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# Retinal Optical Coherence Tomography in Neuromyelitis Optica

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## Abstract

### Background and Objectives

To determine optic nerve and retinal damage in aquaporin-4 antibody (AQP4-IgG)-seropositive neuromyelitis optica spectrum disorders (NMOSD) in a large international cohort after previous studies have been limited by small and heterogeneous cohorts.

### Methods

The cross-sectional Collaborative Retrospective Study on retinal optical coherence tomography (OCT) in neuromyelitis optica collected retrospective data from 22 centers. Of 653 screened participants, we included 283 AQP4-IgG-seropositive patients with NMOSD and 72 healthy controls (HCs). Participants underwent OCT with central reading including quality

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## Glossary

**AQP4-IgG** = aquaporin-4 IgG; **CROCTINO** = Collaborative Retrospective Study on retinal OCT in Neuromyelitis Optica; **EDSS** = Expanded Disability Status Scale; **GCIP** = ganglion cell and inner plexiform; **HC** = healthy control; **HC-VA** = high-contrast visual acuity; **INL** = inner nuclear layer; **MME** = microcystic macular edema; **MS** = multiple sclerosis; **MOG-IgG** = myelin oligodendrocyte glycoprotein IgG; **NMOSD** = neuromyelitis optica spectrum disorder; **NMOSD-NON** = NMOSD eyes without a history of ON; **OCT** = optical coherence tomography; **ON** = optic neuritis; **pRNFL** = peripapillary retinal nerve fiber layer; **SE** = standard error; **VA** = visual acuity; **VEP** = visually evoked potential.

control and intraretinal segmentation. The primary outcome was thickness of combined ganglion cell and inner plexiform (GCIP) layer; secondary outcomes were thickness of peripapillary retinal nerve fiber layer (pRNFL) and visual acuity (VA).

## Results

Eyes with ON (NMOSD-ON, N = 260) or without ON (NMOSD-NON, N = 241) were assessed compared with HCs (N = 136). In NMOSD-ON, GCIP layer ( $57.4 \pm 12.2 \mu\text{m}$ ) was reduced compared with HC (GCIP layer:  $81.4 \pm 5.7 \mu\text{m}$ ,  $p < 0.001$ ). GCIP layer loss ( $-22.7 \mu\text{m}$ ) after the first ON was higher than after the next ( $-3.5 \mu\text{m}$ ) and subsequent episodes. pRNFL observations were similar. NMOSD-NON exhibited reduced GCIP layer but not pRNFL compared with HC. VA was greatly reduced in NMOSD-ON compared with HC eyes, but did not differ between NMOSD-NON and HC.

## Discussion

Our results emphasize that attack prevention is key to avoid severe neuroaxonal damage and vision loss caused by ON in NMOSD. Therapies ameliorating attack-related damage, especially during a first attack, are an unmet clinical need. Mild signs of neuroaxonal changes without apparent vision loss in ON-affected eyes might be solely due to contralateral ON attacks and do not suggest clinically relevant progression but need further investigation.

Patients with neuromyelitis optica spectrum disorder (NMOSD) experience recurrent optic neuritis (ON),<sup>1</sup> resulting in vision loss and decreased quality of life.<sup>1-4</sup> According to our understanding, there are at least 3 subtypes based on serostatus: Up to 3 of 4 patients manifest as anti-aquaporin-4 IgG (AQP4-IgG) seropositive. Approximately half of the AQP4-IgG-seronegative patients manifest as anti-myelin oligodendrocyte glycoprotein IgG (MOG-IgG) seropositive, and half are double seronegative.<sup>5</sup> Yet, clinical correlates of serologic phenotypes, including subclinical or clinical retinal degeneration and vision loss, remain unclear.<sup>6,7</sup>

Optical coherence tomography (OCT) is an interferometric technique producing high-resolution retinal images.<sup>1,8</sup> OCT has become a reliable tool for diagnosing and monitoring neurologic and neuro-ophthalmologic diseases, especially for quantifying neurodegeneration after ON.<sup>1</sup> Because of limited samples and varying methods, existing OCT studies in NMOSD are inconsistent regarding the amount of retinal neurodegeneration with and without ON. Previous studies also failed to address the influence of retinal neurodegeneration on microcystic macular edema (MME) and function.<sup>2,6,7</sup> These issues together with heterogeneities and the often monocentric character of previous cohorts limit the relevance of meta-analyses.

To overcome these limitations, we performed an OCT analysis of AQP4-IgG-seropositive patients with NMOSD in an international multicenter study, termed Collaborative Retrospective Study on retinal OCT in Neuromyelitis Optica (CROCTINO).<sup>9</sup>

It represents the largest NMOSD OCT data set and additionally validated an OCT postprocessing approach to circumvent differences in acquisition and imaging processing protocols inherent to pooled analyses.<sup>10</sup> Outcomes include: (1) distinguishing retinal neurodegeneration after ON from subtle damage in clinically unaffected eyes, (2) defining frequency of MME, and (3) deriving structure-function correlations.

## Methods

### Study Design

This cross-sectional international multicenter study was performed under the aegis of the CROCTINO study, which was a collaborative effort within the Guthy-Jackson Charitable Foundation network.<sup>9</sup> Participating centers contributed OCT data (acquired between 2008 and 2018) and clinical metadata (acquired between 2000 and 2018, eTable 1, [links.lww.com/NXI/A557](https://links.lww.com/NXI/A557)).

### Cohort Selection

Inclusion criteria for this analysis were (1) patients diagnosed with NMOSD per the 2015 International Panel of NMOSD diagnosis criteria<sup>11</sup> and (2) having confirmed serum AQP4-IgG. Exclusion criteria were (1) comorbidities potentially confounding interpretation of OCT results (e.g., macular degeneration, glaucoma, and intracranial hypertension); (2) > 3 months distance between clinical and OCT data acquisition; (3) < 6 months between OCT imaging and most recent ON, or (4) an uncertain history of ON. The inclusion and

**Table 1** Cohort Description for AQP4-IgG–Seropositive Patients With NMOSD and HCs

	AQP4-IgG-seropositive NMOSD	HC
Subjects (N)	283	72
Eyes (N)	501	136
Sex (male/female, N/N [%/-%])	28/255 (9.9/90.1)	26/46 (36.1/63.9)
Age (y, mean ± SD)	44.1 ± 14.2	30.9 ± 7.7
Ethnicity (N [%])		
Asian	77 (27.2)	16 (22.2)
Black/African American	13 (4.6)	0 (0)
White, Hispanic/Latino	4 (1.4)	1 (1.4)
White, Non-Hispanic	159 (56.2)	55 (76.4)
Other/nonreported	30 (10.6)	0 (0)
Disease-modifying therapy (N [%])		
Rituximab	73 (25.8)	
Azathioprine	67 (23.7)	
Oral prednisolone	53 (18.7)	
Mycophenolate mofetil	48 (17.0)	
Methotrexate	8 (2.8)	
Time since onset (y, mean ± SD)	7.2 ± 6.7	.
EDSS score (median [IQR])	3.5 (2.0–4.5)	.
Patients with a clinical history of ON (N [%])	204 (72.1)	.
Eyes with a clinical history of ON (N [%])	260 (52)	.
No. of ON episodes/eye (N, median [min–max])	1 (1–6)	.
Time since last ON (mo, mean ± SD)	71 ± 57	.

