One-year results of aflibercept treatment for polypoidal choroidal vasculopathy with good visual acuity

Dear Editors:

We are submitting a full-length article entitled “One-year results of aflibercept treatment for polypoidal choroidal vasculopathy with good visual acuity” for consideration for publication in Korean Journal of Ophthalmology.

In most studies of intravitreal anti-VEGF injections for PCV, the initial BCVA is less than 20/40. Whereas there are several reports of intravitreal ranibizumab injections for PCV with good initial visual acuity, studies on intravitreal aflibercept injections for PCV with good initial BCVA are sparse. In this study, we figured out that intravitreal aflibercept injections for PCV with initial BCVA of 20/40 or better were effective in maintaining good visual acuity and reducing CSMT for one year.

This manuscript has not been published or presented elsewhere in part or in entirety and is not under consideration by another journal.

Thank you for your consideration. We look forward to hearing from you.

Sincerely,
One-year Aflibercept in PCV with good vision

One-year results of aflibercept treatment for polypoidal choroidal vasculopathy with good visual acuity

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Abstract

Purpose: To evaluate the one-year efficacy of intravitreal aflibercept injection in polypoidal choroidal vasculopathy (PCV) with best-corrected visual acuity (BCVA) of 20/40 or better.

Methods: This was a multicenter retrospective study. The medical records of patients diagnosed with treatment-naive PCV were retrospectively reviewed. Patients with an initial BCVA of 20/40 or better and who had undergone intravitreal aflibercept injection were included. Patients were treated with 3 consecutive monthly injections, followed by pro re nata regimen according to the clinician’s discretion at variable interval visits. The proportions of eyes for which BCVA was maintained (≤ 0.2 logMAR change) or improved at 12 months were evaluated. The changes of BCVA, central subfield macular thickness (CSMT), pigment epithelial detachment (PED), and subretinal fluid (SRF) also were assessed.

Results: A total of 86 eyes were included. The mean number of injections for 12 months of treatment was 5.4 ± 1.7. BCVA was maintained or improved in 94.2% (81/86) of cases. Mean BCVA (logMAR) had changed from the baseline (0.23 ± 0.09) at 3 months (0.21 ± 0.14), 6 months (0.24 ± 0.22) and 12 months (0.20 ± 0.18), but with no statistical significance. CSMT had improved significantly from the baseline (336.1 ± 97.3 μm) at 3 months (223.6 ± 47.22 μm), 6 months (239.6 ± 64.2 μm), and 12 months (223.8 ± 47.9 μm). PED was observed in 93% of cases at the baseline, 72.1% at 3 months, and 69.8% at 12 months, showing a significant decrease at all observation points. SRF was observed in 91.9% of cases at the baseline, 20.9% at 3 months, and 29.1% at 12 months, showing a significant decrease at all observation points.

Conclusion: In cases of PCV with good visual acuity, intravitreal aflibercept injections decreased CSMT and were effective in maintaining visual acuity.

Keywords: Aflibercept, Intravitreal injection, Polypoidal choroidal vasculopathy
**Introduction**

Polypoidal choroidal vasculopathy (PCV) is a type of neovascular age-related macular degeneration (AMD) that is characterized by a network of branching choroidal vessels with terminal, polyp-like aneurysmal dilations [1, 2]. PCV is more prevalent in the East than in the West. It accounts for up to 54.7% of neovascular AMD in Asians, 22.2–36.3% in South Koreans, while only about 8–13% in Caucasians [3–8]. The natural course and response to treatment often differ from typical neovascular AMD. PCV usually causes fewer fibrotic scars, progresses slowly, and shows a favorable prognosis relative to typical neovascular AMD [2, 9]. However, it often causes persistent bleeding and exudative changes, leading to a chronic and recurrent course that ends in poor visual prognosis due to retinal pigment epithelial (RPE) atrophy [10, 11].

