Introduction

Cotton wool spots (CWS) are not specific for a disease, but rather a sign of vascular damage [1]. Ischemia due to vasoconstriction or occlusive events as well as mechanical pressure affecting the superficial capillary network located in the retinal nerve fiber layer (RNFL) may induce CWS. In turn, ischemic events of the intermediate and deep capillary network may induce paracentral middle maculopathy and acute macular neuroretinopathy. Histologically, CWS consist of accumulated cystoid bodies, which represent the terminal swelling of a disrupted ganglion cell axon with accumulated axonal debris [2, 3].
CWS reveal a characteristic presentation on each image device. On color fundus photographs (CF) or by ophthalmoscopy, CWS appear as fluffy white lesions with frayed edges [1]. In fluorescein angiography (FA), 90% of the lesions appear hyperfluorescent and the remaining 10% appear hypofluorescent in the early phase and become hyperfluorescent in the late phase [4]. In optical coherence tomography (OCT), CWS appear as hyperreflective lesions in the retinal RNFL [5]. On average, CWS disappear within 7–12 weeks from ophthalmoscopic view, although they have been found to be still visible on OCT after vanishing from CF images [5–7]. After resolution on OCT, CWS often leave ganglion cell layer, RNFL, and inner nuclear layer thinning, and functionally CWS resolution is associated with persistent relative scotomas [8, 9].

CWS have been methodically studied in HIV, diabetic macular edema, central retinal vein occlusion and hypertension, but so far not in branch retinal vein occlusion (BRVO) [8–13]. The aim of this study was to assess and describe CWS in BRVO in a longitudinal setting over 3 years using multimodal imaging.

Methods

Patient Selection and Setting

30 eyes of 30 patients with new-onset BRVO, who were recruited for a prospective nonrandomized study at the Department of Ophthalmology, Medical University Vienna, were evaluated [14]. The study adhered to the tenets of the Declaration of Helsinki, was approved by the Ethics Committee of the Medical University of Vienna and all patients gave informed consent. Inclusion criteria were the confirmed diagnosis of treatment-naïve acute BRVO with a decrease of visual acuity and a central retinal thickness >300 μm. Patients received one intravitreal ranibizumab (Lucentis; Novartis, Basel, Switzerland) injection at baseline, followed by an ‘as needed’ regimen (treatment was repeated when a circumscribed, whitish, fluffy lesion with frayed edges was visible hyperfluorescent in the late phase [49]. In optical coherence tomography (OCT), CWS appear as hyperreflective lesions corresponding to the hyperreflective RNFL thickening in SD-OCT or the fluffy whitish lesion on CF images was present. A CWS was graded as present on IR images when a well-demarcated hypo/hyperreflective roundish lesion corresponding to the hypo/hyperreflective roundish lesion corresponding to the hyperreflective RNFL thickening in SD-OCT or the fluffy whitish lesion on CF images was present. A CWS was graded as present on FA when a well-demarcated hypo/hyperreflective roundish lesion corresponding to the hyperreflective RNFL thickening in SD-OCT or the fluffy whitish lesion on CF images was present. A CWS was graded as present on FA when a well-demarcated hypo/hyperreflective roundish lesion corresponding to the hyperreflective RNFL thickening in SD-OCT or the fluffy whitish lesion on CF images was present. A CWS was graded as present on FA when a well-demarcated hypo/hyperreflective roundish lesion corresponding to the hyperreflective RNFL thickening in SD-OCT or the fluffy whitish lesion on CF images was present.

Statistical Analyses

Statistical analyses were performed using SPSS 21 (SPSS Inc., Chicago, III., USA) software. Data were evaluated descriptively and numeric values were indicated as the mean ± SD. Differences in terms of size were evaluated with a paired t test. Factors associated with the duration of visibility of CWS were analyzed using a multivariate regression analysis. p ≤ 0.05 was deemed statistically significant.
Results

Demographic Characteristics

Out of the 30 patients, 24 patients (12 men, 12 women, mean age: 66 ± 11 years, 4 inferotemporal and 20 superotemporal BRVOs) presented with BRVO with an onset ≤8 weeks before the screening visit and were included in the study for further evaluation. Out of these 24 eyes, 18 eyes (75%) showed one or more CWS within the raster scan area (mean: 1.8 ± 1.3 CWS, range: 1–5) per eye. Overall, 29 CWS lesions within the SD-OCT scan area were found and analyzed at baseline. During the 3-year observational period, 3 additional CWS appeared in 3 different patients. A representative example of the presentation of a CWS in respective imaging devices can be found in figure 1. Figure 2 shows the corresponding marked areas. The average number of ranibizumab injections per patient was 4.5 ± 3.7 (range: 1–13) over the 3-year period. Out of the 24 patients, 6 (5 presenting with CWS, 1 not) received segmental argon laser coagulation laser treatment (mean: 1.7 ± 0.7, range: 1–3) during the observational period.