Abbreviations: AQP4-IgG = aquaporin-4 antibodies; NMOSD = neuromyelitis optica spectrum disorder; HC = healthy control; N = number; OCT = optical coherence tomography; ON = optic neuritis episode. Age ( $p < 0.0001$ ) and sex ( $p < 0.0001$ ) were not matched.

exclusion criteria are depicted in eFigure 1, [links.lww.com/NXI/A556](https://links.lww.com/NXI/A556). AQP4-IgG testing was performed at the discretion of each investigator.

### Standard Protocol Approvals, Registrations, and Patient Consents

All participants gave written informed consent, and the study was approved by local ethics committees and conducted in accordance with the applicable laws and the current version of the Declaration of Helsinki. Data are reported according to STROBE reporting guidelines.<sup>12</sup>

### OCT

High-resolution imaging data were acquired using 3 different spectral domain OCT devices: Spectralis SD-OCT (Heidelberg Engineering, Heidelberg, Germany) at 15 centers (194 patients/358 eyes; 72 healthy controls [HCs]/136 eyes); Cirrus HD-OCT (Carl Zeiss Meditec, Dublin, CA) at 6 centers (58 patients/87 eyes); or Topcon 3D OCT-1 (Topcon, Tokyo, Japan) at 1 center (31 patients/56 eyes). All reading of OCT data was performed at Charité–Universitätsmedizin Berlin Translational Neuroimaging Group by 5 graders. Image quality was assessed using modified OSCAR-IB criteria by one of the graders, respectively.<sup>13,14</sup> OCT segmentation for the combined ganglion cell and inner plexiform (GCIP) layer and inner nuclear layer (INL) thicknesses was corrected semi-automatically using an in-house software.<sup>10,15</sup> In brief, GCIP layer and INL thicknesses were calculated from a 5-mm diameter cylinder around the fovea excluding the central 1-mm diameter cylinder from a macular volume scan.<sup>10</sup> The peripapillary retinal nerve fiber layer (pRNFL) thickness was measured and corrected according to the device protocol (Spectralis: peripapillary ring scan with 12° or approximately 3.4 mm diameter around the optic disc; Topcon and Cirrus: extraction from optic disc volume scan). For the current analysis, eyes were excluded from the analysis if neither ring nor macular scan passed quality control. We further excluded data from the less common instrument for 1 center. The final cohort included 364 macular and 481 peripapillary scans of 501 eyes from 283 patients and 136 eyes from 72 HCs. Lower numbers of macular scans compared with peripapillary scans were due to both lower submission of macular data and more exclusions based on quality concerns.

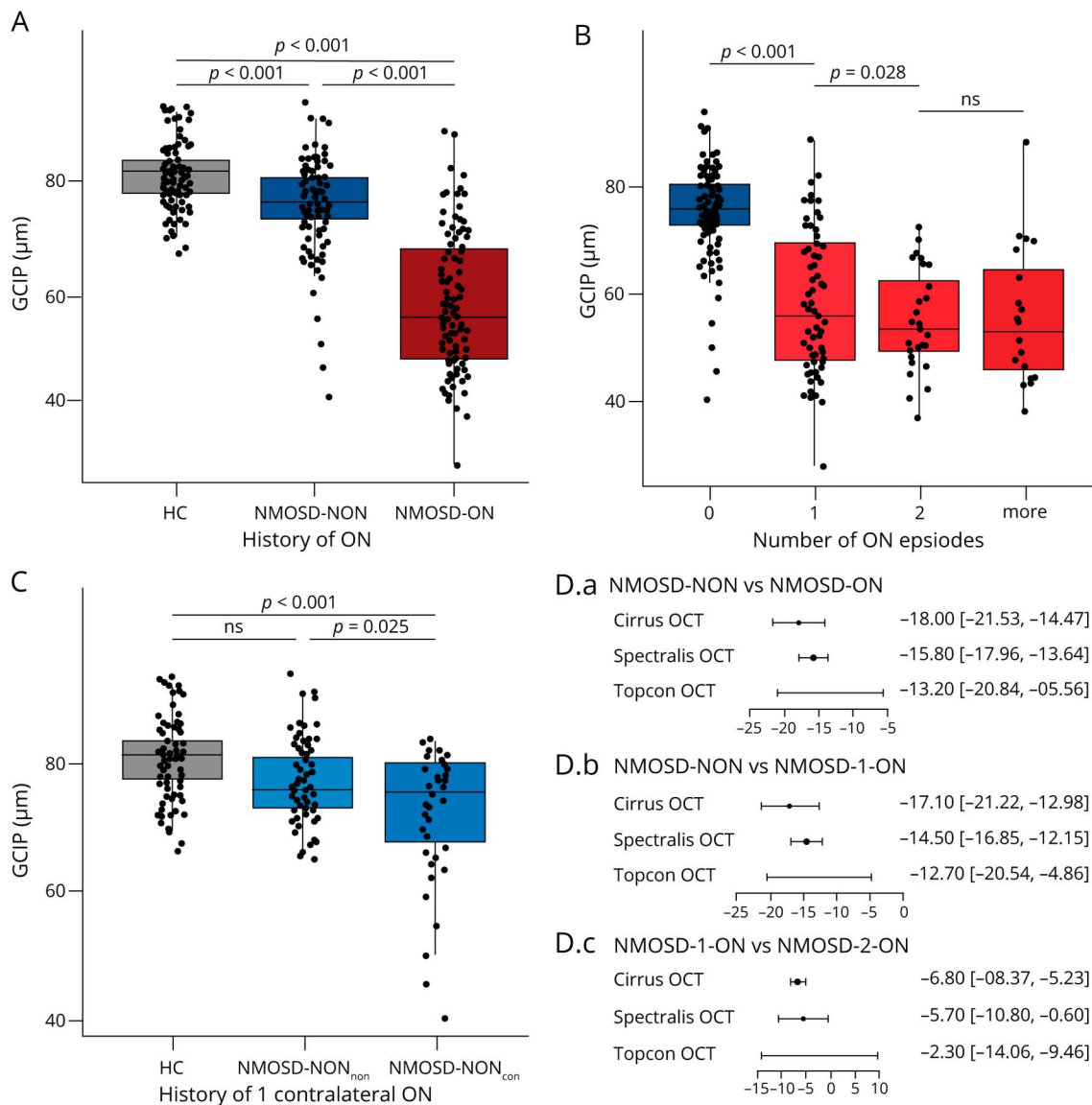
### Visual and Global Function Testing

High-contrast visual acuity (HC-VA) was available for 497 (99.2%) patient and 10 (13.9%) HC eyes. HC-VA was best corrected for 212 (42.3%) patient and 56 (41.2%) HC eyes, habitually corrected for 145 (28.9%) patient and 2 (<0.1%) HC eyes, and without correction for 140 (27.9%) patient eyes and 17 (1.3%) HC eyes. All VA data are reported as logMAR. VA measurement method was decided on discretion of each center. Visually evoked potentials (VEPs) were available for 167 (33.3%) patient eyes and 40 (29.4%) HC eyes, with P100 latency recorded as a binary value (normal/prolonged). Expanded Disability Status Scale (EDSS) scores were determined at the discretion of each center with data available for 180 (63.6%) patients.