Photodynamic therapy (PDT) is an effective treatment for closure of polypoidal lesions but is less effective for regression of branching vascular networks; indeed, only limited improvement of visual acuity has been reported from long term follow-up studies [12, 13]. Monotherapy with ranibizumab (Lucentis, Genentech, South San Francisco, CA, USA), an anti-vascular endothelial growth factor (anti-VEGF) monoclonal antibody, has effected significant visual improvement [14] but only a limited polyp regression rate of 25–40% [15–18]. Aflibercept (Eylea, Regeneron, Tarrytown, NY, USA), a recombinant fusion protein of anti-VEGF, has been reported to have similar efficacy and safety to ranibizumab with longer half-time in neovascular AMD, and there have been reports of the administration of intravitreal aflibercept injections for PCV as well. In fact, many studies have noted visual and anatomical improvements associated with intravitreal aflibercept injections, but in most cases, the initial best-corrected visual acuity (BCVA) was worse than 20/40 [19, 20].

While many studies have been reported using aflibercept in AMD with good visual acuity, there were few studies only on PCV patients. In addition, the number of PCV patients was small in studies of AMD patients including PCV. The purpose of the present study was to evaluate the 1-year efficacy of intravitreal aflibercept injections for PCV in eyes with BCVA of 20/40 or better.

**Materials and Methods**

This was a multi-center retrospective observational study that adhered to the tenets of the Declaration of Helsinki and was approved by the Institutional Review Board (IRB) of ***** Hospital (Approval number: 2019-03-016-001). We retrospectively reviewed the medical records of patients with PCV who had received intravitreal aflibercept injections and been followed up for 12 months between January 2014 and December 2018 at ***** Hospital, ***** Hospital, ***** Medical Center, and ***** Hospital. The diagnosis of PCV, made by
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Indocyanine green angiography (ICGA), was based primarily on findings of polypoidal hyperfluorescent lesions and branching vascular networks, and was confirmed when intraretinal or subretinal fluid was accompanied on optical coherence tomography (OCT). The study-inclusion criteria were initial BCVA of 20/40 or better, and no change of treatment drug for at least 1 year after the initial administration. The exclusion criteria were as follows: previous history of intravitreal anti-VEGF or PDT; severe media opacity, previous history of vitreoretinal surgery, proliferative diabetic retinopathy, retinal vascular occlusion, macular hole, or any other retinal disorder that could influence macular microstructure and/or function.

At baseline, patients’ history of systemic and ophthalmic diseases was checked prior to initiation of treatment. BCVA was measured on the Snellen visual acuity chart. Slit-lamp microscopy, fundus examinations, fundus photography, OCT, and fluorescein and indocyanine green angiography were performed. Fundus photographs and OCT images were obtained according to each center’s setting (including TRC-NW8, Nonmyd7, TRC-50DX for fundus photographs and Cirrus HD-OCT, DRI OCT-1 for OCT images). All hospitals used Spectralis HRA (Heidelberg Engineering GmbH, Heidelberg, Germany) for fluorescein and indocyanine green angiography.

BCVA and OCT were measured at 3, 6, 9, and 12 months after treatment.

All of the patients were given an initial 3 loadings of intravitreal aflibercept (2.0 mg in 0.05 mL) at 1-month intervals. After the 3 loading injections, follow-up visits were performed every 1 to 3 months, according to the clinician’s judgment. BCVA, fundus examinations, fundus photography, and OCT also were checked or performed at each visit. Additional injections were performed on an “as-needed” basis (i.e., in cases where OCT-detected subretinal or intraretinal fluid threatened the fovea, or retinal or subretinal hemorrhage involved the macula) (Figure 1).

The BCVA was converted to the logarithm of minimal angle of resolution (logMAR) scale for analysis. BCVA 3, 6, 9, and 12 months after treatment were compared with the baseline, and the proportion of eyes that maintained or improved visual acuity was evaluated. The change of BCVA was classified as improvement, maintenance and worsen with a cut-off value of 0.2 logMAR [21]. Central subfield macular thickness (CSMT) 3, 6, 9, and 12 months after treatment were compared with the baseline, as were changes of pigment epithelial detachment (PED), subretinal fluid (SRF), and intraretinal fluid (IRF). CSMT was defined as the average retinal thickness in the 1mm-diameter center of the fovea. PED was defined as an elevation of the RPE layer over Bruch’s membrane, SRF was defined as a hyporeflective space between neurosensory retina and RPE. Both were assessed based on the OCT line scans of fovea, and only cases that violated the fovea were counted.

