Changes in the foveal microstructure after intravitreal bevacizumab application in patients with retinal vascular disease

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Purpose: To investigate changes in the area of the foveal avascular zone (FAZ) in patients with retinal vascular disease.

Patients and methods: This retrospective, consecutive study examined 53 eyes of 53 patients with macular edema due to branch retinal vein occlusion in 25 patients (47.2%) and nonproliferative diabetic retinopathy in 28 patients (52.8%). The macular edema was treated with an intravitreal injection of 0.05 mL equal to 1.25 mg bevacizumab. Before and 6–8 weeks after the injection, best corrected visual acuity, slit lamp biomicroscopy of the anterior segment and fundus, optical coherence tomography, and fluorescein angiography were conducted. The FAZ was manually circumscribed on early-phase angiography images and the area of the FAZ was measured.

Results: The preoperative overall mean FAZ area was \(0.327 \pm 0.126 \text{ mm}^2\) (median 0.310 mm\(^2\)). At the control consultation, the overall mean area was significantly larger (\(0.422 \pm 0.259 \text{ mm}^2\); median 0.380 mm\(^2\); \(P < 0.001\)). In the nonproliferative diabetic retinopathy subpopulation, the mean area was 0.361 \(\pm 0.129 \text{ mm}^2\) (median 0.330 mm\(^2\)) before bevacizumab application and 0.434 mm\(^2\) at the follow-up visit (mean increase 0.071 mm\(^2\)/19.7%). In the branch retinal vein occlusion group, the baseline FAZ area was 0.290 \(\pm 0.115 \text{ mm}^2\) and 0.407 \(\pm 0.350 \text{ mm}^2\) at follow-up (median 0.330 mm\(^2\); mean increase 0.117 mm\(^2\)/40.3%). No cases of severe operation-associated complications were observed.

Conclusion: The results confirm the safety of intravitreal bevacizumab injection in patients with macular edema due to nonproliferative diabetic retinopathy and branch retinal vein occlusion. The enlargement of the FAZ could be equivalent to an increase in retinal ischemia. These results may be transient; a potential vascular risk, however, when applying antivascular endothelial growth factor therapy in eyes with preexistent vascular disease must be considered.

Keywords: foveal avascular zone, ischemia, diabetic retinopathy, branch retinal vein occlusion, anti-VEGF

Introduction
Management of macular edema following branch retinal vein occlusion (BRVO) and nonproliferative diabetic retinopathy (NPDR) is challenging. Except photocoagulation, which used to be the initial therapy of choice, the pharmacologic inhibition of vascular endothelial growth factor (VEGF) shows promise.\(^1,2\) In the meantime, ranibizumab has become the therapy of choice for BRVO with macular edema and a visual acuity of 20/40 and less.

VEGF is a multifunctional cytokine. It is not only crucial for cell proliferation, cell migration, proteolysis, cell survival, and maintenance of the choriocapillaris,
it also facilitates survival and development of retinal neurons following ischemia and it is necessary for preservation of normal vessels.\textsuperscript{3–7} Its role in angiogenesis and organogenesis is vital; the loss of one single VEGF allele results in embryonic death.\textsuperscript{8} One of its major inducers is hypoxia.\textsuperscript{9} Therefore, it plays an important role in ocular diseases in which retinal malperfusion is developing, eg, diabetic macular disease or BRVO.\textsuperscript{10} Furthermore, VEGF stimulates endothelial mitogenesis, promotes endothelial survival, and controls vascular permeability. It is 50,000 times more potent in inducing vascular leakage than histamine, causing macular edema when released in the vitreous cavity of monkeys.\textsuperscript{11,12} Development of macular edema causes impairment of visual acuity in these pathological conditions.

