

Impact of Dietary Intervention on Serum Neurofilament Light Chain in Multiple Sclerosis

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Abstract

Background and Objectives

Adapted ketogenic diet (AKD) and caloric restriction (CR) have been suggested as alternative therapeutic strategies for multiple sclerosis (MS), but information on their impact on neuroaxonal damage is lacking. Thus, we explored the impact of diets on serum neurofilament light chain (sNfL) levels in patients with relapsing-remitting MS.

Methods

We retrospectively evaluated a prospective randomized controlled trial of 60 patients with MS who were on a common diet or ketogenic diet or fasting. We examined sNfL levels of 40 participants at baseline and at the end of the study after 6 months using single molecule array assay.

Results

sNfL levels were investigated in 9 controls, 14 participants on CR, and 17 participants on AKD. Correlation analysis showed an association of sNfL with age and disease duration; an association was also found between sNfL and the Multiple Sclerosis Functional Composite. AKD significantly reduced sNfL levels at 6 months compared with the common diet group ($p = 0.001$).

Discussion

For clinical or study use, consider that AKD may incline sNfL levels independent of relapse activity up to 3 months after initiation. At 6 months, AKD, which complements current therapies, reduced sNfL levels, therefore suggesting potential neuroprotective effects in MS. A single cycle of seven-day fasting did not affect sNfL. AKD may be an addition to the armamentarium to help clinicians support patients with MS in a personalized manner with tailored diet strategies.

Trial Registration Information

Clinical trial registration number NCT01538355.

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Glossary

AKD = adapted ketogenic diet; **BMI** = body mass index; **CD** = common diet; **CR** = caloric restriction; **DMT** = disease-modifying therapy; **MS** = multiple sclerosis; **MSFC** = Multiple Sclerosis Functional Composite; **PASAT** = Paced Auditory Serial Addition Test; **SiMoA** = single molecule assay; **sNfL** = serum neurofilament light chain.

In multiple sclerosis (MS), effective therapeutic strategies and sensitive biomarkers to evaluate drug efficacy and disease course are urgently needed. Serum neurofilament light chain protein (sNfL) has recently been suggested as a promising candidate for a reliable, easy-to-use biomarker of neuroaxonal damage that accurately detects changes over both long and short time intervals in MS.^{1,2} After axonal injury, NfL is elevated in both the CSF and in the peripheral blood, where it can be measured by highly sensitive single molecule assays (SiMoA).^{3,4} Studies have suggested a possible role of caloric restriction (CR) and adapted ketogenic diet (AKD) on neuroinflammation in MS and other neurologic disorders.⁵⁻⁷ The observed diet-induced improvement of neuronal resistance and axonal survival are clearly neuroprotective, but the underlying mechanisms remain elusive.^{5,8} In the present study, we investigated whether AKD and CR in comparison to common diet (CD) may affect neurodegeneration as measured by sNfL levels in MS.

Methods

Clinical Trial Design, Patients, Study Setting, Interventions, and Primary Outcome

All study details were described previously^{6,7,9} and are presented in eMethods, links.lww.com/NXI/A653. This study was a three-armed parallel group, single-centered, controlled and randomized clinical trial. This study was registered at clinicaltrials.gov as NCT01538355.

Outcome Measure

The sNfL concentration of 40 patients was quantified in 2020/2021.

sNfL Measurements

We applied the same method as described recently³ and presented in eMethods, links.lww.com/NXI/A653. In brief, sNfL was measured in several rounds by SiMoA HD-1 (Quanterix) using the NF-Light Advantage Kit (Quanterix) from the same batch according to the manufacturer's instructions. sNfL analysis was performed in a blinded fashion without clinical information about patients.

Control Diet

Patients on CD met the criteria of a common diet in the German population as described in the National Nutrition Survey II (mri.bund.de/de/institute/ernaehrungsverhalten/publikationen/forschungsprojekte/nvsii/).

Caloric Restriction

As previously described, a single cycle of 7-day CR (200–350 kcal/d) was performed at study outset; a 3-day stepwise

reintroduction to an isocaloric common diet was performed starting on day 8.⁶

Adapted Ketogenic Diet

Patients followed an AKD for 6 months from study outset. We recommended an average daily intake of <50 g carbohydrates, > 160 g fat (omega 6 vs omega 3 ratio 2:1), and average protein intake ≤100 g per day. Compliance during CR and AKD was monitored via ketosis self-measured in blood and urine.

Standard Protocol Approvals, Registrations, and Patient Consents

This study was approved by the local ethics committee in the frame of the IGEL study (ethics no. EA1/105/11). All patients included in this study signed specific informed consent.