Changes in BCVA and CSMT from the baseline were assessed by Wilcoxon signed-rank test. The proportions
of eyes that maintained or improved BCVA at 3 and 12 months after treatment, and the proportions of eyes with PED, SRF, and IRF at 3, 6, and 12 months after treatment, were compared by McNemar test. The visual outcomes were analyzed using Mann–Whitney U test by subgrouping patients according to baseline BCVA, baseline CSMT, presence of SRF at 3 months, and total injection numbers. To analyze factors affecting final BCVA, multiple linear regression analysis was performed. The analyses were conducted using SPSS software version 22.0 (SPSS, Chicago, IL, USA), and \( P \) values < 0.05 were considered statistically significant.

**Results**

A total of 86 eyes of 86 patients, 60 males and 26 females, were included in the study. The mean age was 65.6 ± 7.26 years (Table 1).

The mean number of injections was 5.4 ± 1.7. BCVA was maintained or improved in 94.2% (81/86) of cases: 83.7% (72/86) maintenance; 10.5% (9/86) improvement. In eyes with maintained BCVA (changes in BCVA ≤ 0.2 logMAR), 12 (14%) showed a gain of 0.2 logMAR, and 21 (24.4%) a gain of 0.1 logMAR (Figure 2). The proportion of eyes that maintained or improved BCVA at 12 months after treatment (94.2%) was not significantly different from that at 3 months (97.7%) (\( P = 0.371 \)).

The mean BCVA was 0.23 ± 0.09, 0.21 ± 0.14, 0.24 ± 0.22, 0.21 ± 0.20, and 0.20 ± 0.18 at baseline and 3, 6, 9, and 12 months after treatment, respectively (Figure 3a). There was no significant relative-to-baseline improvement at any of the observation points after treatment (\( P = 0.116, P = 0.544, P = 0.080, \) and \( P = 0.068, \) respectively). The mean CSMT was 336.1 ± 97.3 μm, 223.6 ± 47.22 μm, 239.6 ± 64.2 μm, 229.4 ± 47.1 μm, and 223.8 ± 47.9 μm at the baseline and 3, 6, 9, and 12 months after treatment, respectively (Figure 3b). Significant improvement was seen at all observation points after treatment relative to the baseline (\( P < 0.001, P < 0.001, P < 0.001, \) and \( P < 0.001, \) respectively).

PED was observed in 93% (80/86) of cases at the baseline, 72.1% (62/86) at 3 months, 72.1% (62/86) at 6 months, and 69.8% (60/86) at 12 months. The proportion of PED was significantly decreased at all observation points relative to the baseline (\( P = 0.001, P = 0.001, P < 0.001, \) respectively, Figure 4). SRF was observed in 91.9% (79/86) of cases at the baseline, 20.9% (18/86) at 3 months, 30.2% (26/86) at 6 months, and 29.1% (25/86) at 12 months. The proportion of SRF was significantly decreased at all observation points relative to the baseline (\( P < 0.001, P < 0.001, P < 0.001, \) Figure 4). IRF was observed in 20.9% (18/86) of cases at the baseline, 3.5% (3/86) at 3 months, 5.8% (5/86) at 6 months, and 3.5% (3/86) at 12 months. The proportion of
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SRF was significantly decreased at all observation points relative to the baseline ($P < 0.001$, $P = 0.001$, $P < 0.001$, Figure 4).

Final visual outcomes were analyzed by subgrouping patients according to baseline BCVA, baseline CSMT, presence of SRF at 3 month, and total injection numbers (Figure 5). The patients with baseline BCVA of 0.2 logMAR or better had significantly better final BCVA than the patients with baseline BCVA worse than 0.2 logMAR (Figure 5a). Stratification based on baseline CSMT of $300 \ \mu m$ showed a significant difference in BCVA only before treatment, and no significant difference was observed during the study period (Figure 5b). The group with SRF at 3 months had significantly worse final BCVA (Figure 5c), and stratification according to the number of injections showed no significant difference in BCVA (Figure 5d). Similarly, in a multiple linear regression analysis, baseline BCVA and SRF at 3 months were found to affect final BCVA (Table 2).

