Outcomes and surgical management of persistent fetal vasculature

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ABSTRACT

Objective To analyse outcomes in different forms of persistent fetal vasculature (PFV).

Methods and analysis Retrospective cohort study at a university-based practice of children presenting with PFV between 2011 and 2020. Exclusion criteria was surgical management outside of our institution and follow-up less than 1 month. Wilcoxon and Student's t-tests were used for statistical analysis.

Results Forty-six eyes of 45 patients presented with PFV at 16.7±31.3 (median 2.8) months old with 32.6±29.8 (median 22.5) months of follow-up. Types of PFV included: mild combined anterior-posterior (23 eyes, 50%), severe combined anterior-posterior (18 eyes, 39%), severe anterior (3 eyes, 7%), mild anterior (1 eye, 2%) and posterior (1 eye, 2%). Thirty-two eyes (70%) underwent PFV surgical correction; lensectomy (13 mild combined), vitrectomy (3 mild combined), sequential lensectomy then vitrectomy (3 severe combined), combined lensectomy-vitrectomy (11 severe anterior or severe combined), laser retinopexy (1 mild combined). Five eyes required additional vitrectomy surgery for retinal detachment, fold or cyclitic membrane. Nine eyes developed glaucoma, six requiring intraocular pressure (IOP)-lowering surgery. At final follow-up, 32 eyes had at least form vision and 6 eyes were aversive to light. Eight eyes, all which were severe combined, and four that did not undergo PFV surgery, were unable to detect light due to phthisis bulbi (7) and optic nerve hypoplasia (1).

Conclusions Classification of PFV is important in determining surgical approach with severe cases often requiring both lensectomy and vitrectomy for optimal anatomic and functional outcomes.

BACKGROUND

Persistent fetal vasculature (PFV), which is a result of failed regression of the embryonic ocular vasculature, causes significant visual morbidity and accounts for 5% of childhood blindness in the USA.^{1 2} The majority of cases are idiopathic and unilateral with no identified genetic cause. Rare autosomal recessive mutations in *ATOH7* have been identified in familial non-syndromic bilateral PFV.^{3 4} Further, bilateral cases can be connected to systemic syndromes such as neurofibromatosis type 2, Aicardi syndrome, osteoporosis-pseudoglioma syndrome and chromosomal aneuploidies including trisomy 13, 15 and 18.^{5–11} PFV can also be associated

Key messages

What is already known about this subject?

- Persistent fetal vasculature (PFV) is classified by location as well as severity.
- Many of these eyes require surgical treatment to optimise visual potential and decrease risk of complications such as retinal detachment and glaucoma.

What are the new findings?

- Classification and severity of the PFV dictates the surgical approach and visual outcome.
- Mild cases may be managed with observation, lensectomy or lens-sparing vitrectomy.
- For severe cases of anterior and combined anterior-posterior PFV, a combined lens extraction with full vitrectomy approach was most effective in restoring anatomy and minimising complications.

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

- Careful classification and determination of severity of PFV is important in managing these challenging cases.
- A combined approach with anterior and posterior segment paediatric ophthalmic surgeons is effective in establishing vision in severe cases of PFV.

with retinopathy of prematurity and other congenital ocular abnormalities including Axenfeld-Rieger Syndrome and Peters Anomaly.^{12–14}

PFV is classified by location as well as severity (table 1) and typically presents as leucocoria with microphthalmia. Additional findings include glaucoma, cataract, corneal opacification, coloboma, retinal fold, retinal detachment, and spontaneous retinal and vitreous haemorrhages.^{15 16} The surgical approach is primarily guided by the type of PFV and secondarily by the associated ocular comorbidities, taking into context the likely visual potential of the eye, the amount of amblyopia and refractive management.^{2 17} Surgery intends to restore anatomy and maximise vision by establishing the visual axis and managing complications. Mild cases of combined anterior-posterior PFV with minimal obstruction of the pupil may be observed. However, vitrectomy to sever

Table 1 Persistent fetal vasculature classification

	Mild	Severe
Isolated anterior	Retrolental plaque without extension onto ciliary processes±cataract. No stalk or attachment to posterior pole. (1 eye (2%) of 1 patient)	Retrolental plaque (often with vessels) with extension onto ciliary processes±cataract. No stalk or attachment to posterior pole. (3 eyes (7%) of 3 patients)
Isolated posterior	Stalk or retinal fold with minimal to no optic nerve or macular distortion. No attachment to lens or ciliary processes. (1 eye (2%) of 1 patient)	Stalk or retinal fold with optic nerve or macular distortion. No attachment to lens or ciliary processes. (0 eyes)
Combined anterior-posterior	Stalk (with minimal to no distortion of the optic nerve or macula) connecting posterior pole to retrolental plaque without extension onto ciliary processes±cataract. (23 eyes (50%) of 23 patients)	Stalk (with moderate to severe distortion of the optic nerve or macula) connecting posterior pole to retrolental plaque with extension onto ciliary processes±cataract. (18 eyes (39%) of 18 patients)

