When is Onuf's nucleus involved in multiple system atrophy? A sphincter electromyography study

T Yamamoto, R Sakakibara, T Uchiyama, Z Liu, T Ito, Y Awa, K Yamamoto, M Kinou, T Yamanishi, T Hattori


Background: External anal sphincter (EAS) electromyography (EMG) abnormalities can distinguish multiple system atrophy (MSA) from Parkinson's disease in the first five years after disease onset. However, the prevalence of the abnormalities in the early stages of MSA is unknown.

Objectives: To present EAS-EMG data in the various stages of MSA.

Methods: 84 patients with 'probable' MSA were recruited (42 men, 42 women; mean age 62 years (range 47 to 78); mean disease duration 3.2 years (0.5 to 8.0; <1 year in 25%); 50 cerebellar form (MSA-C), 34 parkinsonian form (MSA-P)). EAS motor unit potential (MUP) analysis and EMG cystometry were carried out in all patients.

Results: The overall prevalence of neurogenic change of the EAS MUP was 62%—52% in the first year after disease onset, increasing to 83% by the fifth year (p<0.05); it also increased with severity of gait disturbance (p<0.05), storage and voiding disorders, and detrusor sphincter dysphagynergy (NS). The neurogenic change was not correlated with sex, age, MSA-P/C, postural hypotension, constipation, erectile dysfunction in men, underactive or acontractile detrusor, or detrusor overactivity. In 17 incontinent patients without detrusor overactivity or low compliance, urinary incontinence was more severe in those with neurogenic change than in those without (p<0.05).

Conclusions: Involvement of Onuf's nucleus in MSA is time dependent. Before the fifth year of illness, the prevalence of neurogenic change does not seem to be high, so a negative result cannot exclude the diagnosis of MSA.

METHODS

We recruited 84 patients with 'probable' MSA*: 42 men, 42 women; mean age 62 years (range 47 to 78); mean disease duration 3.2 years (0.5 to 8.0; <1 year in 21 patients (25%)); 50 MSA-C (cerebellar form), 34 MSA-P (parkinsonian form). We added an imaging study to ensure the diagnosis of the early cases; all patients had magnetic resonance imaging (MRI) abnormalities including pontocerebellar atrophy, abnormal signal intensity in the cerebellum, cerebellar atrophy, abnormal signal intensity in the posterior putamen, all of which were consistent with MSA and helped to exclude Parkinson's disease and progressive supranuclear palsy. Genetic analyses were carried out as far as possible to exclude hereditary spinocerebellar ataxia. No patient had abnormalities of blood chemistry (including blood sugar) or urinalysis. None had abnormal findings on digital examination or ultrasound echography of the pelvic organs.

We carried out standard EMG cystometry with a pressure-flow analysis in all patients, using an EMG computer (Neurpack Sigma; Nihon Kohden, Tokyo, Japan). We sampled at least 10 single MUPs per patient, manually examining the automatically sampled waves to ensure that the MUPs were indeed single. Neurogenic change was diagnosed when at least one of the following abnormalities was seen:

- more than 20% of MUPs had a duration of >10 ms;
- the average duration of MUP was >10 ms, including the late components in particular.14 18

We carried out standard EMG cystometry with a pressure-flow analysis in all patients. The filling phase abnormalities include detrusor overactivity and low compliance detrusor. The voiding phase abnormalities include detrusor-sphincter dysphagynergy. In the Schafer's nomogram test, we obtained detrusor contractility classed as strong, normal, weak, or very weak; the latter two were designated as underactive detrusor in this study. The detrusor is classed as acontractile when patients with large post-void residuals (PVR) cannot contract the detrusor at all without urinary flow.

Abbreviations: EAS, external anal sphincter; MSA, multiple system atrophy; MSA-C, cerebellar form of multiple system atrophy; MSA-P, parkinsonian form of multiple system atrophy; MUP, motor unit potential; PVR, post-void residual
PVR volume measured by transurethral catheterisation was
enced in 25, monthly in 14, weekly in 13, and daily in 32. The
Urinary incontinence (storage disorder) was never experi-
with aid (score 6–7) in 28, and wheelchair bound (score 8) in
1–3) in 23, walking with one stick (score 4–5) in 22, walking
was absent (score of 0) in none, independent walking (score
postural hypotension, constipation, erectile dysfunction in
men, underactive or acontractile detrusor, or detrusor over-
activity.

Seventeen of the 56 incontinent patients (seven men, 10
women) lacked abnormal bladder contraction during the
filling phase, although 12 of the 17 also had PVR (mean
135 ml (range 30 to 500)). In the 17 patients, urinary
incontinence was more severe in those with neurogenic
change (n = 8; monthly, 0; weekly, 1; daily, 7) than in those
without (n = 9; monthly, 4; weekly, 2; daily, 3) (p < 0.05).