There is great evidence from large prospective randomized trials for the efficacy of anti-VEGF therapy in diabetic macular edema (DME), macular edema following BRVO, and age-related macular degeneration.\textsuperscript{13–22} A novel agent, aflibercept (Eylea\textsuperscript{8}, Regeneron Pharmaceuticals, Inc, Tarrytown, NY, USA), now augments the pharmacological anti-VEGF armamentarium.\textsuperscript{23} This soluble decoy receptor shows a higher affinity and faster association to VEGF-A than ranibizumab or bevacizumab.\textsuperscript{24}

This study presents two groups of patients who were diagnosed with macular disease secondary to DME or BRVO and consequently treated with a single dose of intravitreal bevacizumab. Both groups demonstrated a significantly enlarged foveal avascular zone (FAZ) at the 6–8-week follow-up.

Material and methods
In this retrospective consecutive study, more than 200 patients with NPDR or BRVO who visited the Department of Ophthalmology of Technical University of Munich (Munich, Germany) between June 2008 and December 2009 were screened, and 53 patients were included. These 53 individuals suffered from macular edema due to NPDR or BRVO and also had a well-circumscribed FAZ, both visible on fluorescein angiography. The study was approved by the local institutional review board (Ethics Commission, Faculty of Medicine, Technical University of Munich) and informed consent was obtained from every patient for the intravitreal injection. The subjects were also provided information about the off-label use of the drug.

The mean age of the patients was 67.4 years; 31 female (58.5%) and 22 male (41.5%) patients were included. The macular edema was caused by NPDR in 28 patients (52.8%) and by BRVO within the last 6 months in 25 patients (47.2%). The exclusion criteria were macular disease secondary to causes other than DME or BRVO, any ongoing ocular or periocular inflammation, any kind of ocular surgeries on the studied eye within the last 6 months, or any previous treatment of the macular edema (eg, any intravitreal injection for macular edema in the last 6 months or laser treatment at any time in the past for macular edema).

Examinations at baseline included best corrected visual acuity with a Snellen chart at a distance of 20 feet, slit lamp biomicroscopy of the anterior segment and fundus (Slit Lamp BM 900\textsuperscript{e}; Haag-Streit AG, Koeniz, Switzerland), optical coherence tomodraphy (Spectralis\textsuperscript{8}; Heidelberg Engineering, Heidelberg, Germany), and fluorescein angiography (Heidelberg Retina Angiograph II; Heidelberg Engineering).

The intravitreal injection of 0.05 mL with 1.25 mg bevacizumab was performed in accordance with the guidelines for intravitreal injections published by the Macula Committee of the German Ophthalmological Society and the Professional Association of Ophthalmologists of Germany.\textsuperscript{25} Before injection administration, the eye was washed with povidone-iodine (5%) and the eyelashes and lid region were then wiped, also with povidone-iodine (5%). The drape (as used for intraocular intervention) was applied, a lid speculum was inserted. After dislocation of the conjunctivae, the injection was performed at a distance of 3.5 mm from the limbus. Consecutively, each patient received an injection of 1.25 mg bevacizumab (Avastin\textsuperscript{8}; F Hoffmann-La Roche, Basel, Switzerland). Antibiotic eye drops were then applied. At day one and day three after the injection, a slit lamp examination was performed to rule out intraocular inflammation or elevated intraocular pressure.

In order to visualize the FAZ, the angiography was performed with scanning laser technology, and only patients with good quality early-phase fluorescein angiograms without any bleeds or exudates hindering visualization of the macular region were included for evaluation. The FAZ was manually circumscribed by two experienced retina consultants in these early-phase fluorescein angiograms, once by each of them (Figure 1). Heidelberg Eye Explorer software (version 1.7.0.0; Heidelberg Engineering) yielded the area (in mm$^2$). The mean value was used for further calculations and patients with differing values of more than 0.05 mm$^2$ were excluded.

Patients returned to the outpatient clinic for routine postinjection follow-up. Best corrected visual acuity, slit lamp biomicroscopy of the anterior segment and fundus, optical coherence tomography, and fluorescein angiography
were conducted again at 6–8 weeks postinjection. The area of FAZ was assessed again in the above described manner. Statistical analysis was performed using SPSS version 13 (SPSS Inc, Chicago, IL, USA) with the Wilcoxon matched-pairs test and t-test.

