



OPEN ACCESS

Short-term real-world outcomes following intravitreal brolocizumab for neovascular AMD: SHIFT study

Louisa Maria Bulirsch,¹ Marlene Saßmannshausen,¹ Jennifer Nadal,² Raffael Liegl,¹ Sarah Thiele ,¹ Frank G Holz¹

¹Department of Ophthalmology, University of Bonn, Bonn, Germany

²Institute for Medical Biometry, Informatics and Epidemiology, University of Bonn, Bonn, Germany

Correspondence to

Professor Frank G Holz, Department of Ophthalmology, University of Bonn, 53127 Bonn, Germany; Frank.Holz@ukbonn.de

ST and FGH contributed equally.

Received 14 January 2021

Revised 2 March 2021

Accepted 25 March 2021

ABSTRACT

Background Brolocizumab has recently been approved in Europe as a novel treatment for patients with neovascular age-related macular degeneration (nAMD). We report on early experiences with real-world outcomes of switch to brolocizumab therapy in previously anti-vascular endothelial growth factor (anti-VEGF)-treated patients.

Methods Patients with recalcitrant nAMD were switched to brolocizumab therapy. Functional and structural parameters 4 weeks after first brolocizumab injection were evaluated including best-corrected visual acuity (BCVA (logMAR)), foveal centre point (FCP (μm)), central subfield retinal thickness (CSRT (μm)) and macular volume (mm^3).

Results Sixty-three eyes of 57 patients with nAMD (52.6% females) with a mean (\pm SD) age of 79.5 ± 6.7 years were included. Mean change of BCVA was -0.03 ± 0.14 logMAR ($p=0.115$). Significant reductions were recorded for FCP with a mean (\pm SD) change of -66.81 ± 72.643 μm , -66.76 ± 60.71 μm for CSRT and -0.27 ± 0.24 mm^3 for macular volume (all $p<0.001$). Intraocular inflammation was observed in seven eyes of seven patients, including one case of retinal vasculitis without occlusion.

Conclusions The results of the SHIFT study indicate that switch to brolocizumab may represent a treatment option in patients with nAMD poorly responsive to other anti-VEGF agents. Further long-term analyses appear prudent to assess efficacy and safety of brolocizumab in a routine clinical setting.

INTRODUCTION

Age-related macular degeneration (AMD) is the leading cause of blindness in the elderly in industrialised countries.¹ With the advent of anti-vascular endothelial growth factor (VEGF) therapy, the visual outcome of patients with neovascular AMD (nAMD) has been improved and measurable reductions of legal blindness incidence have emerged.^{2,3}

Besides the anticipated worldwide increase of AMD prevalence due to demographical changes with longer life expectancy, the burden for both patients and caregivers is high when managing patients with repetitive intravitreal injections and monitoring visits over a long period of time in a chronic disease. Various real-world studies have shown visual outcomes to be inferior compared with the results from prospective randomised clinical trials (RCTs).⁴ Undertreatment is one of the major factors in part driven by non-adherence. In addition, some patients and certain subphenotypes

of nAMD do not respond favourably.^{5–12} Therefore, more efficacious agents with longer durability represent an important unmet need.

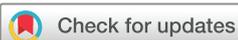
Brolocizumab (Novartis), a single-chain antibody fragment, was recently approved for the treatment of nAMD in October 2019 and in February 2020 by the regulatory agencies in the USA and the European Union, respectively, as well as in other countries.¹³ Potential benefits of brolocizumab are assumed to be related to its low molecular weight with subsequent better tissue penetration as well as higher molar concentration.^{13–15} Two pivotal trials have recently shown non-inferiority of brolocizumab to the comparator aflibercept with regard to visual outcome.¹³ Post hoc analyses demonstrated overall favourable anatomical effects.^{16–18} However, safety signals have been reported in both RCTs and post-marketing reports, which included the occurrence of intraocular inflammation (IOI) and retinal vasculitis with or without occlusion.^{13,18–25}

The aim of the SHIFT study was to report early real-world experiences in a single European clinical centre of brolocizumab treatment for nAMD with regard to both functional and anatomical disease control as well as adverse effects following approval in February 2020 in Europe.

METHODS

The SHIFT study is a retrospective, observational, monocentre study of patients with exudative AMD who received 6 mg brolocizumab intravitreal therapy between 16 March 2020 and 15 October 2020, at the Department of Ophthalmology, University of Bonn, Germany, in routine clinical care. All patients were previously treated repetitively because of recalcitrant fluid accumulations on optical coherence tomography (OCT) despite frequent dosing with other anti-VEGF agents, including ranibizumab, aflibercept and bevacizumab. Recalcitrant fluid was defined as persistent fluid accumulations despite a high frequency of intravitreal injections of other anti-VEGF agents over a longer period of time prior to the switch to brolocizumab. The day of the first intravitreal brolocizumab injection was regarded as the baseline visit.

At each visit, best-corrected visual acuity (BCVA) determination and complete ophthalmic examination, including slit-lamp examination and funduscopy following pupil dilation, was performed. Signs of IOI and/or retinal vasculitis were recorded if present. Retinal imaging was performed at each visit with combined confocal scanning laser ophthalmoscopy and spectral-domain optical coherence



© Author(s) (or their employer(s)) 2021. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

To cite: Bulirsch LM, Saßmannshausen M, Nadal J, et al. *Br J Ophthalmol* Epub ahead of print: [please include Day Month Year]. doi:10.1136/bjophthalmol-2020-318672

tomography (SD-OCT) (Spectralis HRA2+OCT, Heidelberg Engineering, Heidelberg, Germany) with acquisition of central 30°×30° near-infrared reflectance ($\lambda=820$ nm, ART (automatic real time) at least 15 frames). SD-OCT volume raster scans consisted of at least 19 B-scans (distance between neighbouring B-scans ≤ 240 μm) and a field size of at least 20°×15° (centred to the fovea) and included at least 5 frames (ART mode).