Morphologic and Angiographic Appearance of CWS

All 29 CWS (100%) were visible on SD-OCT, but only 19 (65.5%) were visible on the CF images at baseline. 25 (86.2%) of the CWS lesions showed corresponding changes on the IR image. All 29 (100%) CWS lesions showed corresponding fluorescent changes on FA: 9 (31%)
showed hypofluorescent lesions, 4 (13.8%) hyperfluorescent lesions, and the majority, 16 (55.2%), hypofluorescent changes at 2 min and hyperfluorescent changes at 10 min. At baseline, 27 (92.1%) CWS were surrounded by blood visible on CF images, 23 (78.3%) by intraretinal cysts visible on OCT, 2 (6.9%) by subretinal fluid and 1 by an avascular zone visible on FA.

CWS on FA could change their fluorescent behavior on FA as the fluorescence seemed to be associated with the surrounding pathology: CWS in the near vicinity of intraretinal cysts were mainly hyperfluorescent (88%), CWS surrounded by intraretinal hemorrhage were either hypofluorescent (41%) or hypofluorescent in the early phase and hyperfluorescent in late phase (44%), and CWS within an avascular zone presented mostly (83%) as hypofluorescent on FA.

Size and Duration of Visibility
The mean area of CWS for each imaging method is shown in table 1. The mean area of a CWS in SD-OCT was larger than on the CF image (paired t test: 0.26 ± 0.17 mm² vs. 0.13 ± 0.1, p ≤ 0.0001). The CWS area evaluated by the OCT-Tool-Kit software on SD-OCT (but not on CF), and surrounding pathology such as intraretinal cysts, avascular zones and intraretinal hemorrhage were predictive for how long a CWS remained visible (r² =

Fig. 2. The different marked areas of the CWS in SD-OCT (corresponding to fig. 1), FA and CF evaluated by the OCT-Tool Kit software. CWS were marked in each OCT B-scan (a), and the marked area was then projected en-face on the IR picture (b) and the affected area was automatically calculated by interpolation. The CWS (arrow) was 0.14 mm² in SD-OCT (c) 0.09 mm² in CF (arrow) (d) and the corresponding CWS area in FA revealed 0.17 mm².
Large CWS surrounded by intraretinal hemorrhage were visible longer than smaller CWS in the near vicinity of intraretinal cysts and avascular zones.

CWS were visible for 12.4 ± 7.5 months (range: 1–30) on SD-OCT, but only for 4.4 ± 3.2 months (range: 1–12) on CF images. Representative examples can be found in figures 3, 4 and 5. Corresponding CWS lesions on IR images were visible for 4.3 ± 3.4 months (range: 1–12) while corresponding fluorescent changes on FA were visible longer than the CWS itself on SD-OCT (mean: 17.5 ± 7.1 months, range: 3–27). However, after the 15-month follow-up, the margins of the CWS on FA became very indistinct and the lesions were only discernable from their surroundings with the use of an overlay.

**Discussion**

To the best of our knowledge this is the first study to evaluate the appearance of CWS in a very homogeneous group of patients with BRVO using four different imaging modalities in a longitudinal manner. Although CWS have been described in different diseases, including CRVO, only a few case reports have addressed the appearance and behavior of CWSs in BRVO so far [13, 17]. The current study reveals that SD-OCT aims to be a sensitive tool to detect small CWS even when they are invisible on CF, and it demonstrates that the duration of visibility of CWS seems to differ dependent on image modality and appears to be associated with the size of a CWS and with the surrounding pathology.