the stalk connecting the posterior capsule and optic nerve can improve visual function by allowing for axial lengthening and alleviating posterior pole traction.^{18–20} Although there is limited data regarding a lens-sparing approach, maintaining phakic status in infants and young children is advantageous from an amblyopia standpoint.^{21 22} In cases in which the visual axis is blocked by cataract or retrolental membrane, the standard of care is lensectomy.^{2 23} Previous studies have shown that eyes with isolated anterior PFV have a better prognosis than eyes with posterior involvement due to the absence of macular and optic nerve pathology.^{2 20 24–26} Nevertheless, in anterior PFV that extends onto and beyond the ciliary processes, concurrent or subsequent core vitrectomy can decrease the risk of tractional retinal detachment, but may be limited by access to a multi-disciplinary surgical team.^{27 28} Isolated posterior PFV often presents later, and surgical involvement may be dictated by retinal traction or detachment. Although eyes with severe PFV, especially those with retinal detachment, have poor visual prognosis, surgical intervention may help prevent phthisis and glaucoma.^{19 29} With the advent of modern surgical technology, PFV outcomes have improved overall, however, the rarity of the disease prevents large studies to fully assess outcomes.^{19 23 30–35} The purpose of this study is to add to the literature our 9-year predominantly surgical management of PFV and visual outcomes.

METHODS

A retrospective case series identified children younger than 18 years with PFV whose initial presentation to the Kellogg Eye Center at the University of Michigan was between January 2011 and January 2020.

Data collection included age, gender, ocular findings, type of PFV and visual acuity at presentation. Duration of follow-up and anatomic status and visual acuity at final visit were also included. Optotype visual acuity (Allen figures or Snellen) was attempted on cooperative patients typically 3 years of age or older. Optotype visual acuity was converted to LogMAR scale such that 20/200 and 20/20 were equivalent to 1 and 0.1, respectively. Patients

with visual acuity worse than 20/400 were assessed by the ability to detect count fingers, hand motion or light and were not converted into LogMAR equivalents. Patients older than 3 months of age and/or uncooperative with optotype testing were assessed for ocular preference by induced tropia test with 10 base down prism. Individuals younger than 3 months of age were assessed for aversion to light as attentive fix and follow behaviour was not expected. At final follow-up, stable visual acuity was defined as within one line of initial optotype visual acuity or no change in ability to detect count fingers, hand motion or light. Improvement or decline in visual acuity was defined as two or more line change from initial optotype acuity or a change in ability to detect count fingers, hand motion or light.

Additionally, information regarding intraocular and strabismus surgeries, dates of surgeries, surgical procedure details and postoperative complications were collected. Exclusion criteria were surgical correction performed outside of our institution, age greater than 18 years at presentation, follow-up of less than 1 month, or isolated Mitterdorff dot or Bergmeister papilla. PFV was classified clinically by location (anterior, posterior or combined anterior-posterior) and severity (mild or severe) (table 1). At the time of initial surgical intervention, all patients underwent examination under general anaesthesia.

Surgical procedure using the Constellation Vitreoretinal Surgical System (Alcon Laboratories, Fort Worth, Texas, USA) varied depending on ocular pathology and surgeon preference. Briefly for lensectomy, two 23-gauge limbal incisions were created for the vitrector hand piece and infusion port. The lens was removed with the vitrector. The posterior capsule followed by the anterior hyaloid were opened and a limited anterior vitrectomy was performed. For intraocular lens (IOL) placement, keratometry and axial length were obtained at the time of surgery and SRK-T lens equation was used to calculate the appropriate power. A foldable 3-piece (MA60AC, Alcon) or single-piece (SN60WF, Alcon) lens was chosen based on position within the sulcus or the capsule. The

IOL was inserted through a superior scleral tunnel. For lens-sparing vitrectomy surgery, trocars were placed through the pars plana following the nomogram recommended by Lemley and Han.³⁶ In aphakic eyes, trocars were placed just posterior to the limbus and surgery was performed transpupillary through the same trochar cannulae. In combined lensectomy-vitrectomy surgery, both procedures were performed through the same trocars/incisions. The posterior capsule and retrolental plaque were removed and in the process the stalk was severed. Endodiathermy was needed infrequently to cauterise bleeding hyaloidal vessels. A core vitrectomy was performed and posterior hyaloid membrane and cortical vitreous were dissected using aspiration and/or manual membrane peeling. In cases with severe anterior involvement, the 19-gauge or 23-gauge endoscope (Endo-Optiks, Little Silver, New Jersey, USA) was used to visualise the pars plicata, pars plana and ora serrata, identify posterior extension of the anterior plaque and apply photocoagulation to the peripheral retina as indicated.

Aphakic eyes were fit with a contact lens. The refractive target for contact lenses in patients under 2 years of age was -1.00 to -3.00 . After the age of 2 years, the contact lens refractive target was adjusted to plano and glasses with a $+3.00$ bifocal were dispensed. In patients unable to tolerate a contact lens, a secondary IOL, with a refractive target based on age, was placed if there was adequate capsular support. Patching of the preferred eye (amount per day dictated by age and visual acuity) was recommended for all amblyopic patients. Atropine penalisation of the preferred eye was used in some patients who did tolerate or were not compliant with patching.