Assessment of structural outcomes

Morphological effects were quantified to assess structural efficacy of brolocuzumab. Three parameters were determined: foveal centre point (FCP), defined as the distance (μm) between the internal limiting membrane (ILM) and Bruch's membrane (BM); central subfield retinal thickness (CSRT), defined as the mean retinal thickness (μm) between ILM and BM of the circular area within 1 mm diameter around the centre of the fovea; as well as macular volume, defined as the mean volume of the retina in a circular area within 3 mm diameter around the fovea. For thickness and volume quantification, the automated segmentation of the Heidelberg Eye Explorer software V.2 (Heidelberg Engineering, V.2.4.1, Heidelberg, Germany) was carefully reviewed and, if indicated, manually corrected in each B-scan. Furthermore, a thorough evaluation of fluid distribution within the retina was performed with qualitative assessment of presence of subretinal, intraretinal and/or subretinal pigment epithelial (sub-RPE) fluid.

Statistical analysis

Statistical analysis was performed using SPSS (IBM SPSS Statistics V.25). The baseline characteristics are summarised by mean \pm SD and range (minimum–maximum) for normally distributed continuous variables.

BCVA was measured in decimals and for the purpose of this analysis converted to the logarithm of the minimum angle of resolution (logMAR).^{26–28}

Functional and structural outcomes were determined at baseline and at first visit after switch to brolocuzumab (visit 1). The outcomes were differences for each functional (logMAR) and structural (FCP, CSRT and macular volume) parameter between visit 1 and baseline. Descriptive values of these outcomes are shown with mean \pm SD and (95% CI). Linear mixed-effects models with the change of functional or structural parameter as the response variable were used to analyse the effects of treatment with brolocuzumab. To account for dependencies between measurements from the same patient, patient-specific random intercepts and slopes (analysis of longitudinal data) were included in the mixed-effects models. P values smaller than 0.05 were considered significant.

RESULTS

Study cohort

Between 16 March 2020 and 15 October 2020, a total of 207 brolocuzumab injections were performed with a mean (\pm SD) of 3.09 ± 1.55 (range 1–7) brolocuzumab injections per eye. Sixty-three eyes of 57 patients (30 females, 52.6%) with a mean (\pm SD) age of 79.5 ± 6.7 (58–94) years were treated with at least one brolocuzumab intravitreal injection (baseline) and were further assessed at the first visit after brolocuzumab injection (visit 1) after a mean of 4.3 ± 0.8 weeks (2.9–7.7). Table 1 summarises the results of descriptive analysis of the study cohort.

During the entire observation period, patients treated with brolocuzumab were seen across a mean follow-up of 16.04 ± 6.86 (2.86–27.43) weeks. All patients had received other anti-VEGF

Table 1 Study cohort characteristics at baseline (BSL) and visit 1 (V1)

	BSL		V1	
Per patient	n=57			
	Mean\pmSD	Range		
Age (years) (mean \pm SD)	79.46 \pm 6.65	58–94		
	N	%		
Gender, female (n) (%)	30	52.6		
Per eye (mean\pmSD)	n=63		Mean\pmSD	Range
BCVA (logMAR)	–0.39 \pm 0.28	0–1.2	–0.41 \pm 0.31	0–1.3
FCP (μm)	363.32 \pm 133.03	89–826	296.51 \pm 113.96	88–738
CSRT (μm)	409.43 \pm 112.32	224–784	342.67 \pm 99.05	163–707
Macular volume (mm^3)	2.72 \pm 0.51	2.07–4.63	2.46 \pm 0.44	1.9–4.31

BCVA, best-corrected visual acuity; CSRT, central subfield retinal thickness; FCP, foveal centre point.

agents before switching to brolocuzumab. Treatment history comprised a mean of 32.5 ± 25.6 (3–126) anti-VEGF injections per eye over a mean period of 4.2 ± 3.3 (0.3–13.4) years.

Functional outcome

Mean BCVA was -0.39 ± 0.28 (0–1.2) logMAR at baseline. After switch to brolocuzumab, a mean change in visual acuity of -0.03 ± 0.14 (95% CI -0.01 ± 0.06) logMAR was detected at visit 1 ($p=0.115$). See table 1 for absolute values at baseline and visit 1 and table 2 for the change of functional outcome under brolocuzumab treatment. A graphical outcome representation is provided in figure 1.

Anatomical outcomes

Regarding structural quantitative parameters, mean (\pm SD) FCP was 363.32 ± 133.03 (89–826) μm , mean CSRT was 409.43 ± 112.32 (224–784) μm and mean macular volume was 2.72 ± 0.51 (2.07–4.63) mm^3 at baseline. At visit 1, mean change in FCP was -66.81 ± 72.64 (-85.1 ; -48.5) μm , in CSRT -66.76 ± 60.71 (-82.05 ; -51.47) μm and in macular volume -0.27 ± 0.24 (-0.33 ; -0.2) mm^3 (all: $p<0.001$). See table 1 for absolute values at baseline and visit one and table 2 for the changes of structural parameters under brolocuzumab treatment. A graphical representation of the outcomes is provided in figure 1.

