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Prevalence, correlates and impact of pain and cramps in anti-MAG neuropathy: a multicentre European study

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Key Words: cramps, anti-MAG, function, neuropathy, pain, quality of life

Abbreviations: anti-MAG: anti myelin-associated glycoprotein; FSS: Fatigue Severity Score; INCATSSS: Inflammatory Neuropathy Cause and Treatment Sensory Sum Score;

I-RODS : Inflammatory Rasch-Built Overall Disability Scale; MOSS: Medical Outcome Study-Social Support Survey; NPSI: Neuropathic Pain Symptom Inventory; ONLS: Overall Neuropathy Limitation Score; SF-36 HR-QoL: SF-36 health-related quality of life;

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Abstract.

Background: The frequency of pain and cramps is uncertain in anti-MAG (Anti-myelin associated glycoprotein antibody) neuropathy. Whether these symptoms may affect function/quality of life is unknown.

Methods: We performed a cross-sectional study of the prevalence, correlates and impact of pain, pain subtypes and cramps, their severity, frequency and anatomical distribution, in 55 clinically stable patients with anti-MAG neuropathy.

Results: Pain of any type was reported by 80% of subjects. The most common subtype was paraesthesiae and dysaesthesiae (70%). Cramps were reported by >60% of patients, with lower limb cramps in all and upper limb cramps, in about 20%. Cramps affected daily activities in >30% of these

subjects, sleep in 60%, ability to exercise in >30%. Total Pain Score correlated with several SF-36 health-related quality of life (SF-36 HR-QoL) measures ($p < 0.05$), with Inflammatory Rasch-Built Overall Disability Scale (I-RODS) ($p = 0.006$) and 10-metre timed walk ($p = 0.019$). An independent association was ascertained with I-RODS ($p = 0.002$). Different pain subtypes showed multiple associations with SF-36 HRQoL measures and/or functional scales. Upper limb cramps had multiple SF-36 HRQoL/functional correlates, with an independent association with the Overall Neuropathy Limitation Score (ONLS) ($p = 0.004$). Cramp severity correlated with ONLS ($p = 0.04$) and I-RODS ($p = 0.028$) and inversely with level of physiotherapy input ($p = 0.009$). Cramp frequency was associated with tremor score ($p = 0.004$) and multiple SF-36 HRQoL subsections.

Conclusions: Neuropathic pain and cramps may affect function and quality of life in anti-MAG neuropathy. Optimizing treatments of these symptoms, including by adequate levels of physiotherapy, may be beneficial in affected patients and requires further research.

Introduction.

Anti-myelin associated glycoprotein antibody (Anti-MAG) neuropathy is a paraproteinaemic demyelinating neuropathy of usually slow progression resulting in disability through distal sensory more than motor deficits, as well as tremor. Neuropathic pain and cramps can be prominent and disabling [1-4]. We have previously described our findings of a study on the functional determinants of quality of life in this multicentre cohort [5]. This current analysis focused on the issue of pain and cramps in patients with anti-MAG neuropathy.

There are currently no effective, evidence-based immunotherapy treatments for anti-MAG neuropathy [6]. Immunoglobulins may have some short-term effect [6] but their appropriateness in this setting is highly questionable. In addition, following unconvincing initial results of a randomized controlled trial of Rituximab which had suggested a possible, albeit moderate, beneficial effect [7], a subsequent

larger trial showed globally negative results [8]. Globally, a recent meta-analysis provides however globally, low-quality evidence for Rituximab and further longer-duration trials are needed [6].

Consideration of potential determinants of disability and quality of life is important in this medically currently untreatable disorder as may help shed light on potentially useful symptomatic therapies and ways of improving patients' daily lives.

The main objective of the current analysis was to evaluate the impact pain and its severity and quality, as well as of cramps, their severity, frequency and anatomical location, on neurological function and quality of life in patients with anti-MAG neuropathy.

Patients and Methods.

We performed a cross-sectional study of patients with a clinically, electrophysiologically and immunologically confirmed diagnosis of anti-MAG neuropathy, attending our respective neuromuscular clinics for their routine follow-up assessments, between March 2015 and February 2016. Inclusion criteria were (i) mainly distal sensory-predominant neuropathy (ii) presence of an IgM monoclonal protein (iii) presence of anti-MAG antibodies >1500 BTU [9] (iv) distal predominant demyelinating physiology (v) clinically stable for 3 months at time of assessment. Patients were evaluated irrespective of previous immunomodulatory therapies received and of current or previous treatments for pain and cramps.