### Statistical Analysis

Statistical analyses were performed with R version 3.6.1 using RStudio and R Markdown (RStudio Inc., Boston, MA).<sup>16</sup>  $p$  Values less than 0.05 were considered significant. We considered  $p$  values less than 0.10 a trend. Group matching by age and sex for confirmatory analyses was performed using automatic matching by R package MatchIt (method: exact). Group comparisons and correlations of OCT and VA values were performed using linear mixed-

**Figure 1** Group Comparisons of GCIP Layer Thickness



Boxplots of GCIP layer thicknesses ( $\mu\text{m}$ ) acquired by Heidelberg SD-OCT with values of individual eyes (jitter) for (A) HC (gray/left), NMOSD-NON (dark blue/middle), NMOSD-ON (dark red/right); for (B) number of ON episodes (NMOSD-NON dark blue/left, NMOSD-1-ON light red/left-middle, NMOSD-2-ON medium-red/right-middle, NMOSD- $\geq 3$ -ON medium-dark red/right); and for (C) HC (gray/left), NMOSD-NON<sub>non</sub> (light blue/middle), NMOSD-NON<sub>con</sub> (blue/right). (D) Forest plots for results from different OCT devices for (D.a) NMOSD-NON vs NMOSD-ON, (D.b) NMOSD-NON vs NMOSD-1-ON, and (D.c) NMOSD-1-ON vs NMOSD-2-ON (eFigure 2 and eTable 2, [links.lww.com/NXI/A556](https://links.lww.com/NXI/A556) and [links.lww.com/NXI/A557](https://links.lww.com/NXI/A557)). GCIP = ganglion cell and inner plexiform; HC = eyes of HCs; NMOSD-NON = eyes of patients with neuromyelitis optica without a history of ON; NMOSD-NON<sub>con</sub> = eyes of patients with neuromyelitis optica without a history of ipsilateral or contralateral ON but a history of contralateral ON; NMOSD-NON<sub>non</sub> = eyes of patients with neuromyelitis optica without a history of ipsilateral or contralateral ON; NMOSD-ON = eyes of patients with neuromyelitis optica with a history of ON; NMOSD-1-ON = eyes of patients with neuromyelitis optica with a history of 1 ON episode; NMOSD-2-ON = eyes of patients with neuromyelitis optica with a history of 2 ON episodes; ON = optic neuritis.

effect models. Intereye within-subject effects and effects of the center were included as random effects. Sex, ethnicity, and age were included as fixed effects for the analyses in the entire cohort. Age and sex were not included for the matched subset. For OCT parameters, the model was used for data from all devices separately and combined by Fisher combined probability test. The marginal and conditional coefficients of determination of the linear models were calculated with pseudo R-squared. All results are reported

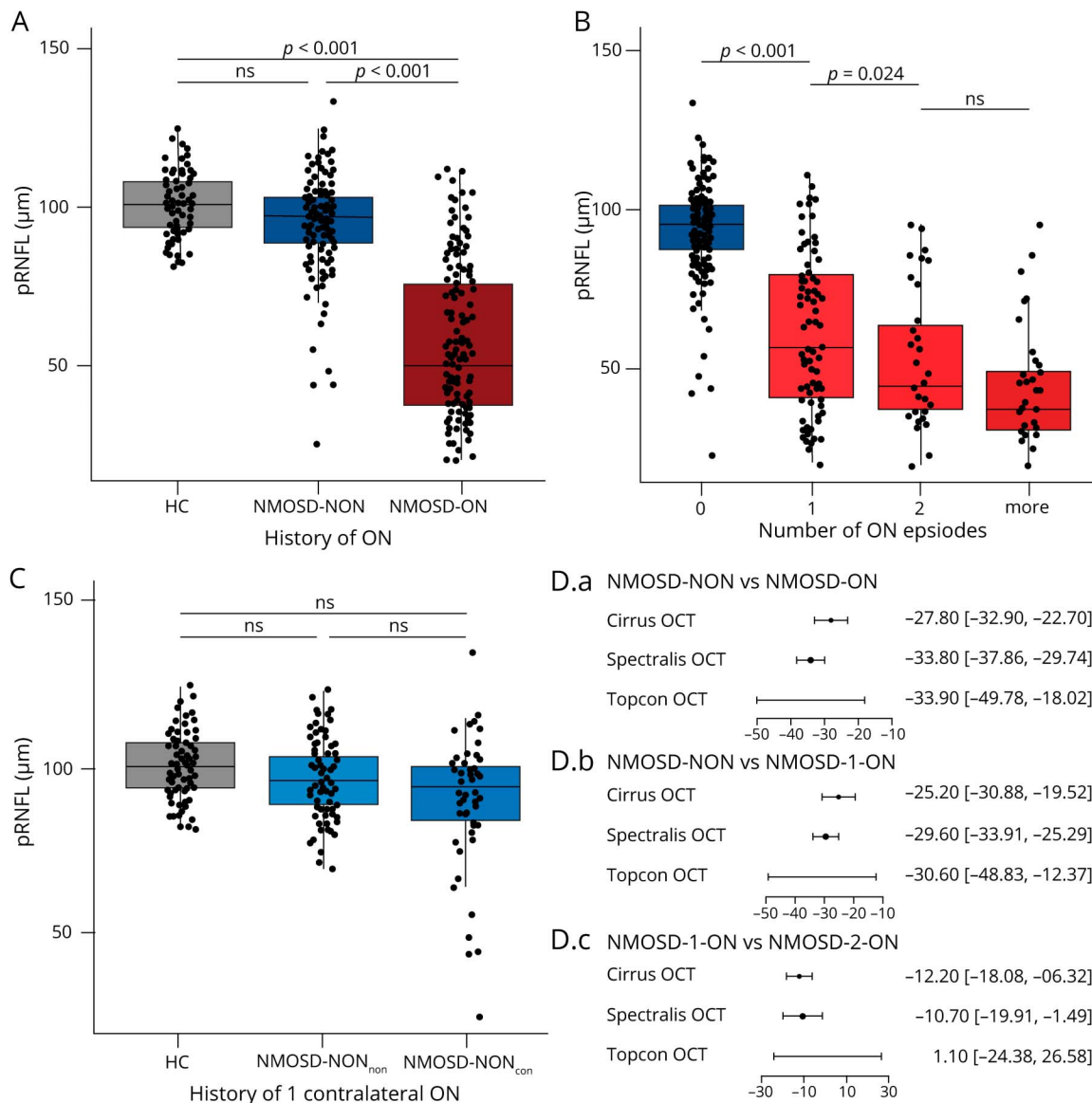
combined and individually for Spectralis SD-OCT; the analyses of data acquired by Cirrus HD-OCT or Topcon 3D OCT-1 and for the matched subset are reported as online-only supplement.