Assessment of qualitative SD-OCT parameters

For the following analysis, two eyes of two patients with frequent previous injections but without intraretinal, subretinal fluid or sub-RPE fluid at baseline were excluded. Sixty-one eyes (96.8%) of 55 patients presented at baseline with intraretinal and/or subretinal and/or sub-RPE fluid (eyes with subretinal, intraretinal and sub-RPE fluid: $n=7$; eyes with only subretinal and intraretinal fluid: $n=4$; eyes with subretinal and sub-RPE fluid: $n=18$; eyes with intraretinal and sub-RPE fluid: $n=6$; eyes with only subretinal fluid: $n=13$; eyes with only intraretinal fluid: $n=12$, eyes with only sub-RPE fluid: $n=1$) detected by SD-OCT imaging. At visit 1, absence of fluid in any retinal compartment was detected in 18 eyes (29.5%) of 18 patients.

At baseline, 27 eyes of 27 patients demonstrated (with or without fluid in any other compartment) intraretinal and 42 eyes of 38 patients demonstrated subretinal fluid, respectively. After initiation of brolocuzumab treatment (visit 1), 11 eyes (40.7%) of 11 patients with intraretinal and 24 eyes (57.1%) of 21 patients with subretinal fluid at baseline showed complete resolution. Out of the 30 eyes of 27 patients demonstrating sub-RPE fluid at

Table 2 Functional and structural outcomes after switch to brolucizumab

Outcome	Mean±SD	95% CI	P value
Change BCVA (logMAR)	-0.03±0.14	(-0.01 to 0.06)	0.115
Change FCP (µm)	-66.81±72.63	(-85.10 to -48.52)	<0.001
Change CSRT (µm)	-66.76±60.71	(-82.05 to -51.47)	<0.001
Change macular volume (mm ³)	-0.27±0.24	(-0.33 to -0.20)	<0.001

BCVA, best-corrected visual acuity; CSRT, central subfield retinal thickness; FCP, foveal centre point.

baseline, complete sub-RPE fluid resolution at visit 1 was detectable in only 6 eyes (20%) of 6 patients. For exemplary cases of structural response to brolucizumab treatment, see [figures 2–5](#).

Adverse events

During the observation period, adverse events were reported in eight eyes (12.7%) of eight patients. In one patient, a vitreous haemorrhage occurred immediately following the intravitreal injection, which was considered as a procedure-related and not drug-related event. Vitreous haemorrhage resolved spontaneously and required no further therapeutic intervention.

Out of the 63 eyes of 57 patients treated with brolucizumab, 7 eyes (11.1%) of 7 patients (3.38% per 207 given injections) with a mean age of 80.7±9.1 years (4 females, 57.1%) developed various degrees of IOI, which was suspected to be drug related. Out of the seven eyes with IOI, anterior uveitis with anterior chamber cells only was noted in two patients. Four eyes had additional signs of intermediate uveitis with occurrence of

anterior and vitreous cells. In one patient, segmental periarteriolar sheathing was noted without concurrent retinal vascular occlusion ([figure 6](#)). Due to IOI development, four patients sought medical attention at an unscheduled visit, while three patients presented at a regularly scheduled visit. Once signs of IOI development were detected, treatment with brolucizumab for nAMD was discontinued and patients were switched back to an alternate anti-VEGF drug.

Regarding the time of IOI development, four patients developed signs of IOI after the first brolucizumab injection with occurrence of first symptoms after a mean of 19.0±6.5 (12–25) days. Medical attention was sought after a mean of 23.8±6.4 (13–29) days after injection. In the three patients with IOI development after the second brolucizumab injection, first symptoms occurred after a mean of 7.3±5.6 (2–15) days. Medical advice was sought after 8.7±5.2 (5–16) days. During the observation period, no patient receiving at least three brolucizumab injections developed any signs of IOI.

Initial symptoms differed between the patients with IOI development. Eye redness as the initial symptom was reported by four patients, while three reported floaters and/or cloudy vision. One patient reported a decrease in visual acuity. None of the patients with documented adverse events reported earlier episodes of uveitis or ocular autoimmune disease, except for one patient who had a history of bilateral macular oedema interpreted as Irvine-Gass syndrome following routine cataract surgery with intraocular lens implantation.