Currently, the main clinical image modality to observe CWS is CF photography. This study, however, shows that only 65.5% of all examined CWS were detectable on CF, whereas all CWS were visible on SD-OCT and FA. Furthermore, the area of a CWS was larger on SD-OCT compared to CF. Previous studies have already demonstrated that CWS are visible longer on OCT than by ophthalmoscopy [2, 9]. The higher prevalence of CWS on SD-OCT may therefore be a result of the longer duration of visibility. It may also be possible that a CWS has to reach a certain size to be clearly visible by ophthalmoscopy.

Many previous studies have evaluated the mean ‘lifetime’ of CWS in funduscopy in different kinds of retinal diseases. These studies showed they were visible anywhere from 4 weeks to 17.2 months depending on underlying disease, patient’s age and medication [12, 13, 18]. Besides these factors, our analysis revealed that the area of CWS as well as the surrounding pathology such as intraretinal cysts, intraretinal hemorrhage and avascular zones seem to be factors associated with the duration of visibility. On average, CWS were visible on CF images for 4.4 months in our study, which is right in the middle of previous findings. However, CWS were visible three times longer in SD-OCT than on CF images and IR, with a mean of 12 months. Previous studies have already shown that CWS are visible longer on OCT than on CF images and that the hyperreflectivity of the CWS lesions decreases gradually on OCT [5, 9]. Similar to CF, the presence of CWS was shorter on IR than on OCT.

CWS result from ischemic events at the level of the superficial capillary network and present as hyporeflective lesions on IR and as localized hyperreflectivity and thickening of the RNFL on OCT. This presentation corresponds to the appearance of acute macular neuroretinopathy and paracentral middle maculopathy lesions on IR and SD-OCT, which are thought to be induced by vasoocclusive events at the level of the intermediate and deep retinal capillary network, and are therefore sometimes reported as ‘deep CWS’ [19–21]. However, unlike the current finding in CWS that the hyperreflectivity on OCT was visible longer than the hyporeflectivity on IR, acute macular neuroretinopathy and paracentral middle maculopathy lesions were reported to diminish simultaneously in respective imaging devices [19–22]. This discrepancy may be due to the fact that the CWS we examined were mainly found near other pathologies such as cysts, avascular zones or intraretinal hemorrhage, which may

<table>
<thead>
<tr>
<th>Visits, months</th>
<th>SD-OCT, μm²</th>
<th>CF, μm²</th>
<th>FA, μm²</th>
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</thead>
<tbody>
<tr>
<td>Baseline</td>
<td>0.22±0.16</td>
<td>0.12±0.09</td>
<td>0.27±0.29</td>
</tr>
<tr>
<td>3</td>
<td>0.19±0.13</td>
<td>0.08±0.05</td>
<td>0.25±0.21</td>
</tr>
<tr>
<td>6</td>
<td>0.18±1.9</td>
<td>0.17±0.12</td>
<td>0.3±0.22</td>
</tr>
<tr>
<td>9</td>
<td>0.09±0.07</td>
<td>0.06</td>
<td>0.12±0.1</td>
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<tr>
<td>12</td>
<td>0.08±0.03</td>
<td>0.03</td>
<td>0.12±0.7</td>
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<tr>
<td>15</td>
<td>0.08±0.07</td>
<td>0.03</td>
<td>0.29±0.47</td>
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<tr>
<td>18</td>
<td>0.09±0.06</td>
<td>n.a.</td>
<td>n.a.</td>
</tr>
<tr>
<td>21</td>
<td>0.06±0.04</td>
<td>n.a.</td>
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<tr>
<td>24</td>
<td>0.06±0.04</td>
<td>n.a.</td>
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<tr>
<td>27</td>
<td>0.1±0.06</td>
<td>n.a.</td>
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<tr>
<td>30</td>
<td>0.1</td>
<td>n.a.</td>
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<tr>
<td>33</td>
<td>n.a.</td>
<td>n.a.</td>
<td>n.a.</td>
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<tr>
<td>36</td>
<td>n.a.</td>
<td>n.a.</td>
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</tr>
</tbody>
</table>

Values are given as means ± SD. Area of CWS on FA was evaluated until month 15. After that, the margins of CWS on FA were too blurry to allow accurate area evaluation.
aggravate the evaluation of the CWS in different image devices and alter their presentation and behavior.