### Data Availability

The data supporting the findings of this study are available within the article and from the corresponding author by reasonable request.



**Figure 2** Group Comparisons of pRNFL Thickness



Boxplots of pRNFL thicknesses acquired by Heidelberg SD-OCT [µm] with values of individual eyes (jitter) for (A) HC (gray/left), NMOSD-NON (dark blue/middle), and NMOSD-ON (dark red/right); for (B) number of ON episodes (NMOSD-NON dark blue/left, NMOSD-1-ON light red/left-middle, NMOSD-2-ON medium-red/right-middle, and NMOSD-≥3-ON medium-dark red/right); and for (C) HC (gray/left), NMOSD-NON<sub>non</sub> (light blue/middle), and NMOSD-NON<sub>con</sub> (blue/right). (D) Forest plots for results from different OCT devices for (D.a) NMOSD-NON vs NMOSD-ON, (D.b) NMOSD-NON vs NMOSD-1-ON, and (D.c) NMOSD-1-ON vs NMOSD-2-ON (eFigure 3 and eTable 3, [links.lww.com/NXI/A556](https://links.lww.com/NXI/A556) and [links.lww.com/NXI/A557](https://links.lww.com/NXI/A557)). HC = eyes of HCs; NMOSD-NON = eyes of patients with neuromyelitis optica without a history of optic neuritis; NMOSD-NON<sub>con</sub> = eyes of patients with neuromyelitis optica without a history of optic neuritis but a history of contralateral optic neuritis; NMOSD-NON<sub>non</sub> = eyes of patients with neuromyelitis optica without a history of ipsilateral or contralateral optic neuritis; NMOSD-ON = eyes of patients with neuromyelitis optica with a history of optic neuritis; NMOSD-1-ON = eyes of patients with neuromyelitis optica with a history of 1 optic neuritis episode; NMOSD-2-ON = eyes of patients with neuromyelitis optica with a history of 2 optic neuritis episodes; ON = optic neuritis; pRNFL = peripapillary retinal nerve fiber layer.

## Results

Five hundred one eyes of 283 AQP4-IgG-seropositive patients and 136 eyes of 72 HCs were included in the analysis (Table 1).

### Neuroaxonal Damage After ON

GCIP layer and pRNFL were reduced in NMOSD-ON compared with NMOSD eyes without a history of ON

(NMOSD-NON) and HC (GCIP layer:  $81.4 \pm 5.7$  µm, pRNFL:  $101.1 \pm 9.0$  µm) (Figures 1A and 2A; Tables 2 and 3). The absolute (GCIP layer:  $-22.7$  µm; pRNFL:  $-38.5$  µm) and relative (GCIP layer:  $-38.8\%$ ; pRNFL:  $-61.6\%$ ) loss in eyes with 1 ON episode (NMOSD-1-ON) compared with NMOSD-NON was higher than in eyes with 2 ON episodes (NMOSD-2-ON) compared with NMOSD-1-ON (GCIP layer absolute loss:  $-3.5$  µm, relative loss:  $-6.0\%$ ; pRNFL absolute loss:  $-9.1$  µm, relative loss:  $-14.5\%$ , n.s.). The loss in

**Table 2** Group Comparisons of GCIP Layer Thickness for Heidelberg SD-OCT

	No. of eyes	Thickness ( $\mu\text{m}$ , mean $\pm$ SD)	Comparison to	Absolute difference ( $\mu\text{m}$ , mean)	Relative difference (% mean)	B	SE	p Value	R <sup>2</sup> <sub>marg</sub>	R <sup>2</sup> <sub>cond</sub>	Combined p value
<b>NMOSD</b>	268	67.3 $\pm$ 13.6	HCS	-14.0	-20.8	15.6	2.0	<b>&lt;0.0001</b>	0.143	0.690	—
<b>NMOSD-ON</b>	124	57.4 $\pm$ 12.2		-24.0	-41.8	25.0	2.0	<b>&lt;0.0001</b>	0.482	0.839	—
<b>NMOSD-NON</b>	144	75.9 $\pm$ 7.7		-5.4	-7.2	5.8	1.4	<b>&lt;0.0001</b>	0.088	0.947	—
<b>NMOSD-NON<sub>non</sub></b>	99	77.3 $\pm$ 5.9		-4.0	-5.0	2.69	1.42	0.061	0.189	0.924	—
<b>NMOSD-NON<sub>con</sub></b>	45	72.9 $\pm$ 10.2		-8.5	-10.4	8.44	1.77	<b>&lt;0.0001</b>	0.260	0.985	—
<b>NMOSD-ON</b>	124	57.4 $\pm$ 12.2	NMOSD-NON	-18.5	-24.4	-15.8	1.1	<b>&lt;0.0001</b>	0.430	0.795	<b>5.9e<sup>-45</sup></b>
<b>NMOSD-1-ON</b>	76	58.6 $\pm$ 12.9		-17.3	-22.8	-14.5	1.2	<b>&lt;0.0001</b>	0.417	0.803	<b>1.4e<sup>-33</sup></b>
<b>NMOSD-2-ON</b>	28	55.1 $\pm$ 9.4	NMOSD-1-ON	-3.5	-6.0	-5.7	2.6	<b>0.028</b>	0.143	0.729	<b>0.038</b>
<b>NMOSD-<math>\geq</math>3-ON</b>	20	55.9 $\pm$ 12.8	NMOSD-2-ON	0.8	1.4	4.9	2.6	0.070	0.311	0.781	0.222
<b>NMOSD-NON<sub>con</sub></b>	45	72.9 $\pm$ 10.2	NMOSD-NON <sub>non</sub>	-4.4	-6.1	-3.8	1.7	<b>0.025</b>	0.111	0.701	<b>0.004</b>
<b>NMOSD-NON<sub>1-con</sub></b>	37	73.6 $\pm$ 9.4		-3.7	-5.1	-3.5	1.6	<b>0.037</b>	0.149	0.906	0.154

Abbreviations: AQP4-IgG = aquaporin-4 IgG; B = estimate; GCIP = ganglion cell and inner plexiform; NMOSD = AQP4-IgG-seropositive NMOSD; NMOSD-NON = AQP4-IgG-seropositive NMOSD eyes without a history of ON; NMOSD-NON<sub>non</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON or contralateral ON; NMOSD-NON<sub>con</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON but with a history of contralateral ON; NMOSD-NON<sub>1-con</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON but with a history of 1 contralateral ON; NMOSD-ON = AQP4-IgG-seropositive NMOSD eyes with a history of ON; NMOSD-1-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 1 ON; NMOSD-2-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 2 ONs; NMOSD- $\geq$ 3-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 3 or more ONs; NMOSD = neuromyelitis optica spectrum disorder; NMOSD-NON = NMOSD eyes without a history of ON; OCT = optical coherence tomography; ON = optic neuritis episode; Rcon = conditional R-squared; Rmarg = marginal R-squared; SE = standard error.

eyes with  $\geq$ 3-ON episodes (NMOSD-3-ON) was lower compared with NMOSD-2-ON (n.s.) (Figures 1B and 2B). Five NMOSD-NON eyes had pRNFL values  $<$  60  $\mu\text{m}$ ; these patients had no relevant comorbidities but a history of contralateral ON.<sup>17,18</sup> In sensitivity analyses to account for device differences, all significant analyses within the NMOSD cohort were confirmed for data acquired by Cirrus and Topcon OCT devices (eFigures 2 and 3; eTables 3 and 4, links.lww.com/NXI/A556; links.lww.com/NXI/A557).