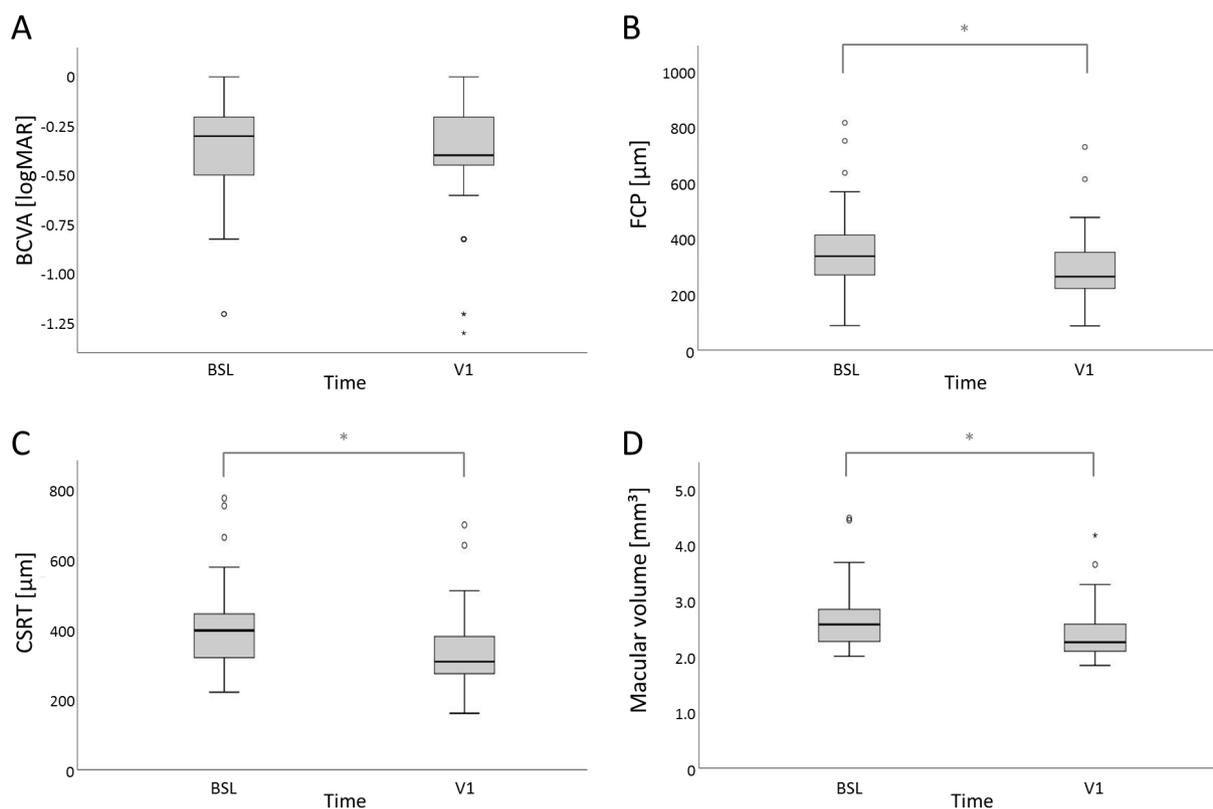


Figure 1 Boxplots (each box: median; 75% and 25% quartiles; whiskers: 10% and 90% quantiles; outliers are defined as sample outliers 1.5 times (circles) and 3 times (star) the IQR above the upper quartile and below the lower quartile) at baseline (BSL) and visit 1 (V1) for (A) best-corrected visual acuity (BCVA (logMAR)), (B) focal centre point (FCP (µm)), (C) central subfield retinal thickness (CSRT (µm)) and (D) macular volume (mm³). Structural outcomes (B–D) showed a significant change after brolucizumab treatment.

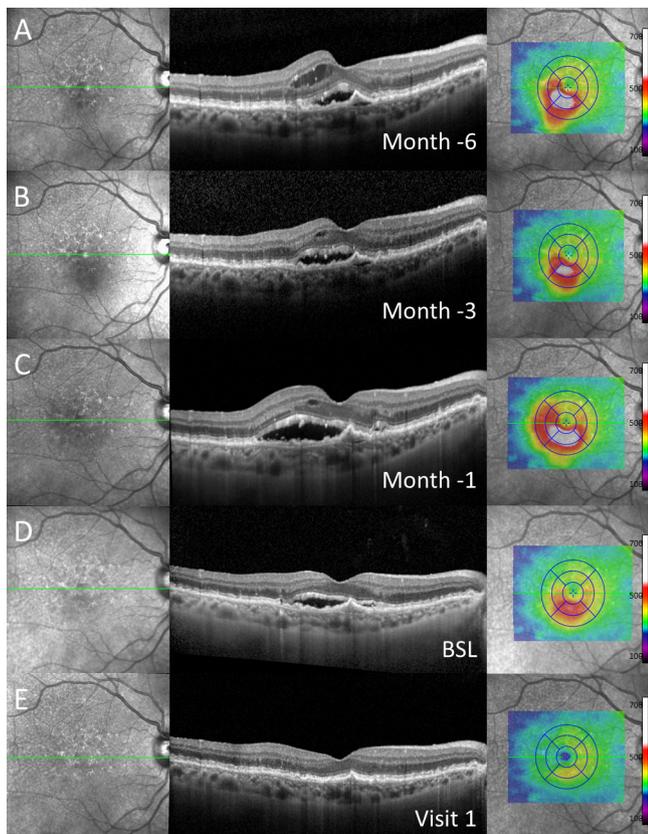


Figure 2 Exemplary case of a patient with subretinal fluid at baseline (BSL) (D) and both intraretinal and subretinal fluid 1 (C), 3 (B) and 6 (A) months in historical imaging before baseline despite regular anti-vascular endothelial growth factor (anti-VEGF) treatment as demonstrated in (from left to right) near-infrared imaging, spectral-domain optical coherence tomography through the fovea and colour-coded two-dimensional thickness map for total retinal thickness. One month after switch to brodalumab (E, visit 1), complete resolution of subretinal fluid was detected. Note: Before switch to brodalumab, the patient received repetitive, high-frequency intravitreal injections of other anti-VEGF agents over a longer period of time.