Interestingly, CWS remained visible for $17.5 \pm 7.1$ months on FA, longer than on SD-OCT. Previous studies have described superficial capillary nonperfusion with vascular leakage and staining in the late phase to correspond to the CWS on FA in the acute phase [21]. In the chronic phase, CWS were mainly described as invisible on FA or detectable as an indistinct hypofluorescent area, visible due to the leakage from the surrounding perfused capillaries [21]. In our study, CWS were detectable on FA longer than on SD-OCT. It has to be taken into account, however, that hyperreflectivity and thickening of the RNFL, but not the ensuing localized atrophy of the RNFL, was graded as CWS on SD-OCT and that during follow-up the margins of the CWS on
FA became very indistinct and the lesion was only discernable from the surroundings when an overlay was used. It seems reasonable that the affected area remains distinguishable for a long period of time on FA as it resembles an area of capillary nonperfusion which remains nonperfused [21].

Indeed, the surrounding pathology together with the size of a CWS has an impact on the duration of visibility. Large CWS surrounded by intraretinal hemorrhage were visible longer than smaller CWS close to intraretinal cysts or an ischemic zone. The surrounding pathology not only affected the duration of visibility, but also influenced the fluorescent behavior of a CWS. In previous reports, CWS in FA were mainly described as hyperfluorescent lesions with late staining and leakage around their margins [1, 10, 13, 23]. Lesions that were hypofluorescent in the early

**Fig. 4.** Another example of the presentation of a CWS in CF (a–c) and SD-OCT (d–f). 

a Baseline (area: 0.22 mm²).  
b 6 months after initial presentation, the CWS has faded but is still visible (area: 0.1 mm²).  
c 12 months after initial presentation, the CWS is no longer visible in CF.  
d Axoplasmic debris in the RNFL at baseline (area: 0.3 mm²).  
e After 6 months: besides the atrophy of the inner retinal layers, a hyperreflective sign is still visible (area: 0.13 mm²).  
f After 12 months: concordant to the CF-images, the hyperreflective sign faded and the atrophy of the inner retinal layers remains.
phase but became hyperfluorescent in the late phase were suggested to be a sign of ischemia and were observed in a study with experimental retinal ischemia [4]. In the current study, CWS were either hyperfluorescent or hypofluorescent in the early and late phases or hypofluorescent in the early phase and hyperfluorescent in the late phase. Their fluorescent behavior changed over time and was associated with the surrounding pathology: CWS surrounded by an avascular zone or by intraretinal hemorrhage presented as hypofluorescent and either remained hypofluorescent or became hyperfluorescent in the late phase due to staining and leakage. CWS surrounded by intraretinal cysts and edema, however, mainly presented as hyperfluorescent in the early and late phases. The discrepancies between the presentation of CWS on FA in experimental retinal ischemia and in BRVO in the current study may be due to different findings in the area surrounding a CWS or due to different underlying pathologies. However, both have in common that hypofluorescent rather than pure hyperfluorescent CWS lesions seem to be a sign of hypoperfusion and ischemia. Intraretinal hemorrhage and focal ischemia were pathologies often described in close vicinity to CWS [23, 24], and since intraretinal hemorrhage is associated with local ischemia and often results in inner retinal atrophy, intraretinal hemorrhage could therefore be a concrete sign of the presence and the development of an avascular zone with similar results as an avascular zone in a hypofluorescent lesion.

In summary, this study indicates that the ’lifetime’ of CWS in BRVO seems comparable to other diseases and that the duration of visibility of CWS varies among dif-

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Fig. 5. Another example of the presentation of a CWS in CF (a–c) and SD-OCT (d–f).

- **a** Baseline (area: 0.22 mm²).
- **b** 6 months after initial presentation: the CWS area is even bigger compared to baseline (area: 0.28 mm²).
- **c** 12 months after initial presentation: CWS is no longer visible in CF.
- **d** Axoplasmic debris in RNFL at baseline (area: 0.51 mm²).
- **e** 6 months: the CWS area has increased in size (area: 0.79 mm²).
- **f** 18 months: a hyperreflective sign is still visible (area: 0.17 mm²).
different image modalities and depends on the surrounding pathology and the size of the CWS. Overall, SD-OCT seems to be a precise tool to detect, measure and observe central CWS, and it may be more sensitive in the detection of a CWS than CF.

Disclosure Statement

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References