GraphPad Prism 8 (GraphPad, La Jolla, California, USA) was used for statistical analyses using Wilcoxon and Student's *t*-tests. The reported *p* values were two-tailed. *P* values of <0.05 were considered statistically significant.

RESULTS

Forty-five patients with PFV initially presented between January 2011 and January 2020. Median age at presentation was 2.8 months (mean 16.7 ± 31.3 months, range 0.7–138.0 months). Nineteen patients (42%) were women and 26 (58%) were men. Thirty-three patients (72%) were Caucasian, seven (15%) were African-American, four (9%) were Asian and one (2%) was Hispanic. Two patients had bilateral PFV, and both had confirmed genetic syndromes. Both eyes of one patient, who had mucopolysaccharidosis type VII (OMIM #253220) were included. Only one eye of a patient with Microphthalmia, Syndrome 12 (OMIM #180220) and Microcornea, Posterior megalolenticonus, Persistent fetal vasculature and Coloboma (MPPC) syndrome was included as the contralateral eye underwent initial surgery at an outside institution. Six patients with unilateral PFV had systemic congenital abnormalities (online supplemental table 1). Two patients had septo-optic dysplasia, both of whom had optic nerve hypoplasia in the contralateral eye. One patient with neurofibromatosis type 2 had retina/retinal

pigment epithelium combined hamartomas in both eyes and an optic nerve glioma in the PFV eye. In three patients with systemic findings (Goldenhar syndrome, craniosynostosis and ventricular septal defect/dysmorphic features), the fellow eye was normal. One patient with no systemic abnormalities had MPCC syndrome in the PFV eye and showed optic nerve hypoplasia and chorioretinal coloboma in the contralateral eye. Another patient with unilateral PFV had a history of retinopathy of prematurity that required laser in both eyes.

Forty-six eyes of 45 patients were included in the analysis (online supplemental table 2) and there was a split between right and left eyes (23 eyes each). Mild combined anterior-posterior PFV was most common (23 eyes, 50%), followed by severe combined anterior-posterior PFV (18 eyes, 39%, table 1). Three eyes (7%) had severe anterior and one eye (2%) each had mild anterior or mild posterior PFV. At initial presentation, cataract (30 eyes, 65%) and microphthalmia (16 eyes, 35%) were most frequently associated with PFV (online supplemental table 3). Due to age and cooperation, only six patients (six eyes, 13%) were cooperative for optotype visual acuity testing (LogMAR 0.3 ± 0.2 , range 0.2–0.7) at presentation (online supplemental table 2). Twenty-two eyes of 22 patients were averse to light, all of whom were under 11 months of age including 16 who were under 3 months of age. Nine eyes of nine patients were not averse to light while seven eyes of seven patients preferred the contralateral eye by induced tropia test. One eye each of the two patients with bilateral PFV was the preferred eye by induced tropia test. Patients who were able to cooperate for optotype testing were significantly older (79.9 ± 30.8 months, range 57.1–138 months, $p < 0.0001$) than the 40 patients who could not complete optotype testing (7.2 ± 17.4 months, range 0.7–105 months).

Fourteen eyes of 13 patients did not undergo surgery (table 2). Nine eyes of eight patients had poor visual potential, seven of which were due to severe combined anterior-posterior PFV in combination with extreme microphthalmia and/or chronic retinal detachment. Two eyes had mild combined anterior-posterior PFV, but vision was limited either due to corneal opacification from mucopolysaccharidosis type VII or combined hamartoma of the retina/retinal pigment epithelium and optic nerve glioma due to neurofibromatosis type 2. Surgery was not required due to minimally visually significant PFV in four eyes of four patients with mild combined anterior-posterior PFV and one eye of one patient with mild posterior PFV.

Thirty-two eyes (70% of 46 eyes) of 32 patients underwent surgery for PFV at 10.1 ± 20.9 months (median 2.4 months, range 0.7–101 months) of age (table 2). Thirteen eyes (41% of 32 eyes) of 13 patients all with mild anterior-posterior combined PFV underwent lensectomy alone. Eleven eyes (34% of 32 eyes) of 11 patients with either severe anterior PFV (three eyes) or severe combined anterior-posterior PFV (eight eyes) had combined lensectomy-vitrectomy. Three eyes (9% of 32

Table 2 Initial persistent fetal vasculature surgeries

	Mild anterior n=1 (% of n)	Mild posterior n=1 (% of n)	Mild combined n=23 (% of n)	Severe anterior n=3 (% of n)	Severe combined n=18 (% of n)	Total n=46 (% of n)
No surgery		1 eye (100%)	6 eyes (26%)		7 eyes (39%)	14 eyes (30%)
Laser retinopexy			1 eye (4%)			1 eye (2%)
Lensectomy	1 eye (100%)		12 eyes (52%)			13 eyes (28%)
Lensectomy-IOL			1 eye (4%)			1 eye (2%)
Vitrectomy			3 eyes (13%)			3 eyes (7%)
Lensectomy then vitrectomy*					3 eyes (17%)	3 eyes (7%)
Lensectomy/vitrectomy†				3 eyes (100%)	8 eyes (44%)	11 eyes (24%)

*Sequential lensectomy then vitrectomy.