Pain was evaluated by a validated Pain Questionnaire, the NPSI (Neuropathic Pain Symptom Inventory) [10]. The questionnaire is detailed in its English language version, in Table 1 (Online Supplementary Material). The Total Pain Score was the sum of the intensity scores for the different pain subtypes. We also established, for each subject, [10], pain sub-scores.

Cramps were evaluated by a validated questionnaire [11]. This questionnaire is detailed in its English language version in Table 2. (Online Supplementary Material) and ascertained severity, frequency and precise anatomical location of cramps.

All participants received a clinical assessment as part of their routine evaluation with: 80-point Medical Research Council (MRC) sum scores, Inflammatory Neuropathy Cause and Treatment (INCAT) Sensory Sum Score (INCATSSS) [12], Jamar grip dynamometry for dominant and non-dominant hands, Overall Neuropathy Limitation Score (ONLS) [13], Inflammatory Rasch-built Overall Disability Score (I-RODS) [14], 10-metre timed walk, ataxia score [8], Whiget tremor score [15]. Health-related Quality of Life (HR-QoL) was ascertained by an interviewer-administered SF-36 Questionnaire [16] and fatigue by the Fatigue Severity Scale (FSS) [17]. Social support was determined by a modified abbreviated version of the Medical Outcome Study-Social Support Survey (MOSS) [18]. Level of physiotherapy input determined by a questionnaire, based on current recommendations for access to therapy in the U.K. [19]. This is detailed in Table 3 (Online Supplementary Material).

No further evaluations were performed. Patients did not undergo any additional electrophysiological tests, further blood tests, imaging or cerebrospinal fluid examinations for the current analysis.

Biological and electrophysiological data were obtained from available records.

The SF-36 HRQoL Questionnaire was individually categorized before data analysis, as previously established, in different domains, consisting of “Physical Function” (PF), “Role Physical (RP), “Bodily Pain” (BP), “General Health” (GH), “Vitality” (VT), “Social Functioning” (SF), “Role Emotional” (RE), “Mental Health”, “Physical Component Score” (PCS) and “Mental Component Score” (MCS). Inter-correlations were ascertained and independent associations were sought.

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This work was approved and registered by our relevant respective institutional review boards. All patients provided informed written consent. Statistical analyses were performed using SPSS 23.0 software (SPSS Inc., Chicago, IL, USA). Correlations were determined by using Spearman's 2-tailed correlation. As this study was exploratory, correction for multiple testing was not performed. Multivariate logistic regression was used to determine eventual independent associations. P values <0.05 were considered as significant.

Results.

We recruited 55 patients (32 from Marseille, France, 7 from Angers, France, and 16 from Birmingham, U.K.). There were 38 males and 17 females. Mean age was 71.5 years (S.D.: 10.3). Mean disease duration was 7.4 years (S.D.: 5.9). Mean anti-MAG antibody titer was 43813 BTU (S.D.: 26827.5).

There were no significant differences between the French and U.K. cohorts in terms of general prevalence and/or severity of pain ($p=0.55$), and cramps ($p=0.54$), gender distribution ($p=0.14$), age ($p=0.62$) or disease duration ($p=0.23$). All physical assessments, including MRC sum score ($p=0.87$), Jamar grip dynamometry ($p=0.54$), ONLS ($p=0.11$), I-RODS ($p=0.66$), INCATSSS ($p=0.60$), 10-meter timed walk ($p=0.46$), ataxia score ($p=0.65$), tremor score ($p=0.25$), Fatigue Severity Score ($p=0.86$), were comparable in the French and U.K. cohorts.

Prevalence of evaluated symptoms are described in Table A. Most patients had a chronic sensory neuropathy with sensory ataxia (34/55, 62%) and upper limb tremor (27/55, 49%). Pain of any type was described by 44/55 patients (80%). Burning superficial spontaneous pain was reported by 21/55 (38.2%), pressing deep spontaneous pain by 19/55 (34.5%), paroxysmal pain by 27/55 (49.1%), evoked pain by 28/55 (50.1%) and paraesthesiae/dysaesthesiae by 39/55 (70.1%). Cramps were

reported by 35/55 (63.6%). Upper limb cramps were present in only 10 of these 35 patients with cramps (28.6%) whereas lower limb cramps were described by all 35. Cramps affected daily activities in 11/35 (31.4%) of these subjects, sleep in 21/35 (60%), ability to exercise in 11/35 (31.4%) and the patient perception of quality of life in 12/35 (34.3%).