A total of 466 injections were performed over 12 months, with no serious adverse events such as endophthalmitis or retinal detachment. Five eyes showed decreased BCVA despite treatment, due to a submacular hemorrhage in 1 eye and macular atrophy in the other 4.

A representative case is shown in Figure 6. A 62-year-old man diagnosed with PCV maintained good visual acuity of 20/40 with a total of 7 intravitreal aflibercept injections.

Discussion

PCV and neovascular AMD share similarities in their association with choroidal neovascularization and genetic variations, but they also show some differences [13, 14]. Choroidal thickness is thicker in PCV, and fibrotic progression is more common in neovascular AMD. Although the expression level of VEGF is increased in both PCV and neovascular AMD as compared with healthy eyes, the VEGF level in eyes with PCV is considerably lower than in AMD eyes [22, 23]. There has also been an increase in inflammatory cytokines, including interleukin-1β and interleukin-23, in the anterior chamber and vitreous bodies, suggesting that they play an inflammatory-response role in PCV [24, 25]. These differences might be among the reasons why the therapeutic response of anti-VEGF agents in PCV is not as good as in typical neovascular AMD.

Treatment strategy for PCV varies according to studies. PDT causes polyp regression by selectively closing neovascularization, and was widely used for PCV before the development of anti-VEGF agents. However, as it was less effective in the closure of branching vascular networks than for polyps, limitations in visual improvement were identified in reports with long follow-up periods [12]. Rare but serious complications, such as subretinal hemorrhage, choroidal infarction, and RPE rupture, also have been reported [26–28]. After
confirmation of the effects of ranibizumab on neovascular AMD in large clinical studies, intravitreal anti-VEGF injection has replaced PDT as the primary treatment for PCV [29, 30]. Early studies reported that ranibizumab monotherapy showed an effective visual improvement [15], but the rate of polyp closure was low, ranging from 25 to 40% [16–18, 31]. In the EVEREST study comparing the efficacy of PDT monotherapy, ranibizumab monotherapy, and a PDT/ranibizumab combination therapy, the combination therapy and PDT monotherapy were superior to ranibizumab monotherapy in achieving complete regression of polyps at 6 months and 12 months, and improvement in BCVA at 12 months [19, 32]. However, in the LAPTOP study comparing the effects of PDT and ranibizumab monotherapy, ranibizumab monotherapy was better than PDT in PCV patients, and possible damage to the RPE and photoreceptors by PDT could account for the results [21]. Aflibercept binds not only to VEGF-A but also to VEGF-B and placental growth factor (PIGF) with greater affinity, and thus, it was expected to be more effective than bevacizumab (Avastin, Genentech, South San Francisco, CA, USA) or ranibizumab, both of which bind only to VEGF-A [33]. Cho et al. compared the effectiveness of intravitreal injection of aflibercept and ranibizumab for patients with PCV [34]. The mean number of injections for 1 year did not differ between the two groups (P = 0.152). After 12 months of injection, there was no significant difference between the groups concerning either BCVA improvement or decreased central foveal thickness, though polyp regression was significantly more frequent in the aflibercept-treated group.

The therapeutic effects of ranibizumab have been reported for eyes with good initial BCVA. In 2012, Saito et al. performed an average of 4.9 injections of intravitreal ranibizumab in 18 PCV eyes with a baseline BCVA exceeding 20/40, and reported a significant improvement in BCVA from the baseline (0.17 logMAR) to 12 months (0.07 logMAR) [35]. Mori et al. performed an average of 4.7 injections of intravitreal ranibizumab in 50 PCV eyes with a baseline BCVA of 0.6–1.0, and reported a significant improvement in BCVA from the baseline (0.11 logMAR) to 12 months (0.07 logMAR) [36]. Kato et al. administered intravitreal ranibizumab in PCV with baseline BCVA of 0.10 logMAR [37]. With 3.05 injections, the absolute value of visual acuity improved to 0.09, but there was no statistical significance.