†Combined lensectomy/vitrectomy surgery.
IOL, intraocular lens.

eyes) of three patients with severe combined anterior-posterior PFV had sequential lensectomy then vitrectomy and three eyes (9% of 32 eyes) of three patients with mild combined anterior-posterior PFV underwent vitrectomy alone. One eye (3% of 32 eyes) of one patient with mild combined anterior-posterior PFV each underwent lensectomy-IOL or laser retinopexy.

Twenty eyes of 20 patients, 63% of the surgical cohort (20/32 patients), required additional retinal surgery, glaucoma surgery, other anterior intraocular or strabismus surgery (2.0 ± 1.4 surgeries/eye, range 1–6, median 2) during follow-up (table 3). Five eyes, 16% of

the surgical cohort (5/32 patients), all of which initially either underwent combined lensectomy-vitrectomy or sequential lensectomy then vitrectomy, required at least one additional vitrectomy surgery (online supplemental table 4, 1.6 ± 0.5 vitrectomies/eye, range 1–2, median 2) for retinal detachment (three eyes), retinal fold (one eye), cyclitic membrane (one eye) an average of 14.9 ± 24.2 months (range 1.2–58.1 months, median 5.4 months) after the initial PFV surgery. In our surgical cohort, 25% (8/32 patients) were diagnosed with glaucoma (online supplemental table 5). In contrast, 7% (1/14 patients) were diagnosed with glaucoma in the observation cohort.

Table 3 Complications and additional surgeries

	Additional retinal surgeries	Glaucoma (# eyes requiring glaucoma surgery)	Additional anterior intraocular surgeries	Strabismus (# patients requiring strabismus surgery)
No PFV surgery (14 eyes)				
Mild posterior (1 eye)				
Mild combined (6 eyes)				3 patients (0)
Severe combined (7 eyes)		1 eye (0)		3 patients (0)
PFV surgery (32 eyes)				
Mild anterior (1 eye)				1 patient (0)
Mild combined (17 eyes)		3 eyes (1)	5 eyes	15 patients (8)
Severe anterior (3 eyes)	1 eye	1 eye (1)	1 eye	1 patient (0)
Severe combined (11 eyes)	4 eyes	4 eyes (4)	2 eyes	6 patients (1)
Total	5 eyes	9 eyes (6)	8 eyes	29 patients (9)
No PFV surgery per cent (14 eyes)	0	7	0	43
PFV surgery per cent (32 eyes)	16	25	25	72

PFV, persistent fetal vasculature.

Six eyes, all of which underwent surgery for PFV (75%, 6/8 patients), required glaucoma surgery to obtain intraocular pressure control (2.0 ± 0.6 glaucoma surgeries/eye, range 1–3, median 2) an average of 33.4 ± 39.1 months (range 4.4–105.7 months, median 39.5 months) after the initial PFV surgery. Eight patients had additional anterior intraocular (non-glaucoma) surgeries (1.2 ± 0.5 surgeries/eye, range 1–2, median 1) for secondary IOL placement (five eyes), pupilloplasty/synechiolysis (three eyes), repeat posterior capsulotomy (one eye) and attempted IOL placement (one eye) (online supplemental table 2). Three aphakic eyes required surgery for posterior capsular opacification, synechiae or membrane formation while one pseudophakic eye required a pupilloplasty. Twenty-nine of the patients had strabismus (19 esotropia, 10 exotropia), nine of which underwent eye muscle surgery (1.2 ± 0.4 strabismus surgeries/patient, range 1–2, median 1).

At final follow-up (32.6 ± 29.8 months, range 1.0–108.8 months, median 22.5 months), 42 eyes of 42 patients were amblyopic due to unilateral disease and had undergone part-time occlusion and/or atropine penalisation. One eye with mild posterior PFV had no amblyopia. In the patient with mucopolysaccharidosis type VII, although neither eye underwent surgery, the preferred eye had milder PFV compared with the non-preferred eye. In the patient with microphthalmia syndrome 12 and bilateral MPCC, only the preferred eye was included as the non-preferred eye had undergone initial surgery at an outside institution. Twenty-two eyes of 22 patients were aphakic,

6 eyes of 6 patients had IOLs and 18 eyes of 17 patients were phakic. Of the 18 eyes that were phakic, 14 did not have surgery, 1 had laser retinopexy and 3 had vitrectomy alone.

Measurement of visual acuity was limited at presentation and final follow-up in many of the patients by age, cooperation and level of vision (online supplemental table 2). The majority of patients in the study, 91% (42/46 eyes), showed either improvement or stability of visual acuity at final follow-up (table 4). Within the PFV surgical cohort 66% (21/32 eyes) had improvement in visual acuity, 25% (8/32 eyes) showed stable visual acuity and 9% (3/32 eyes) displayed worse visual acuity. The three eyes that had worse vision developed unreparable retinal detachments and eventually became phthisical. Within the cohort that did not undergo surgical management for PFV, at final follow-up, 21% (3/14 eyes) had improved visual acuity, 71% (10/14 eyes) showed no change in visual acuity and 7% (1/14 eyes) had worsened vision. At final follow-up, 12 eyes of 12 patients cooperated for optotype testing (online supplemental table 6, LogMAR 0.5 ± 0.3 , range 0.1–1.1, median 0.3). Eight of these 12 eyes were phakic (LogMAR 0.3 ± 0.1) and had significantly better vision ($p=0.004$) than the four eyes (LogMAR 0.8 ± 0.4) that were aphakic (two eyes) or pseudophakic (two eyes). Thirty-three patients (34 eyes) were unable to complete optotype testing. Six eyes were averse to light and two eyes were able to count fingers at 4–8 inches. Eighteen eyes of 18 patients were able to fix and follow, with 16 of the patients showing a preference for