Statistical Analysis

We used analyses of covariance to test for group differences using sNfL baseline data, number of clinical relapses over the course of the study, and body mass index (BMI) as covariates. The Wilcoxon matched-pairs signed-rank test analysis was used for intragroup statistics.

Data Availability

Anonymized data will be made available on request from any qualified investigator.

Results

Situation Before Dietary Intervention

Baseline Intergroup Comparison: No Differences in Outcome

Patient demographics did not differ significantly when the different intervention groups were compared (Table 1). The use of the disease-modifying therapy (DMT) glatiramer acetate was significantly different between the groups, but this is clinically not relevant. sNfL levels did not differ significantly between the 3 groups before dietary interventions. These data indicate that the dietary habits of the patients involved in the study did not influence the sNfL concentration at baseline.

Next, we correlated disease-related and participant-reported parameters across all groups involving all individuals of all experimental groups. We found a significant association between sNfL and age ($r = 0.347$, $p < 0.05$) and disease duration ($r = 0.313$, $p < 0.05$) confirming observations of

Table 1 Baseline Characteristics and Associations

Baseline characteristics	CD (n = 9)	SD IQR	CR (n = 14)	SD IQR	AKD (n = 17)	SD IQR	^a p Value					
Age, y	50	6	45.71	10.5	43.06	9.7	0.3					
Sex, F/M	6/3 (67/33)		12/2 (86/14)		13/4 (76/24)		0.6					
Expanded Disability Status Scale score	3.0	1.75–4	3.5	2.4–4.0	3.0	2.5–3.5	0.5					
Disease duration, y	11.0	10.3	10.6	7.8	7.1	5.3	0.3					
Relapse rate 12 mo prior study outset	0.22	0.4	0.26	0.4	0.47	0.5	0.3					
No immune modulating drugs	1 (11)		2 (14)		5 (29)		0.4					
Glatiramer acetate	7 (78)		4 (29)		4 (24)		0.02					
Interferon beta	0		6 (43)		4 (24)		0.07					
Fingolimod	0		1 (7)		3 (18)		0.3					
Natalizumab	1 (11)		1 (7)		1 (6)		0.9					
BMI, kg/m ²	26.6	7.3	26.8	4.7	26.3	4.9	0.7					
Percent body fat	36.2	11.3	37.1	9.6	36.3	9.5	0.9					
Fasting blood sugar, mmol/L	4.6	0.5	4.5	0.8	4.4	0.5	0.9					
β-hydroxybutyrate, μmol/L	105.9	72.1	111.3	118.3	112.4	100.5	0.6					
MSFC-3	0.17	0.7	-0.13	0.7	-0.02	0.9	0.3					
MSFC-2	0.34	0.8	-0.08	0.6	-0.14	0.9	0.05					
PASAT-3	49.56	11.3	40.5	12.5	46.79	10	0.09					
sNfL, ng/mL	8.9	3.6	10.3	3.6	8.5	2.0	0.3					
Associations	Age	Disease Duration	PASAT-3	MSFC-3	MSFC-2	BMI	EDSS score	MS-54 PH	MS-54 MH	FSS score	MFIS score	BDI
sNfL												
Spearman rho	0.347	0.313	-0.406	-0.317	-0.298	-0.133	0.013	0.099	0.031	-0.122	-0.109	-0.186
p (2 tailed)	0.01	0.02	0.004	0.03	0.04	0.41	0.93	0.55	0.84	0.41	0.47	0.22

Abbreviations: AKD = adapted ketogenic diet; BMI = body mass index; CD = common diet; CR = caloric restriction diet; MSFC = Multiple Sclerosis Functional Composite; PASAT = Paced Auditory Serial Addition Test; sNfL = serum neurofilament light chain. Baseline intergroup analysis: data are presented as mean (SD), number (%), or median (interquartile range, IQR), as appropriate. Baseline data were available for 40 patients.

^a Nonparametric Kruskal-Wallis test to compare between all 3 groups.

Abbreviations: BDI = Beck Depression Inventory; BMI = body mass index; EDSS = Expanded Disability Status Scale; FSS = Fatigue Severity Scale; MFIS = Modified Fatigue Impact Scale; MSFC = Multiple Sclerosis Functional Composite; PASAT = Paced Auditory Serial Addition Test; sNfL = serum neurofilament light chain.

Baseline cross-group correlation: Analysis between sNfL and disease-related or subject-reported parameters.

Multiple Sclerosis Quality of Life-54 physical/mental health (MS-54 PH/MH).

Bold indicates significance, defined as $p < 0.05$.

