Chronic proximal spinal muscular atrophy of childhood and adolescence: sex influence

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SUMMARY Segregation analysis was performed on 354 cases of chronic proximal spinal muscular atrophy of childhood and adolescence (CPSMA) in the total series and in a number of subgroups formed according to the age at onset and sex.

The analysis provided evidence of sex influence in the series studied, particularly in a subgroup of the milder form of the disease with onset between the 37th month and 18th year of life. In the latter subgroup, females were affected much less frequently. This was particularly striking after age at onset of 8 years, and only exceptionally were females affected after the age of 13 years. These facts point to incomplete penetrance of the gene.

The problem of the predominance of males in spinal muscular atrophy (SMA) of childhood and adolescence has been infrequently discussed in publications. However, closer analysis of reported data shows that, although unnoticed, the sex dependent differences have been present in many series.1-3 Only a few authors have stressed the predominance of males or the milder course of the disease in females.4-6

No sex differences were present in the series composed mainly of acute cases of SMA7-10 or in the cases of predominantly early onset.11 However, some authors have noticed that the course of the infantile form of the disease is more severe in males.6-12-17 Furukawa et al6 were of the opinion that SMA, and in particular the Kugelberg-Welander form, was a sex influenced condition.

We have previously presented data showing the influence of sex in a chronic proximal form of spinal muscular atrophy of childhood and adolescence (CPSMA)*.17 18 The segregation ratio in our material18 appeared higher for males than for females and the distribution of age at onset of the disease was clearly different in each sex. In an attempt to explain this we postulated a tentative hypothesis of genetic modification.18 Our analysis has shown that sex differences were particularly pronounced in the benign cases of CPSMA.19

The aim of the present study was a further analysis, partly based on results obtained by one of us,20 of the sex influence in CPSMA.

Material and methods

The present study is based on 354 cases originating from 264 sibships registered by the Department of Neurology, Medical School of Warsaw, in the years 1960 to 1982. Grouping of the cases according to the classification of Hausmanowa-Petrusewicz et al21 is shown in table 1. As can be seen in the table,

<table>
<thead>
<tr>
<th>SMA form</th>
<th>M</th>
<th>F</th>
<th>M and F</th>
</tr>
</thead>
<tbody>
<tr>
<td>IA</td>
<td>25</td>
<td>21</td>
<td>46</td>
</tr>
<tr>
<td>IB</td>
<td>71</td>
<td>68</td>
<td>139</td>
</tr>
<tr>
<td>II</td>
<td>30</td>
<td>40</td>
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</tr>
<tr>
<td>III</td>
<td>63</td>
<td>36</td>
<td>99</td>
</tr>
<tr>
<td>Total</td>
<td>189</td>
<td>165</td>
<td>354</td>
</tr>
</tbody>
</table>

Form IA—acute, onset recorded at birth, never able to walk, survival up to 4 years.

Form IB—chronic, onset recorded at birth or in early months of life, never able to walk, survival up to 30 years.

Form II—age at onset first to fifth year, immobilised usually between tenth and fourteenth year, long survival.

Form III—age at onset from first year to adolescence, never complete immobilisation, long survival.

Relevant nomenclature:

Acute form of Werdnig-Hoffmann—corresponding to IA.

Intermediate form—corresponding to IB and II.

Mild form of Kugelberg-Welander—corresponding to III.

*Terms used for the same condition in other publications: (1) chronic form of childhood spinal muscular atrophy, (2) chronic forms of infantile and juvenile spinal muscular atrophy, (3) benign juvenile pseudodystrophic spinal muscular atrophy. This is a synonym of juvenile muscular atrophy or the Kugelberg-Welander form of the disease.