Treatment was initiated promptly according to the severity of the documented IOI. In cases with only mild IOI and without signs of posterior uveitis or vasculitis, topical therapy with corticosteroid eye drops (dexamethasone or prednisolone) was initiated with application ranging from four times a day up to every hour and ointment (prednisolone) before hours of sleep. In four patients, topical therapy was supplemented by a subconjunctival injection of dexamethasone (8 mg). In the single patients who presented with vasculitis and in three patients with more severe IOI, additional systemic therapy with corticosteroids was initiated in a body weight adapted dose (ie, 60 or 80 mg) and subsequently tapered off according to IOI activity. None of the seven patients demonstrated persistent clinically relevant change in visual acuity once IOI had ceased.

DISCUSSION

Efficacious therapy with maintaining visual acuity gains during the upload phase in the long-term management of patients with chronic nAMD still represents an unmet need. Recently, brodalumab was approved as a new anti-VEGF agent for intravitreal administration in patients with nAMD. However, only limited data of its use in a real-world setting are available to date.^{16 17 20–22}

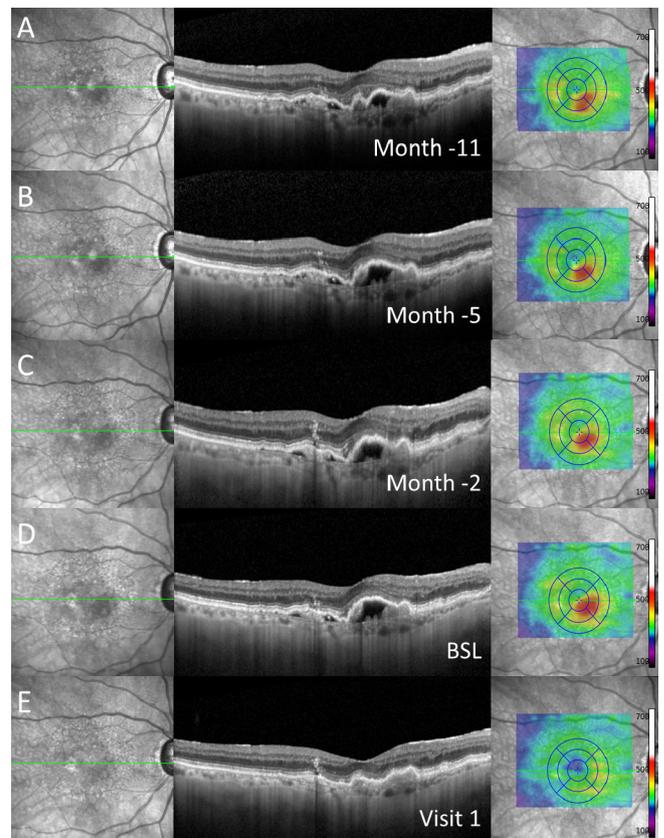


Figure 3 Exemplary case of a patient with subretinal pigment epithelial (sub-RPE) fluid at baseline (BSL) (D) and in historical imaging up to 11 months (A–C) before switch to brodalumab as demonstrated in (from left to right) near-infrared imaging, spectral-domain optical coherence tomography through the fovea and colour-coded two-dimensional thickness map for total retinal thickness. At visit 1 (E), complete resolution of sub-RPE fluid was detected. Note: Before switch to brodalumab, the patient received repetitive, high-frequency intravitreal injections of other anti-vascular endothelial growth factor agents over a longer period of time.

Herein, we report early experiences with brodalumab use in clinical routine in a single centre in Europe following approval in February 2020 (SHIFT study). Our findings indicate that initiation of intravitreal brodalumab therapy in previously treated patients with nAMD (switch) may be an option in particular with regard to the morphological effects in recalcitrant cases who previously received multiple anti-VEGF injections without satisfactory resolution of fluid in various anatomical compartments. A significant reduction on average of retinal thickness parameters, including FCB, CSRT and macular volume, was observed demonstrating a favourable response on morphological signs for disease activity. Although differences in study design and analysis methods limit a direct comparison, our findings mirror a recent real-world report by Sharma *et al*, who also showed improvement in structural outcomes in 42 eyes of 42 patients with nAMD over a mean (\pm SD) observational period of 7.2 ± 3.6 weeks.¹⁶ In accordance with our study, Avaylon and colleagues have also found beneficial structural outcomes at first visit following switch to brodalumab.¹⁷ Although the analysis of this group is based on a rather small cohort of only six patients, our results confirm the observations by Avaylon *et al* of improvement of various OCT characteristics.

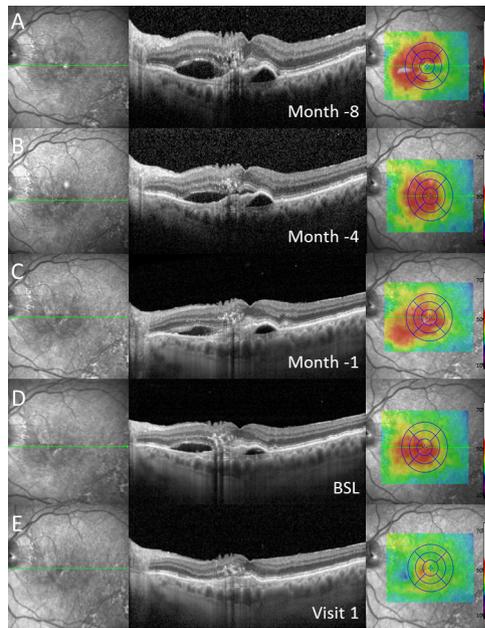


Figure 4 Exemplary case of a patient with subretinal pigment epithelial (sub-RPE) and subretinal fluid at baseline (BSL) (D) as well as in historical imaging up to 8 months (A–C) before switch to brodalumab as demonstrated in (from left to right) near-infrared imaging, spectral-domain optical coherence tomography through the fovea and colour-coded two-dimensional thickness map for total retinal thickness. Retinal imaging at visit 1 (E) revealed complete resolution of subretinal and sub-RPE fluid. Note: Before switch to brodalumab, the patient received repetitive, high-frequency intravitreal injections of other anti-vascular endothelial growth factor agents over a longer period of time.