### Neuroaxonal Damage Without ON

NMOSD-NON eyes had a reduced GCIP layer (75.9  $\pm$  7.7  $\mu\text{m}$ ,  $p <$  0.001), but not pRNFL (95.3  $\pm$  14.4  $\mu\text{m}$ ) compared with HC (GCIP layer: 81.4  $\pm$  5.7  $\mu\text{m}$ , pRNFL: 101.1  $\pm$  9.0  $\mu\text{m}$ ; Figures 1A and 2A; Tables 2 and 3). By comparison, only 28 NMOSD-NON eyes (5.1%) had a GCIP layer  $\leq$  the 5th percentile of HC. GCIP layer was also reduced in NMOSD-NON with a history of contralateral ON (72.9  $\pm$  10.2  $\mu\text{m}$ ) compared with NMOSD-NON without a history of contralateral ON (77.3  $\pm$  5.9  $\mu\text{m}$ ,  $p =$  0.025, NMOSD-NON<sub>non</sub>). This effect vanished when only considering NMOSD-NON<sub>con</sub> with a history of 1 contralateral ON (73.6  $\pm$  9.4  $\mu\text{m}$ ). However, only NMOSD-NON<sub>con</sub> ( $p <$

0.001) but not NMOSD-NON<sub>non</sub> ( $p =$  0.061) had thinned GCIP layer compared with HC (Figures 1C and 2C).

### INL Changes

INL was thicker in NMOSD-ON compared with HC (39.4  $\pm$  2.6  $\mu\text{m}$ ) and NMOSD-NON (eTable 4, links.lww.com/NXI/A557). Specifically, INL was thicker in eyes with 1 ON episode compared with NMOSD-NON but did not differ between eyes with different numbers of ON episodes. Also, INL did not differ between NMOSD-NON<sub>con</sub> (38.9  $\pm$  3.2  $\mu\text{m}$ ) and NMOSD-NON<sub>non</sub> (38.5  $\pm$  3.2  $\mu\text{m}$ ,  $p =$  0.931). In all patients, INL thickness (as the dependent variable in the linear mixed model described above) was correlated with thinner GCIP layer (B = -0.11, standard error [SE] = 0.01,  $p <$  0.001) and pRNFL (B = -0.06, SE = 0.01,  $p <$  0.001). This correlation existed in NMOSD-ON (GCIP layer: B = -0.08, SE = 0.03,  $p =$  0.005; pRNFL: B = -0.06, SE = 0.01,  $p <$  0.001) and not present in NMOSD-NON (eFigure 4, links.lww.com/NXI/A556).

Scans of 363 NMOSD eyes were clearly suitable for MME investigations (high quality, eFigure 4C, links.lww.com/NXI/A556). MMEs were visible in 24 (6.6%) eyes of 21 patients.

**Table 3** Group Comparisons of pRNFL Thickness With (A) Spectralis SD-OCT Data for Comparisons vs HC (B) Data From All Devices for Inpatient Cohort Comparisons

	No. of eyes	Thickness ( $\mu\text{m}$ , mean $\pm$ SD)	Comparison to	Absolute difference ( $\mu\text{m}$ , mean)	Relative difference (% mean)	B	SE	p Value	R <sup>2</sup> <sub>margin</sub>	R <sup>2</sup> <sub>cond</sub>	Combined p value
<b>NMOSD</b>	344	76.3 $\pm$ 27.4	HCs	-14.0	-32.6	24.2	4.3	<b>&lt;0.0001</b>	0.143	0.690	—
<b>NMOSD-ON</b>	170	56.9 $\pm$ 23.7		-44.3	-77.9	43.1	4.1	<b>&lt;0.0001</b>	0.482	0.839	—
<b>NMOSD-NON</b>	174	95.3 $\pm$ 14.4		-5.9	-6.2	4.6	3.0	0.129	0.088	0.947	—
<b>NMOSD-NON<sub>non</sub></b>	116	97.4 $\pm$ 10.9		-3.7	-3.7	-2.9	2.2	0.204	0.107	0.893	—
<b>NMOSD-NON<sub>con</sub></b>	58	91.0 $\pm$ 19.1		-4.7	-4.6	7.1	4.2	0.097	0.100	0.975	—
<b>NMOSD-ON</b>	170	56.9 $\pm$ 23.7	NMOSD-NON	-38.4	-40.3	-33.8	2.1	<b>&lt;0.0001</b>	0.436	0.741	<b>4.4e<sup>-60</sup></b>
<b>NMOSD-1-ON</b>	97	62.6 $\pm$ 24.9		-32.7	-34.3	-29.6	2.2	<b>&lt;0.0001</b>	0.379	0.712	<b>4.6e<sup>-40</sup></b>
<b>NMOSD-2-ON</b>	35	53.5 $\pm$ 21.3	NMOSD-1-ON	-9.1	-14.5	-10.7	4.7	<b>0.024</b>	0.087	0.528	<b>0.010</b>
<b>NMOSD-<math>\geq</math>3-ON</b>	38	45.2 $\pm$ 17.8	NMOSD-2-ON	-8.3	-15.5	-0.4	4.3	0.924	0.087	0.719	0.838
<b>NMOSD-NON<sub>con</sub></b>	58	91.0 $\pm$ 19.1	NMOSD-NON <sub>non</sub>	-6.5	-7.1	-4.0	2.9	0.173	0.061	0.939	0.157
<b>NMOSD-NON<sub>1-con</sub></b>	47	93.2 $\pm$ 18.1		-4.2	-4.6	-3.2	2.9	0.276	0.056	0.925	0.616

Abbreviations: AQP4-IgG = aquaporin-4 IgG; NMOSD = AQP4-IgG-seropositive NMOSD; NMOSD-NON = AQP4-IgG-seropositive NMOSD eyes without a history of ON; NMOSD-NON<sub>non</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON or contralateral ON; NMOSD-NON<sub>con</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON but with a history of contralateral ON; NMOSD-NON<sub>con</sub> = AQP4-IgG-seropositive NMOSD eyes without a history of ON but with a history of 1 contralateral ON; NMOSD-ON = AQP4-IgG-seropositive NMOSD eyes with a history of ON; NMOSD-1-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 1 ON; NMOSD-2-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 2 ONs; NMOSD- $\geq$ 3-ON = AQP4-IgG-seropositive NMOSD eyes with a history of 3 or more ONs; NMOSD = neuromyelitis optica spectrum disorder; NMOSD-NON = NMOSD eyes without a history of ON; OCT = optical coherence tomography; ON = optic neuritis episode; pRNFL = peripapillary retinal nerve fiber layer; VEP = visually evoked potential.