Results
All patients tolerated the injection well. There were no cases of operation-associated complications such as retinal detachment, endophthalmitis, or persistent elevated intraocular pressure. Two patients reported ocular discomfort for a few days, and seven patients (13.2%) showed a mild hyposphagma at the injection site that disappeared without further treatment within 10 days. The mean FAZ was significantly larger at the control visit after 6–8 weeks compared to the previous values for all patients (Wilcoxon: \( P < 0.001 \)). Preoperative parameters showed a mean area of 0.327 ± 0.126 mm\(^2\) (median 0.310 mm\(^2\)). After the first visit, the mean area increased to 0.422 ± 0.259 mm\(^2\) (median 0.407 ± 0.350 mm\(^2\); Table 1; Figure 2).

An analysis to differentiate between the underlying diagnoses was also performed. In the NPDR group (n = 28), the baseline mean FAZ area was 0.361 ± 0.129 mm\(^2\) (median 0.330 mm\(^2\)). At the follow-up, the mean area increased by 0.071 mm\(^2\) (or 19.7%) to 0.434 mm\(^2\) (Table 1; Figure 2).

The subjects with BRVO (n = 25) showed a slightly smaller FAZ area at baseline compared to the diabetic group (0.290 ± 0.115 mm\(^2\) versus 0.361 ± 0.129 mm\(^2\)). At the follow-up, the mean area increased by 0.117 mm\(^2\) (or 40.3%) to 0.407 ± 0.350 mm\(^2\) (median 0.330 mm\(^2\); Table 1; Figure 2).

Discussion
In February 2004, the US Food and Drug Administration approved bevacizumab as the first angiogenesis inhibitor for first-line treatment of cancer; and favorable results led to extended use in non-small-cell lung cancer, kidney cancer, and glioblastoma. As a nonselective inhibitor of all isoforms of the VEGF, this humanized antibody also suppresses vital physiological functions like its vasodilatory effect or the stimulated expression of plasminogen activators, resulting in arterial thromboembolic events.\(^{26-31}\) Ultrastructural analyses showed the reduction of choriocapillaris endothelial cell fenestrations and emerging thrombosis.\(^{32-34}\) Evidence for vasoconstriction of retinal vessels was published, altogether raising concerns about retinal vascular events.\(^{35-37}\)

This retrospective study found a significant increase in the area of the FAZ 6–8 weeks after a single intravitreal injection of 1.25 mg bevacizumab. There were no significant complications after intravitreal bevacizumab, including endophthalmitis, retinal detachment, cataract formation, increased intraocular pressure, or retinal artery occlusion. The presented results are in accordance with previous observations.\(^{38}\) Several multicenter clinical trials also did not detect any elevated incidence of severe side effects in anti-VEGF therapy; and reports on increased intraocular pressure and ischemia are seldom, with no adverse event rate exceeding 0.21%.\(^{39-45}\) Mansour et al determined the overall risk of ocular vascular events following VEGF antagonist injection as 0.108% in a general population and 2.61% in diabetic patients, and refer to the patients’ perioperative stress and the natural history of their conditions.\(^{46}\)

There are several publications that attempt to measure retinal ischemia on fluorescence angiography as the size of the FAZ. Neubauer et al used a high-resolution ultra wide-field scanning ophthalmoscopy system for counting retinal fields of nonperfusion in a group of 19 nonresponders to photocoagulation.\(^{47}\) The number of avascular fields

![Figure 1](https://www.dovepress.com/)

Figure 1 The foveal avascular zone was manually circumscribed on an early-phase fluorescein angiogram and the software measured the area (in mm\(^2\)).