In most studies of intravitreal aflibercept injections for PCV in Korea, the initial BCVA is less than 20/40, and eyes with good visual acuity of 20/40 or better are not included [20, 34, 38]. Cho et al. showed significant improvement in visual acuity at 12 months (0.44 logMAR) relative to the baseline (0.63 logMAR) after intravitreal aflibercept injections in 38 PCV eyes [34]. Kim et al. reported a retrospective review of 46 PCV eyes treated with intravitreal aflibercept monotherapy for 24 months [38]. An average of 4.7 injections was administered in the first 12 months, and 2.3 injections in the second 12 months. BCVA was significantly
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improved at 12 months relative to the baseline values. At 24 months there was no significant difference, but BCVA was maintained or improved in 76.1% of cases. In the VAULT study, Lee et al. administered 7 injections of intravitreal aflibercept in 40 PCV eyes for 1 year (at 2-month intervals after loading doses), and found significant improvement in visual acuity at 12 months (20/53) relative to the baseline (20/80) [20]. In Japan, Oshima et al. performed 2-month interval fixed treatments after loading doses of aflibercept in 50 PCV eyes [39]. BCVA had significantly improved at 12 month (0.12 logMAR) compared to the baseline (0.33 logMAR). However, no independent analysis of eyes with good initial vision was performed in any of the above reports.

The effects of intravitreal aflibercept injection in PCV patients with good visual acuity have been reported in Japanese studies. Minami et al. injected aflibercept in AMD patients with initial visual acuity greater than 0.6, and revealed that there was significant improvement in BCVA at 6 months (0.03 logMAR) and 12 months (0.04 logMAR) compared to baseline (0.07 logMAR) [40]. Of the 29 eyes, 17 PCV eyes were included, but no subgroup analysis was implemented for PCV. Sakamoto et al. injected aflibercept in AMD patients with initial visual acuity greater than 0.5 (typical AMD 18 eyes and PCV 21 eyes) [41]. While BCVA improved significantly at 12 months (0.06 logMAR) compared to baseline (0.10 logMAR) for the entire study group, PCV eyes showed a lack of significance compared to the baseline (0.07 logMAR) in 12 months (0.05 logMAR).

Although the above two studies included PCV with good visual acuity in the study population, the number of PCV patients was not large enough for subgroup analysis. This is different from our study, which was conducted exclusively with PCV patients.

In the present study, BCVA increased from 0.23 logMAR (at baseline) to 0.20 logMAR (12 months) with an average of 5.4 intravitreal aflibercept injections. However, the values were not significantly different relative to the baseline. In reports on the utilization of the pro re nata method, the number of injections of aflibercept and ranibizumab for PCV with poor visual acuity averaged, comparably to the present study, 4.7 to 4.9 over the course of 12 months [34–36, 38]. There was no significant difference in the number of injections in 12 months between studies on good-visual-acuity and poor-visual-acuity eyes. The reason seems to be that most studies are conducted based on 3 monthly loading injections, making it difficult to show difference between the remaining 9 months due to the recommended 2-month-interval administration. More long-term studies are needed for more meaningful comparisons.

Although this study did not show significant visual improvement at 12 months after treatment, the change of visual acuity was comparable to similar study by Mori et al. [36]. And whereas this study adopted the same pro re nata administration, no significant visual improvement was shown, unlike the reports of Saito et al’s and Mori
et al’s, which showed significant visual improvement by implementation of ranibizumab for PCV with good visual acuity [35, 36]. The reason seems to be baseline BCVA and follow-up intervals. In the current study, a subgroup analysis showed that final visual outcome was poor when baseline BCVA was worse than 0.2 logMAR, and the multiple linear regression analysis showed that final BCVA was associated with baseline BCVA. The average baseline BCVA of our study was 0.22 logMAR, while those of Saito et al. and Mori et al. were 0.14 and 0.11, respectively. Therefore, the fact that initial BCVA in this study was worse than that of the previous studies suggests that the disease was relatively in advanced state compared to the previous studies, and the prognosis after treatment may have been worse. As for follow-up intervals, both of those previous studies had follow-up examinations monthly, while in the current study, follow-up visits were scheduled only every 1 to 3 month. Therefore, recurrence could have been detected late, which would have resulted in delayed administration.