Twenty-three eyes (13.1%) of NMOSD-ON and 1 eye (0.5%) of NMOSD-NON (with a history of contralateral ON) were affected. The number of ON episodes did not influence the incidence of MMEs (NMOSD-1-ON N = 16 (14%), NMOSD-2-ON N = 4 (11.4%), NMOSD- $\geq$ 3-ON N = 3 (11.1%)).

Including only the most frequent ethnicities in our data set (Asian and non-Hispanic White), results did not differ, and no ethnicity was singled out regarding its pattern of injury (data not shown).

### OCT and Vision Loss

HC-VA was numerically reduced in NMOSD (0.25  $\pm$  0.48) compared with HC (-0.01  $\pm$  0.08) and known healthy reference values. NMOSD-ON also had reduced HC-VA (0.44  $\pm$  0.58) compared with NMOSD-NON (0.04  $\pm$  0.20,  $p < 0.001$ ,  $p = 0.401$ ) and HC/healthy reference populations. HC-VA was correlated with GCIP layer (B = -0.016, SE = 0.002,  $p < 0.001$ ) and pRNFL thicknesses (B = -0.010, SE = 0.001,  $p < 0.001$ ) in AQP4-IgG-seropositive NMOSD. AQP4-IgG-seropositive NMOSD eyes with prolonged VEP latency had a thinned GCIP

layer (B = -11.647, SE = 3.628,  $p = 0.002$ ) and pRNFL (B = -21.965, SE = 3.724,  $p < 0.001$ ). EDSS score as a metric of global disability was inversely correlated with GCIP layer (B = -1.370, SE = 0.566,  $p = 0.017$ ) and pRNFL thicknesses (B = -3.148, SE = 1.080,  $p = 0.004$ ).

## Discussion

Our study specifies a severe and functionally relevant decrease of GCIP layer and pRNFL in NMOSD-ON compared with NMOSD-NON and HCs in AQP4-IgG-seropositive patients. Neuroaxonal damage is particularly large from the first episode of ON, where contribution to retinal damage in subsequent episodes of ON is still considerable but smaller. In contrast to previous smaller studies, the current study ascertains GCIP layer but not pRNFL thinning in NMOSD-NON compared with HC.<sup>6,19</sup> This effect was driven by eyes with contralateral ON and not statistically significant in eyes without a history of ON. INL was thicker in NMOSD-ON and was inversely correlated with GCIP layer. Of note, 13.1%



of NMOSD-ON eyes showed MME indicative of secondary inflammatory changes.<sup>20,21</sup> This investigation overcame limits of small samples and OCT data heterogeneity of earlier studies through use of an international consortium of NMOSD specialists. This framework substantiated the CROCTINO study—a large, multicenter, retrospective evaluation of retinal pathology in NMOSD using OCT.

The neuroaxonal degeneration in NMOSD-ON demonstrated here is substantially greater than reported changes in multiple sclerosis (MS), a common differential diagnosis.<sup>2,22</sup> In a meta-analysis, an average pRNFL loss of 20  $\mu\text{m}$  was estimated in MS after ON, which is nearly 2-fold higher in our NMOSD-ON cohort ( $-38.4 \mu\text{m}$ ).<sup>22</sup> For GCIP layer, our data suggested approximately 1.5-fold higher loss in NMOSD ( $-24.0 \mu\text{m}$ ) compared with MS.<sup>22</sup> These differences not only result from a higher ON frequency, but may also be caused by more severe retinal damage in NMOSD after a singular ON.<sup>23</sup> It is intriguing that the damage is particularly large after the first ON episode with smaller losses after subsequent episodes, which might be due to less neuroaxonal content in subsequent episodes.<sup>2,24</sup> Alternatively, although the analysis of treatment effects exceeds the scope of this study, it is possible that the longer time to effective anti-inflammatory therapy and the typical choice of less effective therapies (e.g., steroids instead of plasma exchange) at the first attack compared with following attacks may significantly contribute to this difference. Independent of the number of ONs, the percent loss is smaller in GCIP layer compared with pRNFL—pointing toward either (1) stronger loss of retinal nerve fibers than retinal ganglion cells, (2) impairment of ganglion cells not leading to extinction but axonal loss; (3) a larger amount of non-neuronal tissue in GCIP layer, or (4) RNFL loss in the periphery beyond the macular area measured by GCIP layer. These hypotheses are not mutually exclusive, and each might contribute to the effect. Consistent with these concepts, excessive vision loss relative to neuroaxonal content and VEP latency in AQP4-IgG-seropositive patients compared with MOG-IgG-seropositive and MS patients implies damage of the peripheral retina and optic nerve tissues, which are not reflected in the macula and pattern VEP measurements.<sup>4</sup>

Whether attack-independent neurodegeneration in AQP4-IgG-seropositive NMOSD occurs has been controversial.<sup>2,6,7,25-28</sup> The current study identifies decreased GCIP layer in NMOSD-NON compared with HC, but not pRNFL. Such subclinical changes could be caused by contralateral involvement after unilateral ON.<sup>17,18</sup> Indeed, our cohort suggests pronounced neurodegeneration in eyes with contralateral ON. However, eyes of patients without ON also exhibit a trend for thinner GCIP layer compared with HCs. Underscored by longitudinal studies showing ON-independent neurodegeneration and VEP latency prolongation,<sup>7,29</sup> such patterns suggest at least 2 mechanisms of injury: (1) a primary retinopathy or optic neuropathy in context of an astrocytopathy or caused by direct damage to AQP4-expressing cells such as astrocytes

and Müller cells by either AQP4-IgG or AQP4-specific T cells or (2) a global or afferent visual system specific chronic or episodic neurodegenerative process. Because lesions often spare the brain, and most studies outside of predisposed areas such as optic nerve and spinal cord failed to detect effects,<sup>30</sup> a constant global involvement seems unlikely. Several studies described changes affecting AQP4 expression and astrocytic end feet,<sup>31</sup> parafoveal changes in agreement with an involvement of AQP4-expressing Müller cell,<sup>6,26,27,32</sup> and attack-independent spinal cord atrophy.<sup>33</sup> These findings are consistent with tissue infiltration by AQP4-IgG-specific T cells<sup>34</sup> and the attack-independent loss of retinal ganglion cells<sup>7</sup>—further supporting the existence of an ON-independent pathology, which might be restricted to the main disease foci. The latter hypotheses could be addressed by region-specific pathology or advanced imaging studies.