### Table 1 Foveal avascular zone area measurements

<table>
<thead>
<tr>
<th>Foveal avascular zone (mm(^2))</th>
<th>DME</th>
<th>BRVO</th>
<th>Overall</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pre OP</td>
<td>FU1</td>
<td>Pre OP</td>
</tr>
<tr>
<td>Mean</td>
<td>0.36</td>
<td>0.43</td>
<td>0.29</td>
</tr>
<tr>
<td>Median</td>
<td>0.33</td>
<td>0.42</td>
<td>0.28</td>
</tr>
<tr>
<td>First quartile</td>
<td>0.26</td>
<td>0.34</td>
<td>0.21</td>
</tr>
<tr>
<td>Third quartile</td>
<td>0.46</td>
<td>0.51</td>
<td>0.36</td>
</tr>
<tr>
<td>Minimum</td>
<td>0.14</td>
<td>0.16</td>
<td>0.04</td>
</tr>
<tr>
<td>Maximum</td>
<td>0.63</td>
<td>0.71</td>
<td>0.62</td>
</tr>
<tr>
<td>Standard deviation</td>
<td>0.13</td>
<td>0.14</td>
<td>0.11</td>
</tr>
</tbody>
</table>

Abbreviations: BRVO, branch retinal vein occlusion; DME, diabetic macular edema; FU1, first follow-up visit; Pre OP, before injection of bevacizumab.
dropped significantly, but the diameter of the FAZ remained quasi unchanged, as it did in a prospective, consecutive, noncomparative case series with 126 diabetic pretreated patients at a 12-month follow-up.14 Another group measured an area of capillary nonperfusion in the context of intravitreal bevacizumab for macular edema following BRVO; they found an unchanged mean area of nonperfusion.48 One reason for this could be the slightly different period of time between intervention and the second data acquisition. Nevertheless, the current study yielded different results.

A statistically significant enlargement of the FAZ 6 to 8 weeks after one intravitreal injection of 1.25 mg bevacizumab in patients with macular edema both secondary to NPDR and BRVO was found. This effect may be transient. Papadopoulou et al found a vasoconstrictor impact of ranibizumab on retinal arteries after application of ranibizumab, suggesting that closure of the smallest vessels approaching the FAZ could enlarge this area on fluorescein angiography.35 A recent report defined the FAZ as a factor of macular ischemia and found a negative effect of an enlarged FAZ on short-term visual outcomes after intravitreal bevacizumab for DME.49

Some limitations are inherent in the current study. First, the design was retrospective, the sample size was small (both groups totaled 53 eyes), and there was no control group. As the natural course of the disease also leads to increasing vascular stress and enlargement of the FAZ, it is not known without doubt that the results are secondary to the use of bevacizumab. Second, this one-time follow-up of the patients does not describe the development of the FAZ over time and the results may be transient, especially since the effects of intravitreal bevacizumab have been reported to vanish within weeks after a single application.50 Third, the period from BRVO or the onset of diabetes to the application of bevacizumab was not defined. It is likely that different durations of ischemia yield a varying response to anti-VEGF treatment.

**Conclusion**

The results show that intravitreal injection of anti-VEGF drugs may induce an increase in the area of the FAZ, a morphological retinal change visualized by fluorescein angiography. In the light of retinal ischemia, this may cause further problems. Altogether, the findings emphasize only one single aspect of intravitreal anti-VEGF treatment in patients with preexistent vascular disease; they underlie the necessity to consider the potential vascular risk when applying anti-VEGF therapy in eyes with preexistent vascular disease, which is common in patients with DME or BRVO. A standardized method to assess the FAZ and further investigations are necessary to describe the relationship between intravitreal bevacizumab, microvascular changes in the FAZ, and long-term visual acuity results.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


![Figure 2 Box plots summarize the increase in foveal avascular zone area (mm²) at FU1 compared to pre OP measurements for all patients (overall) and the patients with DME and BRVO. Abbreviations: BRVO, branch retinal vein occlusion; DME, diabetic macular edema; FU1, first follow-up visit; Pre OP, before injection of bevacizumab.](https://www.dovepress.com/fig2-bpx2.png)