Table 4 Vision comparison from presentation to final follow-up

	Improved vision	Stable vision	Worse vision	Total
No surgery				
Mild posterior		1 eye		1 eye
Mild combined	1 eye	5 eyes		6 eyes
Severe combined	2 eyes	4 eyes	1 eye	7 eyes
Retinal laser				
Mild combined	1 eye			1 eye
Lensectomy or lensectomy-IOL				
Mild anterior	1 eye			1 eye
Mild combined	9 eyes	4 eyes		13 eyes
Vitrectomy				
Mild combined	2 eyes	1 eye		3 eyes
Lensectomy then vitrectomy				
Severe combined	1 eye		2 eyes	3 eyes
Combined lensectomy/vitrectomy				
Severe anterior	3 eyes			3 eyes
Severe combined	4 eyes	3 eyes	1 eye	8 eyes
Total	24 eyes	18 eyes	4 eyes	46 eyes
No PFV surgery per cent (14 eyes)	21	71	7	
PFV surgery per cent (32 eyes)	66	25	9	

IOP, intraocular lens; PFV, persistent fetal vasculature.

the contralateral eye. Eight eyes were unable to detect light due to phthisis bulbi from chronic retinal detachment (seven eye) and optic nerve hypoplasia (one eye).

DISCUSSION

PFV is a significant cause of childhood blindness due to both the primary anatomic malformations and secondary complications such as retinal detachment, glaucoma and amblyopia.^{9 16} We describe our experience with PFV over a 9-year period assessing medical and surgical management and anatomic and functional outcomes in these complex eyes.

While over 90% of cases are sporadic and unilateral, there are rare cases of bilateral PFV which are typically associated with genetic syndromes.¹⁶ Of note, our study included two patients (4%) with bilateral PFV. One patient had mucopolysaccharidosis type VII, an extremely rare form of mucopolysaccharidosis due to β -glucuronidase deficiency that often causes prenatal or perinatal death.³⁷ Mucopolysaccharidosis type VII has not been previously correlated with PFV, thus it is unclear whether this ocular anomaly was coincidental or due to early fetal glycosaminoglycan deposition within the eye. The second patient had microphthalmia, syndrome 12, which is due to retinoic acid receptor beta mutations.³⁸ Our patient also had a constellation of ocular findings known as MPPC, which is a very severe form of microphthalmia and PFV where the lens is posteriorly displaced and occupies the majority of the vitreous cavity.³⁹ Another patient also had MPPC of one eye, while the contralateral eye showed optic nerve hypoplasia and peripheral chorioretinal coloboma. To date, this patient does not have systemic findings or a genetic diagnosis. Further, there are no confirmed genetic mutations associated with MPCC showing that additional studies are required on this severe eye abnormality. It is important to note that in addition to these three cases, five patients with unilateral PFV had systemic findings including neurofibromatosis type 2, septo-optic dysplasia, Goldenhar syndrome, craniosynostosis and ventricular septal defect. While many of these diseases have been previously associated with PFV, this serves as a reminder that even in unilateral cases, systemic diagnoses and contralateral eye findings should be investigated.^{10 40 41}

The decision for surgical management is based on patient age and visual prognosis as dictated by the type of PFV and associated macular and optic nerve abnormalities. Age at presentation has been shown to be a significant predictor of visual outcome, with previous reports demonstrating that early intervention leads to improved visual acuity.^{42 43} While the average age of patients who underwent surgical management in our cohort was 10.1 ± 20.9 months, the median age was 2.4 months which is within the timeframe of critical visual development (6–10 weeks). Further, 23 of the 32 eyes undergoing surgery before 6 months of age, of which at final follow-up, 15 showed improved vision, 5 had stable vision and 3 had worse vision. Of the nine eyes

who underwent surgery, there were two outliers who presented late (5 and 8 years of age) with mild amblyopia and subsequently underwent lens-sparing surgery for mild combined anterior-posterior PFV. Further, despite the higher mean age at surgery, our postoperative visual outcomes were comparable to a handful of studies in which at least 30 eyes with PFV underwent surgical intervention at mean ages between 2 and 3 months (table 5).^{19 20 30}