INL changes have been suggested as a marker of neuroinflammation and potential treatment response in MS.<sup>21,35,36</sup> MME may develop as a consequence of neurodegeneration or other—non-disease-specific—processes.<sup>35</sup> Patients with NMOSD were described to have INL thickening and MME.<sup>20,37</sup> In our cohort, 13.1% of NMOSD-ON eyes were affected by MME, which is higher than the 2%–5% described in MS but comparable to incidences in NMOSD described before by Gelfand and colleagues.<sup>35,37</sup> However, INL thickening itself remained comparable to changes reported in MS.<sup>35</sup> This disparity could reflect a disrupted fluid homeostasis due to Müller cell involvement or loss of content of the INL with a parallel inflammatory reaction and development of MME. The limited accessibility of MMEs, especially in severely affected eyes due to limited image quality, has hindered their detailed assessments and most likely leads to underestimation of their incidence.<sup>38</sup> We demonstrate that the INL thickness is inversely correlated with neuroaxonal content and could be a valuable marker of disease severity also in NMOSD.

Our study cohort was representative of patients with AQP4-IgG-seropositive NMOSD with respect to a female predominance and ON history.<sup>11,39</sup> This OCT study included multiple ethnicity backgrounds, although the distribution was shifted toward Caucasian/White patients and other ethnicities were underrepresented (e.g., Hispanic White and Black), limiting the generalizability of results.<sup>40</sup> Patients from different heritages presented similar findings.

The current study was based on source data instead of a meta-analysis. Using the Guthy-Jackson Charitable Foundation network, this multicentric study was conducted without investigator reimbursement and illustrated how collaboration integrating international perspectives can produce meaningful results. To overcome technical challenges of heterogeneous source data, we developed novel OCT postprocessing techniques allowing us to perform standardized analyses and enabling the uniform analysis of the largest OCT image data set in NMOSD to date.<sup>10</sup> Thus, the strengths of CROCTINO include its established

infrastructure, large international network of experts representing multiple ethnicities and geographic regions, and the use of state-of-the-art OCT postprocessing techniques.<sup>41</sup>

We recognize limitations of the current investigation: The retrospective and heterogeneous data acquisition might have led to biases and impreciseness beyond the ability of our quality control. We addressed this by excluding uncertain cases. HCs were only included from a limited number of centers. The unbalanced data set limited some analyses, such as the influence of ethnicities or acute and disease-modifying treatments. Case-control matching was impossible, particularly with respect to subclinical progression dependent or independent of ON and to ethnicity. Similarly, comparisons with other NMOSD subtypes or MS were beyond the scope of this study. Longitudinal data, acute ON data, and AQP4-IgG-seronegative and MOG-IgG-seropositive patient data are part of the CROCTINO data set and will be analyzed in the future. Also, OCT data processing was performed by multiple raters potentially introducing interrater variability. MRI data and posterior visual pathway involvement were not investigated in this study. However, the current study achieved an unprecedented worldwide assessment of retinal damage in AQP4-IgG-seropositive NMOSD.

To conclude, AQP4-IgG-seropositive NMOSD is characterized by a severe, functionally relevant retinal neurodegeneration as a consequence of ON. Although the majority of damage occurs during the first episode, there is cumulative loss with each succeeding relapse. The ON-associated damage is not limited to the neuroaxonal content but can also induce—likely inflammation-mediated—INL increase and occurrence of MME. Our data also suggest attack-independent retinal damage in AQP4-IgG-seropositive NMOSD. Our study supports that attack prevention is key in avoiding neuroaxonal damage and vision loss in patients with NMOSD. It further suggests that the first ON episode causes the most damage, where only some patients with then established diagnosis will be on immunosuppressive therapy. This highlights the need for effective therapies that can ameliorate an ongoing attack or regenerate attack-generated damage, which is an unmet clinical need. Last, the study emphasizes the utility of OCT as a sensitive structural metric and its potential for monitoring progression and even treatment response in AQP4-IgG-seropositive NMOSD. The international CROCTINO program provides an unprecedented opportunity to apply OCT in a standardized manner to assess pathophysiology, clinical course, and therapeutic efficacy in NMOSD.

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<b>Svenja Specovius, MSc</b>	Experimental and Clinical Research Center, Max Delbrück Center for Molecular Medicine and Charité-Universitätsmedizin, Corporate Member of Freie Universität Berlin and Humboldt-Universität zu Berlin, Germany	Acquisition and analysis of data; conception and design of the study
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<b>Claudia Chien, PhD</b>	Experimental and Clinical Research Center, Max Delbrück Center for Molecular Medicine and Charité-Universitätsmedizin, Corporate Member of Freie Universität Berlin and Humboldt-Universität zu Berlin, Germany	Acquisition and analysis of data
<b>Seyedamirhosein Motamedi, PhD</b>	Experimental and Clinical Research Center, Max Delbrück Center for Molecular Medicine and Charité-Universitätsmedizin, Corporate Member of Freie Universität Berlin and Humboldt-Universität zu Berlin, Germany	Acquisition and analysis of data
<b>Charlotte Bereuter, BSc</b>	Experimental and Clinical Research Center, Max Delbrück Center for Molecular Medicine and Charité-Universitätsmedizin, Corporate Member of Freie Universität Berlin and Humboldt-Universität zu Berlin, Germany	Acquisition and analysis of data

## Appendix (continued)

Name	Location	Contribution
<b>Lawrence Cook, PhD</b>	University of Utah, Salt Lake City, UT	Acquisition and analysis of data and conception and design of the study
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<b>Mariana Andrade Fontanelle, MD</b>	University of Minas Gerais, Medical School, Belo Horizonte, Brazil	Acquisition and analysis of data
<b>Ho Jin Kim, MD, PhD</b>	National Cancer Center, Goyang, Republic of Korea	Acquisition and analysis of data
<b>Jae-Won Hyun, MD, PhD</b>	University of Minas Gerais, Medical School, Belo Horizonte, Brazil	Acquisition and analysis of data
<b>Jacqueline Palace, MD</b>	Oxford University Hospitals, National Health Service Trust, Oxford, United Kingdom	Acquisition and analysis of data
<b>Adriana Roca-Fernandez, MSc</b>	Oxford University Hospitals, National Health Service Trust, Oxford, United Kingdom	Acquisition and analysis of data
<b>Maria Isabel Leite, MD, PhD</b>	Oxford University Hospitals, National Health Service Trust, Oxford, United Kingdom	Acquisition and analysis of data
<b>Srilakshmi Sharma, MD, PhD</b>	Oxford University Hospitals, National Health Service Trust, Oxford, United Kingdom	Acquisition and analysis of data
<b>Fereshteh Ashtari, MD</b>	Isfahan University of Medical Sciences, Isfahan, Iran	Acquisition and analysis of data
<b>Rahele Kafieh, PhD</b>	Isfahan University of Medical Sciences, Isfahan, Iran	Acquisition and analysis of data
<b>Alireza Dehghani, PhD</b>	Isfahan University of Medical Sciences, Isfahan, Iran	Acquisition and analysis of data
<b>Mohsen Pourazizi, PhD</b>	Isfahan University of Medical Sciences, Isfahan, Iran	Acquisition and analysis of data
<b>Lekha Pandit, MD, PhD</b>	KS Hegde Medical Academy, Nitte University, Mangalore, India	Acquisition and analysis of data
<b>Anitha D'Cunha, PhD</b>	KS Hegde Medical Academy, Nitte University, Mangalore, India	Acquisition and analysis of data
<b>Orhan Aktas, MD</b>	Heinrich Heine University Düsseldorf, Düsseldorf, Germany	Acquisition and analysis of data
<b>Marius Ringelstein, MD</b>	Heinrich Heine University Düsseldorf, Düsseldorf, Germany	Acquisition and analysis of data
<b>Philipp Albrecht, MD</b>	Heinrich Heine University Düsseldorf, Düsseldorf, Germany	Acquisition and analysis of data
<b>Eugene May, MD</b>	Swedish Neuroscience Institute Neuro-Ophthalmology, Seattle, WA	Acquisition and analysis of data
<b>Caryl Tongco</b>	Swedish Neuroscience Institute Neuro-Ophthalmology, Seattle, WA	Acquisition and analysis of data