When interpreting the functional outcome in our study, it needs to be considered that the SHIFT cohort comprises previously treated patients only, with up to 126 previous intravitreal anti-VEGF injections (with a mean (\pm SD) of 32.5 ± 25.6). This might not only impact the BCVA before initial injection of brodalumab of -0.37 ± 0.28 logMAR but may also explain the lack of BCVA improvement despite significant change in structurally defined efficacy outcome parameters. This phenomenon has been observed before in other switch studies. Given the long-standing ocular history of exudative AMD in our cohort, underlying structural, in part irreversible damage may contribute to a limited potential of visual recovery.^{29–32} The phenomenon of disconnect between morphology and function has been observed in other switch studies of patients with nAMD.^{29–32}

Since the approval of brodalumab, safety signals of development of IOI following brodalumab treatment, ranging from anterior chamber cells to retinal vasculitis with or without occlusion and with or without moderate and severe visual loss, have been reported.^{13 19 21 23 33} The pathogenesis of the inflammatory events is not yet understood and needs further investigations which are ongoing. In the pivotal phase-III HAWK and HARRIER studies of treatment-naïve nAMD patients, an overall rate of 4.6% of any IOI was reported following a review by a safety review committee (SRC).²⁴ The rate of the development of retinal vasculitis was reported to be 3.3% (36/1088) and that of concomitant retinal vasculitis and retinal vascular occlusion was 2.1% (23/1088), respectively.¹³ The overall incidence of at least moderate vision loss due to IOI was <1%. Recently, the American Society of Retina Specialists referred to post approval,

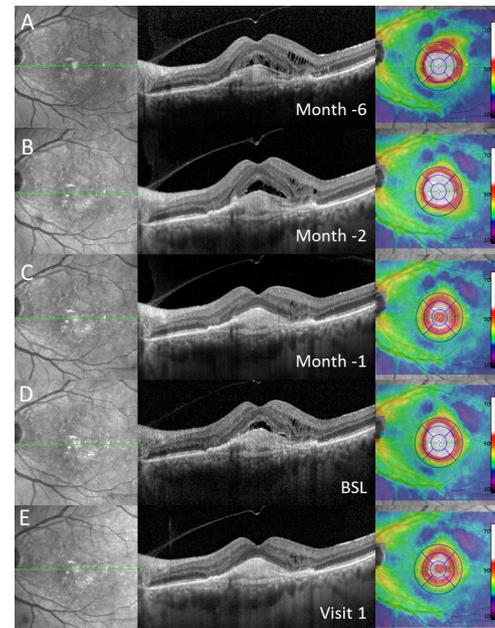


Figure 5 Exemplary case of a patient with intraretinal and subretinal fluid at baseline (BSL) (D) and in historical images 1 (C), 2 (B) and 6 (A) months before switch to brodalumab as demonstrated in (from left to right) near-infrared imaging, spectral-domain optical coherence tomography through the fovea and colour-coded two-dimensional thickness map for total retinal thickness. One month after switch (E, visit 1), complete resolution of subretinal and incomplete resolution of intraretinal fluid was demonstrated. Note: Before switch to brodalumab, the patient received repetitive, high-frequency intravitreal injections of other anti-vascular endothelial growth factor agents over a longer period of time.

reported IOI cases that the majority (88%) was of female gender in accordance with other published reports of brodalumab cases.^{18 20–22} In line with these previous reports, IOI occurred in our small cohort more frequently in female patients, while the case of retinal arteriolar vasculitis was detected in a male patient who showed a favourable outcome in the absence of retinal vascular occlusion and resolution of the intraocular inflammation following steroid therapy.

In the patients reported here, anti-inflammatory treatment was initiated promptly following detection of IOI and included topical, subconjunctival and systemic corticosteroid therapy. In none of the patients functional deficits occurred following IOI resolution. In the meantime, recommendations have been published on the management of patients with nAMD who develop IOI following brodalumab treatment.²⁵

Various limitations need to be considered for this study. This is an observational study in a relatively small cohort of 57 patients. An important limitation of our study is the short review period post switch to brodalumab of 1 month. In this real-world study, we applied a treat & extend scheme immediately after the first injection and did not perform three fixed 4-weekly injections post switch to brodalumab as proposed in other switch studies. Therefore, in this relatively small cohort, data analyses beyond 1 month would be based on a variable number of injections after switch and would not allow a meaningful analysis and comparison between patients. Accordingly, the same applies to an analysis of the treatment history before switch to brodalumab. Furthermore, no treatment-naïve patients were investigated compromising any overall conclusion on structural and visual