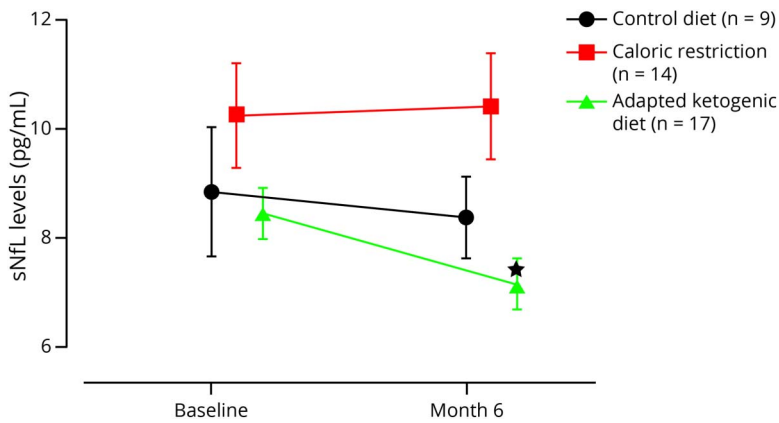
other groups. Moreover, we observed a significant inverse correlation between sNfL and the Multiple Sclerosis Functional Composite 2 and 3 (MSFC 2 + 3) ($r = -0.298$, $p < 0.05$; $r = -0.317$, $p < 0.05$) and the Paced Auditory Serial Addition Test 3 ($r = -0.298$, $p < 0.005$) (Table 1; eFigure 1–eFigure 5, links.lww.com/NXI/A653). Of interest, sNfL levels did not significantly correlate with BMI or with the clinically meaningful neurologic measure Expanded Disability Status Scale. There was also no correlation with the self-report questionnaires Multiple Sclerosis Quality of Life-54 (MS-54), Fatigue Severity Scale, Modified Fatigue Impact Scale, and Beck Depression Inventory.

NfL Alterations as Consequence of Dietary Intervention

Intragroup Comparison: AKD Induced Significant Reductions in sNfL Within 6 months

In the AKD group ($n = 17$), we found a significant decrease in the sNfL level from 8.5 at baseline (no dietary intervention) to 7.1 pg/mL (mean difference -1.307; 95% CI -2.256 to -0.357; $p < 0.05$) at the end of the study after 6 months. The analysis of the CR group ($n = 14$) and CD group ($n = 9$) showed similar sNfL levels at baseline and at the final visit (Figure 1 and eTable 1, links.lww.com/NXI/A653).

Figure 1 Intra-Group Comparison – Adapted Ketogenic Diet Lowers the Serum Concentration of Neurofilament Light Chain II



Alterations of serum neurofilament light chain (sNfL) levels induced by caloric restriction and adapted ketogenic diet. Data represent mean \pm standard error of the mean and were measured at baseline and retested at 6 months for all groups. *p* values indicate Wilcoxon matched-pairs signed rank test analysis, **p* < 0.05.

Intergroup Comparison: AKD Lowers sNfL Levels in Comparison to Common Diet

We used baseline sNfL concentrations, relapse rate, and Δ -BMI as covariates for between-group evaluation. At the end of the study, we observed a significant decline in sNfL levels (Figure 2 and eTable 2, links.lww.com/NXI/A653) when we compared AKD with CD (adjusted difference -2.145 ± 0.615 ; 95% CI -3.393 to -0.897 ; *p* = 0.001) after 6 months. We did not find a difference in sNfL levels between CR and CD groups at month 3 (adjusted difference 1.24 ± 1.71 ; 95% CI -2.236 to 4.716 ; *p* = 0.682).

Relapse Rate and sNfL Over the Course of the Study

Twelve months prior study outset, the relapse rate did not differ significantly between the groups (Table 1). In the course of the study, 3 relapses in the CD group, 2 relapses in the CR group, and 1 relapse in the AKD group were reported. When analyzing sNfL levels at month 3, we detected a significant increase over all groups (eFigure 6 and eTable 1, links.lww.com/NXI/A653). After excluding

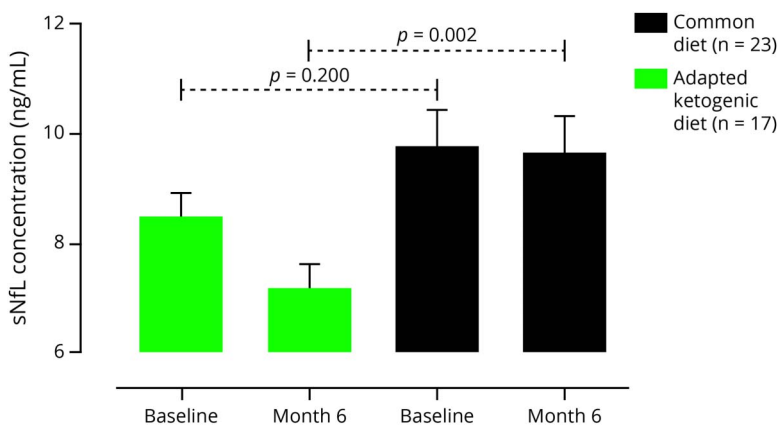
samples from patients with relapse activity, the sNfL levels of the AKD group remained elevated (eFigure 7, links.lww.com/NXI/A653).

