a prospective study with a larger sample size and a longer follow-up period will be required in the future. Though we hypothesize that poor postoperative BCVA is due to macular ischemia, we lack sufficient imaging evidence to support this. In the future, quantification of macular thickness and vascular density on optical coherence tomography or optical coherence tomography angiography should be studied. A comparison of COFDS and ETDRS classifications for PDR is also necessary. In summary, we classified PDR eyes using COFDS grading and discovered that PDR eyes with primary vitreous hemorrhage typically have a better visual acuity prognosis than PDR eyes with primary fibrovascular proliferative type or RD type. These findings may aid retinal surgeons in predicting postoperative function in PDR patients. COFDS classification for PDR may have a significant predictive value for visual outcome and surgical management indications.

ACKNOWLEDGEMENTS

Foundations: Supported in part by the National Science Foundation of Liaoning Province, China (No.2020-MS-360); Shenyang Science and Technology Bureau (No.RC210267).

Conflicts of Interest: Lin TZ, None; Kong Y, None; Shi C, None; Eric Pazo E, None; Dai GZ, None; Wu XW, None; Xu L, None; Shen LJ, None.

REFERENCES

- 1 Teo ZL, Tham YC, Yu M, *et al.* Global prevalence of diabetic retinopathy and projection of burden through 2045: systematic review and meta-analysis. *Ophthalmology* 2021;128(11):1580-1591.
- 2 Grauslund J, Green A, Sjølie AK. Blindness in a 25-year followup of a population-based cohort of Danish type 1 diabetic patients. *Ophthalmology* 2009;116(11):2170-2174.
- 3 Thomas RL, Halim S, Gurudas S, Sivaprasad S, Owens DR. IDF Diabetes Atlas: a review of studies utilising retinal photography on the global prevalence of diabetes related retinopathy between 2015 and 2018. *Diabetes Res Clin Pract* 2019;157:107840.
- 4 Yang QH, Zhang Y, Zhang XM, Li XR. Prevalence of diabetic retinopathy, proliferative diabetic retinopathy and non-proliferative diabetic retinopathy in Asian T2DM patients: a systematic review and Meta-analysis. *Int J Ophthalmol* 2019;12(2):302-311.
- 5 Ip MS, Domalpally A, Hopkins JJ, Wong P, Ehrlich JS. Long-term effects of ranibizumab on diabetic retinopathy severity and progression. *Arch Ophthalmol* 2012;130(9):1145-1152.
- 6 Wilkinson CP, Ferris FL 3rd, Klein RE, Lee PP, Agardh CD, Davis M, Dills D, Kampik A, Pararajasegaram R, Verdaguer JT, Global Diabetic Retinopathy Project Group. Proposed international clinical diabetic retinopathy and diabetic macular edema disease severity scales. *Ophthalmology* 2003;110(9):1677-1682.
- 7 Chinese Ocular Fundus Diseases Society, Chinese Ophthalmological Society, Chinese Medical Association. Guidelines for clinical diagnosis and treatment of diabetic retinopathy in China. *Chin J Ophthalmol* 2014;50(11):851-865.
- 8 Gali HE, Sella R, Afshari NA. Cataract grading systems: a review of past and present. *Curr Opin Ophthalmol* 2019;30(1):13-18.
- 9 Lin T, Zhang A, Pazo EE, Hui Y, Dai G, Shen L. Comparable postoperative myopic shift in eyes with retinal vascular diseases and vitreomacular interface diseases after phacovitrectomy. *Graefes Arch Clin Exp Ophthalmol* 2022; epub ahead of print.

10 Gibbons A, Leung EH, Haddock LJ, Medina CA, Fernandez V, Parel

JMA, Durkee HA, Amescua G, Alfonso EC, Perez VL. Long-term outcomes of the aphakic snap-on Boston type I keratoprosthesis at the Bascom Palmer Eye Institute. *Clin Ophthalmol* 2018; 12:331-337.

- 11 Wong TY, Mwamburi M, Klein R, Larsen M, Flynn H, Hernandez-Medina M, Ranganathan G, Wirostko B, Pleil A, Mitchell P. Rates of progression in diabetic retinopathy during different time periods: a systematic review and meta-analysis. *Diabetes Care* 2009;32(12):2307-2313.
- 12 Flaxman SR, Bourne RRA, Resnikoff S, *et al*, Vision Loss Expert Group of the Global Burden of Disease Study. Global causes of blindness and distance vision impairment 1990-2020: a systematic review and meta-analysis. *Lancet Glob Health* 2017; 5(12):e1221-e1234.
- 13 Stitt AW, Curtis TM, Chen M, Medina RJ, McKay GJ, Jenkins A, Gardiner TA, Lyons TJ, Hammes HP, Simó R, Lois N. The progress in understanding and treatment of diabetic retinopathy. *Prog Retin Eye Res* 2016;51:156-186.
- 14 Kroll P, Meyer-Rüsenberg HW, Busse H. Recommendation for staging of proliferative diabetic retinopathy. *Fortschr Ophthalmol* 1987;84(4):360-363.
- 15 Ratnarajan G, Mellington F, Saldanha M, de Silva SR, Benjamin L. Long-term visual and retinopathy outcomes in a predominately type 2 diabetic patient population undergoing early vitrectomy and endolaser for severe vitreous haemorrhage. *Eye (Lond)* 2011;25(6):704-708;quiz 709.
- 16 Nishi K, Nishitsuka K, Yamamoto T, Yamashita H. Factors correlated with visual outcomes at two and four years after vitreous surgery for proliferative diabetic retinopathy. *PLoS One* 2021; 16(1):e0244281.
- 17 Ramezani A, Ahmadieh H, Rozegar A, Soheilian M, Entezari M, Moradian S, Dehghan MH, Nikkhah H, Yaseri M. Predictors and outcomes of vitrectomy and silicone oil injection in advanced diabetic retinopathy. *Korean J Ophthalmol* 2017;31(3):217-229.
- 18 Shen YD, Yang CM. Extended silicone oil tamponade in primary vitrectomy for complex retinal detachment in proliferative diabetic retinopathy: a long-term follow-up study. *Eur J Ophthalmol* 2007;17(6):954-960.
- 19 Rush RB, del Valle Penella A, Reinauer RM, Rush SW, Bastar PG. Silicone oil versus perfluoropropane gas tamponade during vitrectomy for tractional retinal detachment or fibrous proliferation: a randomized clinical trial. *Retina* 2021;41(7):1407-1415.
- 20 Jackson TL, Johnston RL, Donachie PHJ, Williamson TH, Sparrow JM, Steel DHW. The royal college of ophthalmologists' national ophthalmology database study of vitreoretinal surgery: report 6, diabetic vitrectomy. *JAMA Ophthalmol* 2016;134(1):79-85; quiz 120.
- 21 Christensen UC, la Cour M. Visual loss after use of intraocular silicone oil associated with thinning of inner retinal layers. *Acta Ophthalmol* 2012;90(8):733-737.
- 22 Hernandez-Bogantes E, Abdala-Figuerola A, Olivo-Payne A, Quiros F, Wu L. Cataract following pars plana vitrectomy: a review. *Semin Ophthalmol* 2021;36(8):824-831.