However, similar to current study, Sakamoto et al., who injected intravitreal aflibercept in PCV with good visual acuity, showed no significant improvement in BCVA at 12 months [41]. The mean change of logMAR BCVA they reported was similar to our study.

In earlier studies on PCV with poor visual acuity, the mean BCVA change at 12 months after treatment was 0.11~0.19 logMAR, which was greater than in the present study. This is judged to have been due to the ceiling effect, by which, in the current study, initial vision was relatively better and therefore had a lower potential for improvement. However, the final BCVA at 12 months after treatment was 0.20 logMAR, which was higher than in studies on eyes with poor initial visual acuity (0.42–0.45 logMAR) [20, 34, 38]. The size of lesions would be smaller, the activity of the disease would be less severe, damaged retinal tissues that cannot be recovered would be less prevalent in eyes with good visual acuity, and so, the final vision would be better. Table 3 summarizes the BCVA change at 12 months after intravitreal anti-VEGF treatment of previous studies and current study.

In the present study, visual acuity was maintained or improved in 94.2% of cases at 12 months after treatment, which is higher than the 76.1~87.5% reported in previous studies on eyes with poor initial vision [20, 34, 38]. The CSMT significantly decreased until 12 months after treatment without major changes, similarly to the study on ranibizumab injection for PCV with good visual acuity [36]. As for PED, it decreased significantly at 12 months after treatment but held at 69.8%. SRF showed the maximal improvement at 3 months after treatment and then increased again, but was improved significantly at all observation points. Lee et al. and Mori et al. also reported the greatest improvement of SRF at 3 months after treatment, presumably due to the effect of the 3 consecutive monthly injections, which protocol is common among all studies [20, 36]. After 3 loading doses in this study, the number of eyes with exudative changes was 18 (20.9%). This is similar to the values reported by
Mori et al. (20%, 10/50 eyes) who injected ranibizumab, and Lee et al. (23.8%) and Minami et al. (20%) who injected aflibercept [20, 36, 40].

Through subgroup analysis and multiple linear regression analysis, it was found that the factors affecting final BCVA were baseline BCVA and the presence of SRF at 3 months. Baseline CSMT did not affect the final BCVA. Previous studies have also shown that baseline BCVA independently affects post-treatment BCVA in PCV [16, 42, 43]. As mentioned above, good baseline BCVA indicates that the disease is relatively mild, so it is believed that the prognosis of visual acuity after treatment is expected to be good. The presence of SRF at 3 months means that the exudative change did not completely disappear even after three monthly loading doses, suggesting that the response to anti-VEGF treatment was poor.

The current study, due to its retrospective nature, did not strictly apply consistent retreatment criteria as typically are done in prospective studies. Moreover, since the sizes of PED and SRF were not measured, quantitative comparisons were not possible. Also, ICGA was not performed in all patients at 12 months after treatment; thus, polyp closure was not consistently determined across the patient, even though unclosed polyps are thought to be the main cause of recurrence.

However, to the authors’ best knowledge, this is the first long-term report of aflibercept treatment results for PCV with good visual acuity in Korea. In addition, it is a meaningful report of PCV sole disease excluding other diseases, and a reliable study with sample size of 86. In conclusion, intravitreal aflibercept injections decreased CSMT and were effective in maintaining visual acuity in cases of PCV with good visual acuity, and the baseline good visual acuity independently affected final visual acuity. Therefore, early treatment is necessary to maintain good visual acuity in PCV with good baseline BCVA.
References


43. Ho M, Lo EC, Young AL, Liu DT. Outcome of polypoidal choroidal vasculopathy at 1 year by combined 
   therapy of photodynamic therapy with ranibizumab and predictive factors governing the outcome. *Eye (Lond)* 
   2014;28:1469-76.
Table 1. Baseline characteristics of patients.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>65.6 ± 7.26 (52-81)</td>
</tr>
<tr>
<td>Gender (male : female)</td>
<td>60 : 26</td>
</tr>
<tr>
<td>BCVA (logMAR)</td>
<td>0.23 ± 0.09 (0.0-0.3)</td>
</tr>
<tr>
<td>CSMT (㎛)</td>
<td>336.1 ± 97.3 (139-597)</td>
</tr>
<tr>
<td>PED (%)</td>
<td>93.0</td>
</tr>
<tr>
<td>SRF (%)</td>
<td>91.9</td>
</tr>
</tbody>
</table>

BCVA = best-corrected visual acuity; CSMT = central subfield macular thickness; logMAR = logarithm of minimal angle of resolution; PED = pigment epithelial detachment; SRF = subretinal fluid.
Table 2. Multiple linear regression analysis for final visual acuity.