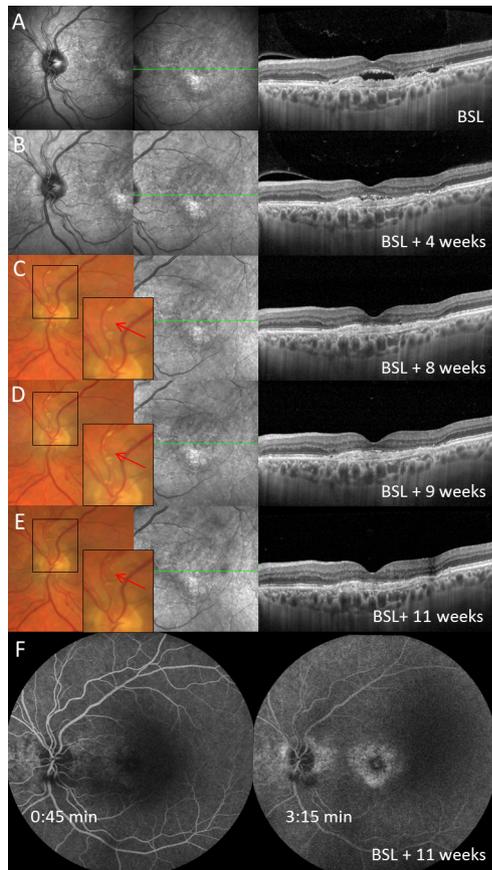


Figure 6 Patient with first brolicizumab injection at baseline (BSL, row A), first follow-up visit (visit 1) and second brolicizumab injection 4 weeks later (row B) and development of non-occlusive retinal vasculitis with segmental perivasculature sheathing 4 weeks following the second brolicizumab injection (row C). Further follow-up visits (row D–F) as presented by multimodal imaging (from left to right: near-infrared imaging (row A–C) or digital colour fundus photography (row D and E) of the optic nerve head, near-infrared imaging of the macula and spectral-domain optical coherence tomography (SD-OCT) through the fovea). The intra-arterial plaque secondary to retinal vasculitis and its disappearance after treatment initiation is marked by a red arrow in row C, D and E. Seven weeks after first diagnosis of retinal vasculitis, fluorescein angiography did not show signs of vascular occlusion or retinal ischaemia (row F). Of note, subretinal fluid completely resolved after two intravitreal injections of brolicizumab (BSL and V1) in SD-OCT imaging.

outcome following treatment. Finally, the study cannot address the questions of potentially longer durability of brolicizumab compared with other anti-VEGF agents in particular during a treat & extend regimen. To determine this, both longitudinal assessments in real-world settings and randomised clinical nAMD trials are warrant.

In summary, the SHIFT study reports real-world early experiences outside RCTs with brolicizumab in previously anti-VEGF-treated patients with nAMD. A beneficial morphological effect was recorded pointing towards a strong antihyperpermeability effect of this agent. Safety issues with regard to IOI remain a concern and require further investigations with regard to the underlying mechanisms as well as risk mitigation measures.

Correction notice This paper has been corrected since it was published online first.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests Non-financial support from Heidelberg Engineering to ST, LB, RL, MS and FGH; from CenterVue to ST and FGH; from Optos to ST, LB, RL, MS and FGH; from Carl Zeiss Meditec to ST, LB, RF, FGH and MS. Grant and personal fees from Allergan, Novartis, Bayer and Heidelberg Engineering to ST and FGH; from Apellis, Carl Zeiss Meditec, Acucela, Genentech/Roche, Boehringer-Ingelheim, LIN Bioscience, Pixium, Kanghiong, Oxurion, Grayburg Vision, Stealth BioTherapeutics, Geuder and Iveric Bio to FGH. JN: None declared.

Patient consent for publication Obtained.

Ethics approval This study does not contain personal information from any identifiable individual. Because this study is a retrospective analysis of data obtained in clinical routine care at an academic university setting, no ethics approval is required.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article or uploaded as supplementary information. All data relevant to this study are included in the article.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iD