RESULTS

Patients’ functional ability and urinary disorders were as
follows. Gait disorder (as measured by the international
cooperative ataxia rating scale, walking capacities subscale21) was absent (score of 0) in none, independent walking (score 1–3) in 23, walking with one stick (score 4–5) in 22, walking with aid (score 6–7) in 28, and wheelchair bound (score 8) in 11. All patients except for two had urinary symptoms. Urinary incontinence (storage disorder) was never experi-
enced in 25, monthly in 14, weekly in 13, and daily in 32. The
PVR volume measured by transurethral catheterisation was
<100 ml in 35, 100–200 ml in 27, and >200 ml in 22.

The overall prevalence of neurogenic change was 62% in
our patients—52% in the first year after disease onset, and
83% by the fifth year. Thus the prevalence of neurogenic change increased during the course of the illness (p < 0.05) (fig 1). Changes in the percentage of MUPs with a duration of
more than 10 ms, including patients undergoing repeated
studies, are shown in fig 2. Many of the patients who
underwent repeated studies had normal to mild abnormality
at the initial assessment, and this became marked during the
course of the illness, although in two cases the EAS EMG
findings remained normal. The prevalence of neurogenic
change was 47% in patients who walked independently, but
82% in those who were wheelchair bound (p < 0.05) (table 1). Similar but non-significant changes were found for urinary incontinence (59% of patients without urinary incontinence
had neurogenic change v 63% with incontinence); post-void
residual (58% with PVR <200 ml v 73% in those with PVR
>200 ml); and detrusor sphincter dyssynergy (60% in
patients without detrusor sphincter dyssynergy v 73% in
those with dysynergy). The neurogenic sphincter EMG
results were not clearly correlated with sex, age, MSA-P/C,

DISCUSSION

Results of the EAS EMG in over 500 MSA patients have
already been reported, with an abnormality rate of more than
70% in many studies.7–15 Compared with those findings, the
overall prevalence rate of neurogenic change in the present
study was slightly lower (62%). This is presumably because
up to 25% of our patients had a disease duration of one year
or less, as early referral to our department has increased
recently, and patients are able to come to us without referral.
Thus the diagnosis of MSA in such early cases should be
made with extreme caution. In addition to the clinical
diagnostic criteria,14 we added an imaging study to ensure the
diagnosis in all patients, and we carried genetic analyses as
far as possible. Although the EAS MUP abnormalities allow
one to distinguish MSA from Parkinson’s disease in the first
five years after disease onset,14 the prevalence of the
abnormalities in the early stages of MSA (or, conversely,
the false negative rate) has not been established up to now.
We report here for the first time that in our patient cohort
the prevalence of neurogenic change was 52% in the first year
after disease onset, increasing to 83% by the fifth year
(p < 0.05). Among the patients who underwent repeated
studies, many were normal or had only mild abnormality
at the initial examination, but the abnormality became marked
during the course of their illness. Therefore, as expected, the
involvement of Onuf’s nucleus in MSA is time dependent.
In the early stages of illness, the prevalence of neurogenic
change in MSA does not seem to be high. In two patients
who underwent repeated studies, the EAS EMG findings
remained normal. We do not know whether there are some

Figure 1  Neurogenic sphincter EMG and duration of illness. The
prevalence of neurogenic sphincter EMG increased during a course
of illness. MUP, motor unit potential.

All patients gave their informed consent before participat-
ing in the study. Statistics were analysed using the χ² test.

Figure 2  Change in percentage of motor unit potentials (MUPs) of
duration >10 ms with duration of illness. Roman figures = the number of
the patients; italic figures = the number of the patients who underwent
repeated study.

In the early stages of the illness, the prevalence of neurogenic change does not seem to be high, so a negative result cannot exclude a diagnosis of MSA.

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Competing interests: none declared

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1 Onufrowicz B. Notes on the arrangement and function of the cell groups in the sacral region of the spinal cord. J Nerv Ment Dis 1899;26 498-504.


### Table 1 Neurogenic sphincter EMG and clinical variables other than duration of illness

<table>
<thead>
<tr>
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<th>Patients with neurogenic sphincter EMG</th>
<th>Patients with neurogenic sphincter EMG</th>
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<tr>
<td></td>
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<td>%</td>
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<tr>
<td>Males</td>
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<td>65</td>
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<td>Age &lt; 60 years</td>
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<td>56</td>
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<td>MSA-C</td>
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<td>58</td>
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<tr>
<td>Postural hypotension</td>
<td>11/23</td>
<td>48</td>
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<tr>
<td>Constipation</td>
<td>40/66</td>
<td>61</td>
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<tr>
<td>Erectile dysfunction</td>
<td>4/5</td>
<td>80</td>
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<tr>
<td>Incontinence</td>
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<td>59</td>
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<td>RU &lt; 200 ml</td>
<td>36/62</td>
<td>58</td>
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<td>Detrusor overactivity</td>
<td>14/26</td>
<td>55</td>
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<tr>
<td>UD/AD</td>
<td>29/52</td>
<td>56</td>
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<tr>
<td>DSD</td>
<td>44/73</td>
<td>60</td>
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</table>

*International cooperative ataxia rating scale, walking capacities subscale.
AD, acontractile detrusor; DSD, detrusor sphincter dyssynergy; MSA, multiple system atrophy; RU, residual urine volume; UD, underactive detrusor.
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