Main results of correlation analyses showed multiple positive associations of pain severity and quality of life and disability. These are detailed in Table B.

Total Pain Score correlated with several SF-36 HR-QoL measures ($p < 0.05$) (see Table B.), with I-RODS ($p = 0.006$) and 10-metre timed walk ($p = 0.019$). The Total Pain Score was independently associated with BP ($p < 0.001$). Considering only the functional parameters, an independent association was found with the I-RODS ($p = 0.002$).

Burning superficial spontaneous pain was independently associated with the INCATSSS ($p = 0.004$) and modified MOSS ($p = 0.008$).

Paroxysmal pain was independently associated with PCS ($p = 0.007$). Amongst the functional parameters, an independent association was found with ataxia score ($p = 0.007$).

Paraesthesiae/dysaesthesiae were independently associated with BP ($p = 0.004$). Considering non-HR-QoL parameters, an independent association was found with mean FSS ($p = 0.029$).

Main correlation results of the cramp characteristics are summarized in Table C. Again, multiple positive associations were ascertained between cramp location, severity and frequency and quality of life and function. Lower limb cramps correlated with ataxia score only, amongst all studied functional parameters ($p = 0.034$).

Upper limb cramps were independently associated with PF ($p=0.011$). An independent association was found with the ONLS ($p=0.004$). More precise cramp location was relevant to performance and function as hand cramps were independently associated with ONLS ($p=0.015$). Cramp frequency was independently associated with RE ($p=0.031$) and BP ($p=0.031$).

Patients describing interference of cramps with their daily activities were significantly more likely to have a lower MRCSS ($p=0.045$), worse arm ONLS score ($p=0.003$) and total ONLS score ($p=0.009$), lower I-RODS ($p=0.035$), higher INCATSSS ($p=0.038$) and higher tremor score ($p=0.027$). Patients who felt cramps interfered with their ability to exercise were more likely to have lower arm ONLS scores ($p=0.016$). Those who described a perceived impact of their cramps on their global quality of life were similarly more likely to have lower ONLS arm scores ($p=0.018$), but also more severe ataxia ($p=0.031$), although SF-36 parameters failed to be influenced.

The level of physiotherapy input correlated with the Total Pain Score ($p=0.024$). Also, level of physiotherapy input was associated the presence of cramps ($p=0.04$). However, conversely and remarkably, cramp severity was strongly inversely associated with the level of physiotherapy input ($p=0.009$). Similarly, an independent negative association was found between level of physiotherapy input and cramps interfering with sleep ($p=0.011$).

The level of social support (modified MOSS) was associated with the presence of burning pain ($p=0.024$), but not with any other pain or cramp characteristics or features.

Discussion.

Our multicentre analysis of 55 patients firstly indicates that both pain and cramps are common features that may significantly impact upon HR-QoL, as well on function in patients with anti-MAG neuropathy. Importantly, both may be more frequent with longer disease duration, indicating the importance and relevance of regular re-assessments.

The frequency of pain was high, occurring in 80% of our cohort. Paraesthesiae and dysaesthesiae, although often considered as non-painful symptoms by patients and physicians, were present in >70%. Consideration of pain management, appears therefore highly relevant in anti-MAG neuropathy, as has been found previously in relation to quality of life more generally in neuropathy [20]. Similarly, we found comparable findings about cramps in our cohort. Although cramps were reported by 63.6% of patients, they affected mostly the lower limbs, with <20% describing upper limb involvement. Hand cramps appeared however the most relevant to disability with an independent association with the ONLS. Interestingly, whereas cramp severity was associated with neurological disability, cramp frequency on the other hand was related to HR-QoL measures, with independent associations with RE and BP. This is in keeping with previous findings in genetic neuropathy [11].