## Appendix (continued)

Name	Location	Contribution
<b>Letizia Leocani, MD, PhD</b>	Institute of Experimental Neurology, Scientific Institute Hospital San Raffaele and University Vita-Salute San Raffaele, Milan, Italy	Acquisition and analysis of data
<b>Marco Pisa, MD</b>	Institute of Experimental Neurology, Scientific Institute Hospital San Raffaele and University Vita-Salute San Raffaele, Milan, Italy	Acquisition and analysis of data
<b>Marta Radaelli, MD, PhD</b>	Institute of Experimental Neurology, Scientific Institute Hospital San Raffaele and University Vita-Salute San Raffaele, Milan, Italy	Acquisition and analysis of data
<b>Elena H Martinez-Lapiscina, MD, PhD</b>	Hospital Clinic of Barcelona-Institut d'Investigacions, Biomèdiques August Pi Sunyer, Barcelona, Spain	Acquisition and analysis of data
<b>Hadas Stiebel-Kalish, MD</b>	Sackler School of Medicine, Tel Aviv University, Tel Aviv, Israel	Acquisition and analysis of data
<b>Sasitorn Siritho, MD</b>	Siriraj Hospital and Bumrungrad International Hospital, Bangkok, Thailand	Acquisition and analysis of data
<b>Jérôme de Seze, MD, PhD</b>	University Hospital of Strasbourg, France	Acquisition and analysis of data
<b>Thomas Senger, MD</b>	University Hospital of Strasbourg, France	Acquisition and analysis of data
<b>Joachim Havla, MD</b>	Ludwig-Maximilians Universität Muenchen, Munich, Germany	Acquisition and analysis of data
<b>Romain Marignier, MD, PhD</b>	Pierre Wertheimer Neurological Hospital, Hospices Civils de Lyon, France	Acquisition and analysis of data
<b>Alvaro Cobo Calvo, MD, PhD</b>	Pierre Wertheimer Neurological Hospital, Hospices Civils de Lyon, France; Hospital Universitari Vall d'Hebron, Universitat Autònoma de Barcelona, Spain	Acquisition and analysis of data
<b>Denis Bichuetti, MD, PhD</b>	Escola Paulista de Medicina, Universidade Federal de São Paulo, São Paulo, Brazil	Acquisition and analysis of data
<b>Ivan Maynart Tavares, MD, PhD</b>	Escola Paulista de Medicina, Universidade Federal de São Paulo, São Paulo, Brazil	Acquisition and analysis of data
<b>Nasrin Asgari, MD, PhD</b>	University of Southern Denmark, Odense, Denmark	Acquisition and analysis of data
<b>Kerstin Soelberg, MD</b>	University of Southern Denmark, Odense, Denmark	Acquisition and analysis of data
<b>Ayşe Altintas, MD</b>	Cerrahpasa Medical Faculty, Istanbul University, Cerrahpasa, Turkey	Acquisition and analysis of data
<b>Rengin Yildirim, MD</b>	Cerrahpasa Medical Faculty, Istanbul University, Cerrahpasa, Turkey	Acquisition and analysis of data

Continued



## Appendix (continued)

Name	Location	Contribution
<b>Uygun Tanriverdi, MD</b>	Cerrahpasa Medical Faculty, Istanbul University, Cerrahpasa, Turkey	Acquisition and analysis of data
<b>Anu Jacob, MD</b>	The Walton Centre for Neurology and Neurosurgery, Liverpool, United Kingdom	Acquisition and analysis of data
<b>Saif Huda, MD, PhD</b>	The Walton Centre for Neurology and Neurosurgery, Liverpool, United Kingdom	Acquisition and analysis of data
<b>Zoe Rimler, BSc</b>	NYU School of Medicine, New York, NY	Acquisition and analysis of data
<b>Allyson Reid, MD</b>	NYU School of Medicine, New York, NY	Acquisition and analysis of data
<b>Yang Mao-Draayer, MD, PhD</b>	University of Michigan Medical School, Ann Arbor, MI	Acquisition and analysis of data
<b>Ibis Soto de Castillo, MD</b>	Hospital Clínico de Maracaibo, Maracaibo, Venezuela	Acquisition and analysis of data
<b>Axel Petzold, MD, PhD</b>	University College London, London, UK	Conception and design of the study
<b>Ari J Green, MD</b>	University of California San Francisco, CA	Conception and design of the study
<b>Michael R Yeaman, MD, PhD</b>	University of California Los Angeles, CA	Conception and design of the study
<b>Terry Smith, MD</b>	University of Michigan Medical School, Ann Arbor, MI	Conception and design of the study
<b>Alexander U Brandt, MD</b>	Experimental and Clinical Research Center, Max Delbrück Center for Molecular Medicine and Charité-Universitätsmedizin Berlin, Berlin, Germany	Acquisition and analysis of data, conception and design of the study, and drafting a significant portion of the manuscript or figures
<b>Friedemann Paul, MD</b>	Charité-Universitätsmedizin Berlin, Corporate Member of Freie Universität Berlin and Humboldt-Universität zu Berlin	Acquisition and analysis of data and conception and design of the study

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## Retinal Optical Coherence Tomography in Neuromyelitis Optica

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In the Article “Retinal Optical Coherence Tomography in Neuromyelitis Optica” by Oertel et al.<sup>1</sup>, the 37th author should be listed as “Alvaro Cobo-Calvo.” The authors regret the error.

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1. Oertel FC, Specovius S, Zimmermann HG, et al. Retinal optical coherence tomography in neuromyelitis optica. *Neurol Neuroimmunol Neuroinflamm*. 2021;8(6):e1068.