<table>
<thead>
<tr>
<th>Variables</th>
<th>$\beta$ coefficient</th>
<th>$P$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>0.098</td>
<td>0.357</td>
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<tr>
<td>Gender</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>0.202</td>
<td>0.052</td>
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<tr>
<td>Female (reference)</td>
<td></td>
<td></td>
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<tr>
<td>Baseline BCVA (logMAR)</td>
<td>0.213</td>
<td>0.047*</td>
</tr>
<tr>
<td>SRF at 3 month</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Absence (reference)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Presence</td>
<td>0.267</td>
<td>0.020*</td>
</tr>
<tr>
<td>Number of injections (n)</td>
<td>-0.088</td>
<td>0.448</td>
</tr>
</tbody>
</table>

* Statistically significant difference from the baseline ($P < 0.05$).

BCVA = best-corrected visual acuity; logMAR = logarithm of minimal angle of resolution; SRF = subretinal fluid.
<table>
<thead>
<tr>
<th>Study</th>
<th>Total sample size (PCV subgroup)</th>
<th>Ethnicity</th>
<th>Regimen</th>
<th>Drug</th>
<th>Only good baseline BCVA</th>
<th>Baseline BCVA (logMAR)</th>
<th>BCVA at 12 month (logMAR)</th>
<th>12-month BCVA change (logMAR)</th>
<th>No. of injection</th>
<th>Follow up interval (at PRN treatment)</th>
</tr>
</thead>
<tbody>
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<td>Saito et al, 2012 [35]</td>
<td>40 (18)</td>
<td>Japanese</td>
<td>3 loadings +</td>
<td>ranibizumab 0.5 mg</td>
<td>O</td>
<td>0.14</td>
<td>0.07*</td>
<td>0.07</td>
<td>4.9</td>
<td>1m</td>
</tr>
<tr>
<td>Mori et al, 2013 [36]</td>
<td>50 (50)</td>
<td>Japanese</td>
<td>3 loadings +</td>
<td>ranibizumab 0.5 mg</td>
<td>O</td>
<td>0.11</td>
<td>0.07*</td>
<td>0.04</td>
<td>4.72</td>
<td>1m</td>
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<tr>
<td>Kato et al, 2015 [37]</td>
<td>36 (17)</td>
<td>Japanese</td>
<td>PRN without</td>
<td>ranibizumab 0.5 mg</td>
<td>O</td>
<td>0.10</td>
<td>0.09</td>
<td>0.01</td>
<td>3.05</td>
<td>1m</td>
</tr>
<tr>
<td>Cho et al, 2016 [34]</td>
<td>98 (60)</td>
<td>Korean</td>
<td>3 loadings +</td>
<td>ranibizumab 0.5 mg</td>
<td>0.66</td>
<td>0.49*</td>
<td>0.17</td>
<td>5.52</td>
<td>1m</td>
<td>1m</td>
</tr>
<tr>
<td>Cho et al, 2016 [34]</td>
<td>98 (38)</td>
<td>Korean</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>0.63</td>
<td>0.44*</td>
<td>0.19</td>
<td>4.94</td>
<td>1m</td>
<td>1m</td>
</tr>
<tr>
<td>Lee et al, 2017 [20]</td>
<td>40 (40)</td>
<td>Korean</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>0.60</td>
<td>0.42*</td>
<td>0.18</td>
<td>7</td>
<td></td>
<td>1m</td>
</tr>
<tr>
<td>Oshima et al, 2017 [39]</td>
<td>50 (50)</td>
<td>Japanese</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>0.33</td>
<td>0.12*</td>
<td>0.21</td>
<td>7</td>
<td></td>
<td>1~2 m</td>
</tr>
<tr>
<td>Kim et al, 2018 [38]</td>
<td>46 (46)</td>
<td>Korean</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>0.56</td>
<td>0.45*</td>
<td>0.11</td>
<td>4.7</td>
<td></td>
<td>1~2 m</td>
</tr>
<tr>
<td>Sakamoto et al, 2018 [41]</td>
<td>39 (21)</td>
<td>Japanese</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>O</td>
<td>0.07</td>
<td>0.05</td>
<td>0.02</td>
<td>7</td>
<td>1~2 m</td>
</tr>
<tr>
<td>Current study</td>
<td>86 (86)</td>
<td>Korean</td>
<td>3 loadings +</td>
<td>aflibercept 2.0 mg</td>
<td>O</td>
<td>0.22</td>
<td>0.19</td>
<td>0.03</td>
<td>5.2</td>
<td>1~3 m</td>
</tr>
</tbody>
</table>