Discussion

In this randomized controlled trial we investigated the impact of intermittent fasting/CR and AKD on sNfL levels in patients with MS. In the examined groups, sNfL levels were comparable with previous studies using the same methodology supporting the reliability of our results and of the applied assay analysis method.³ However, earlier studies have described higher sNfL levels due to various internal protocols, as recently discussed.¹⁰

This setting not only allowed assessment of the relative effects of AKD treatment vs common diet and 7-day fasting in a parallel randomized design but also provided the opportunity to study the relation of sNfL as a specific marker of neuroaxonal damage with other anthropometric, clinical, and

Figure 2 Comparison of sNfL Levels in Patients With MS on Adapted Ketogenic Diet With Common Diet



After 6 months of adapted ketogenic diet, the serum neurofilament light chain (sNfL) concentration declined significantly compared to the common diet group. Data were measured at baseline and retested at 6 months. Data represent mean \pm standard error of the mean. ***p* < 0.01, analysis of covariance (ANCOVA) adjusted for relapse rate, BMI and baseline dependencies.

patient-reported measures of disease activity and severity obtained under good clinical practice conditions.

Baseline characteristics of patients included in this analysis do not suggest any relevant selection bias. sNfL concentrations in patients were similar over all groups at baseline. sNfL levels at baseline were correlated with functional markers of disease activity (e.g., PASAT and MSFC) confirming recent reports.³ Although diet as an environmental factor has been acknowledged in MS, its precise role in the pathogenesis of MS is still far from clear. Moreover, dietary interventions are rapidly gaining popularity within the MS community, but little is known about their safety. AKD and CR have been suggested to induce autophagy and to improve gut microbiota, redox status, and regulatory T cells in a variety of diseases.^{5,7,9,11,12}

It has previously been shown that NfL levels, as a neuroaxonal integrity parameter and marker of treatment response, decrease after initiation of DMTs.³ In line with this, our results indicate that AKD may influence sNfL levels in patients with MS. More studies are necessary to evaluate the longitudinal impact of repeated fasting cycles on sNfL levels in MS.

Our analysis reflects a significant relationship between AKD and declining sNfL levels at 6 months. This suggests that AKD may benefit the course of disease in MS. However, more comprehensive studies with an expanded time span are needed to examine whether effects could relate to the diet itself rather than its impact on MS. The limitations of our study include its short observation time of 6 months, MRI was not performed and a limited power, due to a relatively small number of participants (n = 40 analyzed patients). It should also be noted that the extent of variation indicating a clinically meaningful change in longitudinal sNfL measurements has not yet been defined. However, we also found increased sNfL levels in the AKD group, independent of relapse activity at month 3. Our observation could be due to autophagy of which AKD and CR are strong inductors.^{11,12} Of interest, it was reported that autophagy promotes intracellular degradation of cytoskeleton but also seems to be essential for the survival of neuronal cells.¹³ However, it is likely that release of degradation products would increase sNfL levels extracellularly (and thus in serum) at least in the short term—thus our results would be consistent with the latter results of different groups.

AKD needs more time than CR to activate homeostatic processes, e.g., autophagy, due to the process of keto-adaptation and slower/lower ketone body excess. This may explain why we did not detect an increase in sNfL in the CR group at month 3.

Overall, our study suggests that an AKD offers an avenue to impact sNfL levels, which seems to be a promising biomarker in neuroinflammatory diseases, supporting the use of dietary

interventions as a treatment tool for MS. These findings are of urgent medical interest because such dietetic strategies exhibit few unwanted side effects.

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Disclosure

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Falk Steffen, MD	Department of Neurology, Focus Program Translational Neuroscience (FTN) and Immunotherapy (FZI), Rhine Main Neuroscience Network (rmn2), University Medical Center of the Johannes Gutenberg University Mainz, Germany	Drafting/revision of the manuscript for content, including medical writing for content; study concept or design; and analysis or interpretation of data
Frauke Zipp, MD	Department of Neurology, Focus Program Translational Neuroscience (FTN) and Immunotherapy (FZI), Rhine Main Neuroscience Network (rmn2), University Medical Center of the Johannes Gutenberg University Mainz, Germany	Drafting/revision of the manuscript for content, including medical writing for content; study concept or design; and analysis or interpretation of data

Continued

Appendix (continued)

Name	Location	Contribution
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