We found that presence of cramps was associated with higher levels of physiotherapy input. This may be intuitively expected as probably relates to the naturally higher likelihood of referrals of more affected subjects. However, the concomitant finding of worst cramps in patients with the lowest physiotherapy input may suggest a favorable effect of physiotherapy on cramp severity. We in addition, found that cramps interfering with sleep also inversely correlated with level of physiotherapy. This is in keeping with some recent literature [21], although the benefit of non-medical interventions for cramps is, to our knowledge, not established from meta-analyses performed to date [22].

HR-QoL measures represent the markers that ultimately need to be positively influenced by treatment interventions for meaningful benefit to patients [23]. Pain can be a prominent although under-recognized feature of inflammatory neuropathy [24, 25], and anti-MAG neuropathy may be a likelier cause of pain amongst immune-mediated neuropathies [24]. In that study, positive sensory symptoms of paraesthesiae and dysesthesiae were commonest. Our current results appear confirmatory of these findings and in agreement with the conclusions of that previous study that these are symptoms often not considered as pain in the clinical setting.

We did not evaluate the effects of previous immunotherapy or of medical treatments for pain and cramps in our cohort. This was because our aim was to cross-sectionally ascertain occurrence and correlates of these symptoms rather than retrospectively attempt to determine drug effects. We otherwise did not here specifically consider co-morbidities which may have resulted in arthritic or musculo-skeletal pains and this represents a limitation of our study.

Only properly designed therapeutic drug trials may in future answer the question regarding the eventual efficacy of pain and cramp treatment on function and quality of life in anti-MAG neuropathy. The benefits of physiotherapy input optimization are possible, this despite the cross-sectional and non-controlled nature of our analysis. The scale we used to determine level of physiotherapy input is clearly limited as does not consider intensity of sessions or differences of practice between physiotherapists and institutions, which limits the value of the findings. However, benefit of physiotherapy in anti-MAG neuropathy may merit further study in the setting of an adequate randomized controlled interventional trial. As demonstrated in other neuromuscular disorders, favorable effects of non-drug interventions such as aerobic exercise training and cognitive behavioral therapy may be obtained and objectively measurable [26].

In conclusion, we believe our study brings new insight on practical management of patients with anti-MAG neuropathy, in indicating that pain and cramp severity, frequency, quality and distribution can play a significant part in the impairment of function and quality of life in affected patients. This is important in a disorder for which effective evidence-based treatments remain unavailable. Future research may need to focus on these associated secondary, to date still poorly-recognized symptoms of anti-MAG neuropathy, both in terms of therapeutic benefit of evaluated drugs and other interventions, as well as, eventually, consideration of these manifestations *per se*, as primary therapeutic targets.

Table A. Prevalence of evaluated symptoms in 55 patients with anti-MAG neuropathy.

Symptom	Prevalence in cohort	Percentage
Sensory Ataxia	34/55	62%
Upper Limb Tremor	27/55	49.1%
Pain of any type	44/55	80%
Burning superficial spontaneous pain	19/55	34.5%
Paroxysmal pain	27/55	49.1%
Evoked pain	28/55	50.1%
Paraesthesiae, Dysaesthesiae	39/55	70.1%
Cramps	35/55	63.6%
Upper Limb cramps	10/55	18.2%
Lower Limb cramps	35/55	63.6%
Cramps affecting daily activities	11/55	20%
Cramps affecting sleep	21/55	38.2%
Cramps affecting ability to exercise	11/55	20%
Cramps affecting patient perception of quality of life	12/55	21.8%

Table B. Correlates of Pain (Total Pain

Score and Pain Quality) with functional and Health-Related Quality of Life Parameters in 55 patients with anti-MAG neuropathy.