* Statistically significant difference from the baseline \( P < 0.05 \).

BCVA = best-corrected visual acuity; logMAR = logarithm of minimal angle of resolution; PCV = polypoidal choroidal vasculopathy; PRN = pro re nata.
Figure legends

Figure 1. Schematic flow chart of the treatment regimen using intravitreal aflibercept injections for the treatment of polypoidal choroidal vasculopathy.

Figure 2. Distribution of best-corrected visual acuity (BCVA) changes. There was no significant difference in the “improved” or “maintained” proportion between 3 months (97.7%) and 12 months (94.2%).

Figure 3. Mean changes in best-corrected visual acuity (BCVA) and central subfield macular thickness (CSMT) after intravitreal aflibercept injection. (a) Mean BCVA (logMAR) had changed from the baseline (0.23 ± 0.09) at 3 months (0.21 ± 0.14), 6 months (0.24 ± 0.22) and 12 months (0.20 ± 0.18), but with no statistical significance. (b) CSMT had improved significantly from the baseline (336.1 ± 97.3 μm) at 3 months (223.6 ± 47.22 μm, \( P < 0.001 \)), 6 months (239.6 ± 64.2 μm, \( P < 0.001 \)), and 12 months (223.8 ± 47.9 μm, \( P < 0.001 \)).

* Statistically significant difference from the baseline (\( P < 0.05 \)).

Figure 4. Proportions of eyes with pigment epithelial defect (PED), subretinal fluid (SRF), and intraretinal fluid (IRF). The proportions of PED, SRF, and IRF were significantly decreased at all observation points relative to the baseline.

* Statistically significant difference from the baseline (\( P < 0.05 \)).

Figure 5. Subgroup analysis according to baseline best-corrected visual acuity (BCVA), baseline central subfield macular thickness (CSMT), presence of subretinal fluid (SRF) at 3 month, and total injection numbers. (a) Eyes with baseline BCVA of 0.2 logMAR or better had significantly better final BCVA. (b) Stratification based on baseline CSMT of 300 μm showed a significant difference in BCVA only before treatment, and no significant difference was observed during the study period. (c) Eyes with SRF at 3 months had significantly worse final BCVA. (d) Stratification according to the number of injections showed no significant difference in BCVA during the entire study period.

* Statistically significant difference from the baseline (\( P < 0.05 \)).
Figure 6. A representative case of a 62-year-old man with polypoidal choroidal vasculopathy (PCV) treated with intravitreal aflibercept injections.

(a) Baseline color fundus photograph shows orange polypoidal lesion and large area of subretinal fluid (SRF). (b) Baseline fluorescein and indocyanine green angiography shows polyps and a network of abnormal vessels. (c) Baseline optical coherence tomography image shows large pigment epithelial detachment (PED) and SRF related to PCV. Visual acuity was 20/40. (d) After three monthly loading phase, SRF had resolved, but polypoidal lesions persisted. Visual acuity was 20/25. (e) At 1 year after the first treatment, the patient's visual acuity was 20/40. In total, 7 intravitreal aflibercept treatments were administered during the follow-up period.