Sarah Thiele <http://orcid.org/0000-0002-1123-6700>

REFERENCES

- Lim LS, Mitchell P, Seddon JM, *et al*. Age-related macular degeneration. *Lancet* 2012;379:1728–38.
- Rosenfeld PJ, Brown DM, Heier JS, *et al*. Ranibizumab for neovascular age-related macular degeneration. *N Engl J Med* 2006;355:1419–31.
- Bloch SB, Larsen M, Munch IC. Incidence of legal blindness from age-related macular degeneration in Denmark: year 2000 to 2010. *Am J Ophthalmol* 2012;153:209–13.
- Holz FG, Tadayoni R, Beatty S, *et al*. Multi-country real-life experience of anti-vascular endothelial growth factor therapy for wet age-related macular degeneration. *Br J Ophthalmol* 2015;99:220–6.
- Eghøj MS, Sørensen TL. Tachyphylaxis during treatment of exudative age-related macular degeneration with ranibizumab. *Br J Ophthalmol* 2012;96:21–3.
- Forooghian F, Chew EY, Meyerle CB, *et al*. Investigation of the role of neutralizing antibodies against bevacizumab as mediators of tachyphylaxis. *Acta Ophthalmol* 2011;89:e206–7.
- Wong WL, Su X, Li X, *et al*. Global prevalence of age-related macular degeneration and disease burden projection for 2020 and 2040: a systematic review and meta-analysis. *Lancet Glob Heal* 2014;2:e106–16.
- Schmidt-Erfurth U, Chong V, Loewenstein A, *et al*. Guidelines for the management of neovascular age-related macular degeneration by the European Society of retina specialists (EURETINA). *Br J Ophthalmol* 2014;98:1144–67.
- Boulanger-Scemama E, Querques G, About F, *et al*. Ranibizumab for exudative age-related macular degeneration: a five year study of adherence to follow-up in a real-life setting. *J Fr Ophtalmol* 2015;38:620–7.
- Ehken C, Helms M, Böhringer D, *et al*. Association of treatment adherence with real-life Va outcomes in AMD, DME, and BRVO patients. *Clin Ophthalmol* 2018;12:13–20.
- Yang S, Zhao J, Sun X. Resistance to anti-VEGF therapy in neovascular age-related macular degeneration: a comprehensive review. *Drug Des Devel Ther* 2016;10:1857–67.
- Gasparini JL, Fawzi AA, Khondkaryan A, *et al*. Bevacizumab and ranibizumab tachyphylaxis in the treatment of choroidal neovascularisation. *Br J Ophthalmol* 2012;96:14–20.
- Dugel PU, Singh RP, Koh A, *et al*. Hawk and harrier: Ninety-Six-Week outcomes from the phase 3 trials of Brolicizumab for neovascular age-related macular degeneration. *Ophthalmology* 2021;128:30570–4.
- Dugel PU, Jaffe GJ, Sallstig P, *et al*. Brolicizumab versus aflibercept in participants with neovascular age-related macular degeneration: a randomized trial. *Ophthalmology* 2017;124:1296–304.
- Holz FG, Dugel PU, Weissgerber G, *et al*. Single-Chain antibody fragment VEGF inhibitor RTH258 for neovascular age-related macular degeneration: a randomized controlled study. *Ophthalmology* 2016;123:1080–9.
- Sharma A, Kumar N, Parachuri N, *et al*. Brolicizumab-early real-world experience: brew study. *Eye* 2021;35:1045–7.
- Avaylon J, Lee S, Gallemore RP. Case series on initial responses to intravitreal brolicizumab in patients with recalcitrant chronic wet age-related macular degeneration. *Int Med Case Rep J* 2020;13:145–52.

- 18 Witkin AJ, Hahn P, Murray TG, *et al.* Occlusive retinal vasculitis following intravitreal Brolucizumab. *J Vitreoretin Dis* 2020;4:269–79.
- 19 Brolucizumab Safety information - Post-marketing data. Available: <https://www.brolucizumab.info/post-marketing-data> [Accessed 16 Nov 2020].
- 20 Jain A, Chea S, Matsumiya W, *et al.* Severe vision loss secondary to retinal arteriolar occlusions after multiple intravitreal brolucizumab administrations. *Am J Ophthalmol Case Rep* 2020;18:100687.
- 21 Bauml CR, Spaide RF, Vajzovic L, *et al.* Retinal vasculitis and intraocular inflammation after intravitreal injection of Brolucizumab. *Ophthalmology* 2020;127:1345–59.
- 22 Haug SJ, Hien DL, Uludag G, *et al.* Retinal arterial occlusive vasculitis following intravitreal brolucizumab administration. *Am J Ophthalmol Case Rep* 2020;18:100680.
- 23 Sharma A, Kumar N, Parachuri N. Brolucizumab-Related retinal vasculitis: emerging disconnect between clinical trials and real world. Springer us 2020.
- 24 Monés J, Srivastava SK, Jaffe GJ, *et al.* Risk of inflammation, retinal vasculitis, and retinal Occlusion-Related events with Brolucizumab: post hoc review of hawk and harrier. *Ophthalmology* 2020. doi:10.1016/j.ophtha.2020.11.011. [Epub ahead of print: 15 Nov 2020].
- 25 Holz FG, Heinz C, Wolf A. Intraocular inflammation with brolucizumab use: patient management, diagnosis, therapy. *Ophthalmologie* 2021;118:1–3.
- 26 Khoshnood B, Mesbah M, Jeanbat V, *et al.* Transforming scales of measurement of visual acuity at the group level. *Ophthalmic Physiol Opt* 2010;30:816–23.
- 27 Beck RW, Moke PS, Turpin AH, *et al.* A computerized method of visual acuity testing: adaptation of the early treatment of diabetic retinopathy study testing protocol. *Am J Ophthalmol* 2003;135:194–205.
- 28 Kniestedt C, Stamper RL. Visual acuity and its measurement. *Ophthalmol Clin North Am* 2003;16:155–70.
- 29 Gale RP, Pearce I, Eter N, *et al.* Anatomical and functional outcomes following switching from aflibercept to ranibizumab in neovascular age-related macular degeneration in Europe: SAFARI study. *Br J Ophthalmol* 2020;104:493–9.
- 30 Mantel I, Gillies MC, Souied EH. Switching between ranibizumab and aflibercept for the treatment of neovascular age-related macular degeneration. *Surv Ophthalmol* 2018;63:638–45.
- 31 García-Layana A, Figueroa MS, Araiz J, *et al.* Treatment of exudative age-related macular degeneration: focus on aflibercept. *Drugs Aging* 2015;32:797–807.
- 32 Ho VY, Yeh S, Olsen TW, *et al.* Short-Term outcomes of aflibercept for neovascular age-related macular degeneration in eyes previously treated with other vascular endothelial growth factor inhibitors. *Am J Ophthalmol* 2013;156:23–8.
- 33 Study of safety and efficacy of Brolucizumab 6 Mg dosed every 4 weeks compared to aflibercept 2 Mg dosed every 4 weeks in patients with retinal fluid despite frequent anti-VEGF injections (merlin). Available: <https://clinicaltrials.gov/ct2/show/NCT03710564> [Accessed 16 Nov 2020].