The type of PFV is a critical consideration in management. The majority of cases in many reports and in our series involve both the anterior and posterior fetal vasculatures.^{19 20 23 30 35} Mild cases of combined PFV are characterised by a stalk with minimal optic nerve distortion that connects to the posterior capsule. In our study, the presence of a visually significant cataract or traction on the posterior pole dictated whether surgery was required. Based on this criteria, four eyes with mild combined anterior-posterior PFV did not require surgical management. An additional three eyes underwent lens-sparing vitrectomy to relieve traction and one eye had laser retinopexy. Sparing of the lens in cases where there is not a visually significant cataract prevents aphakic/pseudophakic anisometropia as well as amblyogenic loss of accommodation.^{21 22} Certainly in our study, these three phakic patients had visual acuities of 20/40–20/60 at final follow-up. Other studies have also employed lens-sparing vitrectomy, however, visual outcome data in these patients is limited as this is often a small subgroup of PFV (table 5).^{19 20 30} Nevertheless, It is important to note that even with segmentation of the stalk, there may be astigmatic anisometropia from non-cataractous lens distortion that must be managed to treat amblyopia. Mild combined anterior-posterior PFV with a visually significant cataract is most commonly managed through lensectomy $\pm$ IOL by an anterior segment surgeon or paediatric ophthalmologist.^{23 24 32 34} In the majority of these cases, the visual potential is favourable.^{2 20 24 25} However, unilateral aphakia and pseudophakia are highly amblyogenic, and so persistent and steadfast amblyopia management is required to attain 20/40 or better visual acuity.^{22 44} Similar to previous reports, in our study 3 of 14 eyes with mild combined anterior-posterior PFV which underwent lensectomy $\pm$ IOL had visual acuity of 20/200 or better and only one had visual acuity of 20/40.^{23 25 32 35}

Severe PFV presents a more complicated problem due to inherent anterior and posterior abnormalities which predispose these eyes for retinal complications and glaucoma. In addition, involvement of the optic nerve and macula are associated with worse outcomes.^{19 31 33 45} Due to these inherent congenital abnormalities, electrophysiology has been used to predict visual potential, and thus identify eyes in which surgery may not be beneficial.⁴⁶ As a practice, we do not routinely conduct preoperative visual evoked potential testing, as there may be inaccuracies due to the age and inherent immaturity of the visual system in young infants.⁴⁷ Instead we base the decision to perform surgery in these severe cases on the eye anatomy, initial

Table 5 Comparison of previous series analysing surgical management of PFV

Author	PFV type n (%)	Surgical repair n (%)	Eyes with PFV surgery, n	Mean age, months (range)	Mean follow-up, months (range)	Lens status n (%)	Preoperative visual acuity: n (%)	Postoperative visual acuity: n (%)	Postoperative complications n (%)
Sisk <i>et al</i> ¹⁹	Anterior 26 (37) Posterior 7 (10) Combined 37 (53)	Lensectomy 9 (13) Vitrectomy 3 (4) Lensectomy/ vitrectomy 58 (83)	70	2.25 (0–252)	47* (6–324)	Aphakic 61 (87) Pseudo 6 (9) Phakic 3 (4)	LogMAR 1.2±0.5 6 (14) TAC 0.3–3.2 c/cm 6 (14) CF 7 (10) CSM 1 (1) F&F 3 (4) No F&F 7 (10) BR 40 (57)	LogMAR 1.24±0.6 27 (39) TAC 0.23–9.8 c/cm 7 (10) CF/HM 5 (7) CSM 1 (1) F&F 12 (17) No F&F 5 (7) LP 2 (3) NLP 11 (23)	Retinal detachment 15 (21) Glaucoma 8 (11)
Karacorlu <i>et al</i> ²⁰	Anterior 5 (11) Posterior 5 (11) Combined 34 (78)	Sequential lensectomy then vitrectomy 7 (36) Lensectomy/ vitrectomy 32 (73) Vitrectomy 5 (11)	44	3 (1–36)	37.2 (12–164)	Aphakic 38 (87) Pseudo 1 (2) Phakic 5 (11)	LogMAR 3.0† 2 (5) CSM/F&F 42 (95)	LogMAR 1.15±0.6 14 (32) CSM/F&F 21 (48) LP/NLP- 9 (20)	Retinal tear 3 (7) Retinal detachment 6 (14) Glaucoma 6 (14) Strabismus 26 (63)
Tartarella <i>et al</i> ²³	Anterior 5 (9) Posterior 6 (11) Combined 42 (79)	Lensectomy 20 (5) Lensectomy/ anterior vitrectomy† 18 (47)	38	14 (1–48)	44	Aphakic 5 (13) Pseudo 15 (40) Phakic 15 (40)	LogMAR 0.88 to LP 38 (100)	LogMAR≤0.3 3 (8) LogMAR 0.3–1 17 (45) LogMAR≥1 12 (32) NLP 3 (8)	Retinal detachment 2 (5) Glaucoma 4 (11) Vitreous haemorrhage 2 (5)