	NPSI Total Pain Score	Burning Pain	Pressing Pain	Paroxysmal Pain	Paraesthesiae/ Dyaesthesiae
Disease duration	NS	Rho:0.438 P=0.047	NS	NS	NS
MRC sum score	NS	NS	NS	NS	NS
Dominant Grip	NS	NS	NS	NS	NS
ONLS Total	NS	NS	NS	Rho:0.527; P=0.005	NS
I-RODS	Rho=-0.409; P=0.006	NS	NS	Rho:-0.435; P=0.023	Rho:-0.313; P=0.049
INCAT sensory sum score	NS	Rho: 0.469 P=0.037	NS	Rho:0.447; P=0.025	NS
Ataxia score	NS	NS	NS	Rho:0.461; P=0.015	NS
Whiget Tremor score	NS	NS	NS	NS	NS
Ten meter walk	Rho:0.425; P=0.019	NS	NS	Rho:0.755; P<0.001	NS
Modified MOSS	NS	Rho: 0.491; P=0.024	NS	NS	NS
Fatigue Severity Scale (mean score)	Rho: 0.360; P=0.018	NS	Rho:0.467; P=0.044	Rho:0.444; P=0.020	Rho:0.376; P=0.017
PF	NS	NS	NS	Rho:-0.417; P=0.03	NS
RP	NS	NS	NS	NS	NS
BP	Rho:-0.661; P<0.001	NS	Rho:-0.678; P=0.001	NS	Rho:-0.454; P=0.003
GH	Rho=-0.366 P=0.014	NS	NS	Rho:-0.386; P=0.047	Rho:-405; P=0.010
VT	Rho=-0.307; P=0.043	NS	NS	NS	NS
SF	NS	NS	NS	NS	NS
RE	NS	NS	NS	NS	NS
MH	Rho=-0.316 P=0.037	NS	NS	NS	Rho:-0.391; P=0.013
PCS	Rho=-0.432; P=0.003	NS	NS	Rho:-0.520; P=0.005	NS
MCS	NS	NS	NS	NS	NS

Table C. Correlates of various cramp characteristics with functional and Health-Related Quality of Life Parameters in 55 patients with anti-MAG neuropathy.

	Presence of Cramps	Cramp Frequency (number of days or nights per week)	Severity Of Worst Cramp	Hand Cramps	Cramps Interfering With Daily Activities	Cramps Interfering with Sleep	Cramps Interfering with Quality of Life
Disease duration	Rho:0.289 P=0.034	NS	NS	Rho: 0.367 P=0.030	NS	NS	NS
MRC sum score	NS	NS	NS	Rho=-0.439 P=0.008	Rho=-0.41 P=0.045	NS	NS
Dominant Grip	NS	NS	NS	Rho= -0.447; P=0.015	NS	NS	NS
ONLS Total	NS	NS	Rho=-0.360 P=0.004	Rho:0.46 P=0.005	Rho:-0.433 P=0.009	NS	NS
I-RODS	NS	NS	Rho=0.383 P=0.028	Rho: 0.371 P=0.028	Rho:-0.357 P=0.035	NS	NS
INCAT sensory sum score	NS	NS	NS	NS	Rho:0.369 P=0.038	NS	NS
Ataxia score	Rho: 0.305 P=0.028	NS	NS	Rho: 0.398 P=0.022	NS	NS	Rho:0.376 P=0.031
Whiget Tremor score	NS	Rho:0.418; P=0.024	NS	NS	Rho:0.386 P=0.027	NS	NS
Ten meter walk	NS	NS	NS	NS	NS	NS	NS
Modified MOSS	NS	NS	NS	NS	NS	NS	NS
Level of Physiotherapy Input	Rho:0.28 P=0.040	NS	Rho=-0.449 P=0.009	NS	NS	Rho: -0.423	NS

						P=0.011	
Fatigue Severity Scale (mean score)	NS	Rho:0.472 P=0.007	NS	NS	NS	NS	NS
PF	NS	NS	NS	NS	Rh:-0.457 P=0.006	NS	NS
RP	NS	Rho:-0.55 P=0.001	NS	NS	Rh:-0.4 P=0.017	NS	NS
BP	NS	Rho:-0.405 P=0.024	NS	NS	NS	NS	NS
GH	NS	NS	NS	NS	Rh:-0.353 P=0.038	NS	NS
VT	NS	Rho:-0.453 P=0.011	NS	NS	Rho:-0.372 P=0.038	NS	NS
SF	NS	Rho:-0.614 P<0.001	NS	NS	Rho:-0.486 P=0.003	NS	Rho:-0.374 P=0.027
RE	NS	Rho:-0.590 P=0.001	NS	NS	NS	NS	NS
MH	NS	Rho:-0.511 P=0.003	NS	NS	NS	NS	NS
PCS	NS	NS	NS	NS	Rho:-0.566 P<0.001	NS	NS
MCS	NS	Rho:-0.564 P=0.001	NS	NS	NS	NS	NS

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