Continued

Table 5 Continued

Author	PFV type n (%)	Surgical repair n (%)	Eyes with PFV surgery, n	Mean age, months (range)	Mean follow-up, months (range)	Lens status n (%)	Preoperative visual acuity: n (%)	Postoperative visual acuity: n (%)	Postoperative complications n (%)
Hunt <i>et al</i> ³⁰	Anterior 15 (27) Posterior 7 (13) Combined 33 (60)	Lensectomy 11 (35) Lensectomy/ vitrectomy 20 (65)	31	2.6* (0–15)	28 (1–116)	Aphakic 29 (94) Pseudo 2 (6)	F&F 10 (30) No F&F 2 (6) LP 4 (12)	LogMAR 1.2±0.4 14(45) F&F 2 (6) No F&F 1 (3) LP 4 (13) NLP 7 (23)	Retinal detachment 4 (13) Glaucoma 5 (16) Strabismus 19 (61)
Current study	Anterior 4 (9) Posterior 1 (2) Combined 41 (89)	Lensectomy 14 (44) Vitrectomy 3 (9) Sequential lensectomy then vitrectomy 3 (9) Lensectomy/ vitrectomy 11 (34) Laser retinopexy 1 (3)	32	10 (1–101)	33 (1–108)	Aphakic 22 (69) Pseudo 6 (19) Phakic 4 (13)	LogMAR 0.3±0.1 2 (6) F&F 6 (19) LP 22 (69) NLP 2 (6)	LogMAR 0.54±0.4 8 (25) F&F 15 (47) CF 1 (3) LP 4 (13) NLP 4 (13)	Retinal detachment 3 (9) Glaucoma 8 (25) Strabismus 29 (91)

All cited studies were retrospective in study design.

*Median value as mean not reported.

†Two patients with visual acuity of 20/20 000; LogMAR equivalent 3.0.

‡Anterior vitrectomy=opening of anterior hyaloid, without full core vitrectomy.

BR, blink reflex; CF, count fingers; CSM, central, steady, maintained; F&F, fix and follow; FM, form vision; HM, hand motion; NLP, no light perception; PFV, persistent fetal vasculature; TAC, Teller acuity cards.

vision, status of the contralateral eye and parental desire. Observation is an option especially in cases where there is a closed funnel retinal detachment, severe optic hypoplasia and a normal contralateral eye. However, as the majority of these eyes, without intervention, eventually lose all vision from retinal detachment or glaucoma, we do tend to have a more aggressive approach.^{6 29 31} Of course this pro-surgical avenue requires in-depth discussion with the parents in order to manage expectations regarding visual outcomes and need for multiple surgeries. In our study, of the 18 eyes with severe combined PFV, surgery was not recommended for 7 eyes. In this observational cohort at final follow-up, 57% (4/7 eyes) were unable to detect light, of which three of these eyes (43%) had phthisis from retinal detachment. In contrast, at final follow-up in the 11 severe combined PFV eyes that underwent surgical intervention, 27% (3/11 eyes) were unable to detect light due to phthisis bulbi. While surgical management of combined severe PFV poses challenges due to the risk of postoperative complications, similar to our results, previous studies have noted successful improvement in visual outcomes.^{27 48} Soheilian *et al* noted significant improvement in visual acuity in patients with combined PFV (OR 7.9, p value =0.07), possibly explained by anatomically successful retinal surgery, proper optical correction and amblyopia therapy.²⁷ In addition, Mittra *et al* noted that 71% of patients presenting with combined PFV experienced improvement in final visual acuity of 20/300 or better.⁴⁸

Previous studies have demonstrated modest results with lensectomy±vitrectomy, which may be performed from a pars plana or limbal approach.^{19 20 30 33} Our management for severe anterior and severe combined anterior-posterior PFV was typically a combined lensectomy-vitrectomy surgery coordinated between paediatric anterior segment and retina specialists. The lens was first removed from an anterior approach, followed by a limbal vitrectomy which focused on careful inspection of the peripheral retina and segmentation of a stalk if present. This approach has also been successfully employed by other groups (table 5).^{19 20 30} It is not uncommon for retrolental plaques to extend beyond the ora serrata causing traction and detachment of the peripheral retina.^{17 49} Due to this we often employed the endoscope to visualise the pars plana and pars plicata to rule out posterior plaque extension and retinal traction. In some cases, retrolental plaque demonstrated focal to extensive posterior extension, leading to anterior retinal dragging and tenting of the retina over the pars plana and pars plicata. Endoscopic-guided vitrectomy in these cases reduces the rate of iatrogenic retinal breaks and intraoperative and postoperative retinal detachment. Endoscopy also allows the application of laser photocoagulation to the far periphery to surround focal retinal tractions and defects as needed.⁵⁰ Lack of adequate visualisation and treatment of the peripheral retina predisposes these eyes for retinal detachments complicated by retinal dialyses and proliferative vitreoretinopathy. The combined

lensectomy-vitrectomy approach with anterior and posterior surgeons is advantageous over pars plana lensectomy and vitrectomy in that it often allowed for maintenance of capsular support for a future IOL. However, these eyes require close monitoring as there can be postoperative contracture of the remnant cyclitic/peripheral retinal membrane and capsule resulting in pupil constriction and in some cases tractional retinal detachment, iris bombe and glaucoma.^{17 31 49} In our cohort, with this coordinated approach, 1 of the 11 eyes had worse vision due to an irreparable retinal detachment, while 10 of the eyes showed improved or stable vision at final follow-up. Like ours, other studies which used coordinated lensectomy-vitrectomy surgery for PFV, also employed other surgical approaches (table 5). Nevertheless, there is a lack of visual and anatomic outcomes based on different surgical approaches in these prior reports. Thus, more studies focused on this coordinated approach are needed to verify that the effectiveness of establishing vision in these severely affected eyes that would otherwise likely be completely blind.

The advantage of the combined lensectomy-vitrectomy over sequential lensectomy then vitrectomy is less scarring by using the same incisions/ports and fewer exposures to general anaesthesia. In our series, the sequential surgeries in three eyes were not preoperatively planned, but due to the unanticipated finding that there was greater posterior segment involvement after lens removal. Similarly in Karacorlu *et al*, seven eyes first underwent lensectomy and then were referred to the retinal group for vitrectomy.²⁰ In our cohort, this recategorised these three eyes from mild combined to severe combined PFV and necessitated subsequent vitrectomy which was performed within 2 weeks in two eyes. The third patient was lost to follow-up and vitrectomy was not completed until over a year later. Unfortunately, two of these eyes were phthisical at final follow-up. In Karacorlu *et al*, information regarding time between lensectomy and vitrectomy as well as visual outcomes in this subgroup are not reported. Thus, it is difficult to determine whether sequential surgeries, at least in our hands, had worse anatomic and visual outcomes compared with the combined lensectomy-vitrectomy.

Additional complications associated with PFV include retinal detachment and glaucoma. While mild anterior and mild combined PFV rarely have retinal complications, severe PFV are associated with vision compromising optic nerve and macular abnormalities as well as retinal detachments and folds.^{19 24 33} In our cohort, 16% of eyes which underwent PFV surgery (5/32 eyes), all with severe PFV, developed postoperative retinal detachments or tractional folds (following initial surgical intervention) that required additional vitrectomy. Three of these eyes had anatomic success and vision remained stable, while two became phthisical. Another eye developed an intraoperative irreparable retinal detachment at the time of initial vitrectomy and ultimately also developed phthisis. Our rate of postoperative retinal complications and

phthisis is similar to other studies where these ranged from 5%–21% to 8%–23%, respectively (table 5).^{19 20 23 30} Thus, it is important to carefully monitor for postoperative retinal complications in eyes with PFV.

Glaucoma is a well-recognised complication of PFV and elevated intraocular pressure is often multi-factorial given early lens removal and additional congenital abnormalities.^{24 49 51} Glaucoma following cataract surgery (GFCS) is associated with early cataract removal, with the highest rates in eyes that have undergone surgery before 3 months of age.^{52–54} However, previous studies have found no difference in the rate of glaucoma in aphakic eyes with or without PFV.³⁴ Studies have demonstrated that unlike primary congenital glaucoma, medical therapy can be effective in over 50% eyes with GFCS.^{51 55} However, angle surgery has a lower success rate which is likely due to the different pathologies in eyes with congenital cataracts compared with primary congenital glaucoma.^{56 57} In our cohort, 20% (9/46) eyes were diagnosed with glaucoma, with the rate higher in the PFV surgery cohort (25%, 8/32 eyes). This rate of glaucoma is higher than many of the previously published studies (table 5). This may be due to differences in the severity of PFV within the different reports, although these reports do not give specific details regarding which eyes developed elevated intraocular pressure.^{19 20 23 30} In our study, six of the nine eyes with glaucoma required surgery. Trabeculotomy was successful in one eye which had mild combined PFV. All eyes with severe forms of PFV and glaucoma also had microphthalmia, and required glaucoma drainage devices±cyclophotocoagulation to achieve Intraocular pressure (IOP) control. Few studies have specifically looked at the management of glaucoma in PFV, but rather categorise this with GCFS. A recent study showed that Ahmed glaucoma drainage devices were less successful and had a higher rate of complications in PFV compared with GCFS without PFV.⁵⁸ In our study, Baerveldt glaucoma drainage devices were used in four eyes while an Ahmed was placed in one eye. Additional studies are required to better assess effective management of glaucoma in these severe PFV eyes.

The strengths of our study are the relatively high number of patients given the rarity of PFV. Limitations include the retrospective nature of the case series, variable disease severity at presentation, variable length of follow-up, and lack of control population. Further, although the study included 45 patients, 32 of whom underwent surgery for PFV, the total number of eyes in each surgical category is small. A prospective trial with greater volume of patients is difficult due to the rarity of PFV, although a multi-centre collaborative study would address this problem.

In conclusion, we reviewed our experience with PFV. While 70% of eyes underwent surgical correction for PFV, it is important to note that some eyes do not require surgery either due to non-visually significant findings or poor visual potential. For eyes with visually significant findings, classification of PFV is critical for determining

proper surgical management. While mild cases of PFV may be managed with lensectomy or vitrectomy alone, severe cases typically require both lensectomy and vitrectomy. In our hands, a combined lensectomy-vitrectomy surgical approach was most effective in establishing vision in these complex cases. Surgical management of PFV, particularly in cases of significant posterior involvement, may have limited outcomes due to abnormal neuroretinal anatomy and postsurgical complications, including retinal detachment and glaucoma. In addition, the majority of these eyes also require appropriate treatments for amblyopia and strabismus to maximise visual potential. Our study provides additional information that paediatric ophthalmologists and retinal specialists can use during their consultation with the parents and guide the discussion of the meticulous management of PFV.

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