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The results are presented in Tables 2, 3, and 4. It is apparent that p-values are highest in the lowest age at onset subgroup (0 to 6 months). The p-values are also high in the subset of the low incidence group (0 days to 3 months). In the subset of the low incidence group, the p-values are highest in the lowest age at onset subgroup (0 to 6 months). The sudden fall of the p-value in the lowest age at onset subgroup (0 to 6 months) can be attributed to the assumption that the probability of ascertainment is not affected by the age at onset. If this assumption is false, the p-value can be inflated. The results of the analysis of segregation ratios in different age at onset subgroups are shown in Table 2. The results of the analysis of segregation ratios in different sex at onset subgroups are shown in Table 3. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 4. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 5. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 6. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 7. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 8.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Age at onset</th>
<th>Subgroups</th>
<th>M</th>
<th>F</th>
<th>M</th>
<th>F</th>
<th>M</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0-9 m</td>
<td>10 m-36 m</td>
<td>37 m-8 y</td>
<td>9 y-18 y</td>
<td>25 m-18 y</td>
<td>37 m-18 y</td>
<td></td>
<td></td>
</tr>
<tr>
<td>A</td>
<td>61</td>
<td>55</td>
<td>116</td>
<td>84</td>
<td>10</td>
<td>8</td>
<td>16</td>
<td>14</td>
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<tr>
<td>R</td>
<td>79</td>
<td>72</td>
<td>151</td>
<td>94</td>
<td>16</td>
<td>2</td>
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<td>24</td>
<td>16</td>
<td>40</td>
<td>32</td>
</tr>
<tr>
<td>p</td>
<td>0.025</td>
<td>0.020</td>
<td>0.022</td>
<td>0.036</td>
<td>0.039</td>
<td>0.022</td>
<td>0.016</td>
<td>0.019</td>
</tr>
<tr>
<td>δp</td>
<td>0.003</td>
<td>0.003</td>
<td>0.002</td>
<td>0.002</td>
<td>0.002</td>
<td>0.002</td>
<td>0.002</td>
<td>0.002</td>
</tr>
</tbody>
</table>

M-females, F-females. A-total number of probands (some of the figures are corrected based on the assumption that A is proportional to R). R-total number affected. T-total number of sibs. p-genetic ratio. 

*Difference from 0.25 more than 2 SD.

†Difference from 0.25 more than 3 SD.

**Segregation analysis was carried out on the age at onset of sex separately, but segregation ratios were calculated in six different age at onset subgroups (Table 2) and in five cumulative age at onset subgroups (Table 3). The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 4. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 5. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 6. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 7. The results of the analysis of segregation ratios in different age at onset subgroups and sex at onset subgroups are shown in Table 8.
the disease corresponds roughly to the age at onset 0 to 3 months in table 3. A large proportion of the cases in form II appear to fit well to the age range of 10 to 36 months and in form III to the age range 37 months to 18 years.

Discussion

Most of the estimates of genetic ratios in these series may be considered as characteristic of autosomal recessive transmission of the gene, and in fact we did not observe any pedigrees which would provide evidence of other forms of inheritance, with the exception of one pedigree which suggested X linked transmission. In such a large series, however, it could have occurred simply by chance.

The sex dependent differences in p values become obvious from the age at onset of 37 months upwards. They seem to start in the subgroup between 37 months and 8 years and between 9 and 18 years they are highly significant (p<0.001). They are also significant in some other subgroups, namely 25 months to 18 years (p<0.05), 37 months to 18 years (p<0.01), and in form III of the disease (p<0.01). In the material taken as a whole (age range 0 to 18 years) the differences are not significant owing to the overwhelming majority of the earlier onset cases. Nevertheless it appears to us that our data provide good evidence that CPSMA is a sex influenced condition, based on the lower number of females in the later onset subgroups and, therefore, decreased p values for females. These facts are in keeping with the previous studies of male-female sib pairs, which have shown that the disease has a milder course in females.15 17

In the age range 0 to 9 months the penetrance of the gene is complete and between 10 and 36 months there is a sudden lowering in p values (table 2). The reduction in female cases in the later age at onset ranges starting from 37 months, with clearly retained features of autosomal recessive inheritance, seems to be connected with incomplete penetrance (table 2). In the subgroup 37 months to 8 years, although the p value for females is smaller, the reduced penetrance is difficult to prove because the subgroup is not large enough (25 affected subjects) (table 2). However, in the subgroup 9 to 18 years, although again small (18 affected subjects: 16 males, two females), the reduced penetrance is remarkable, probably owing to a 'female sparing factor': p value for males 0·2 and for the overall subgroup below 0·1. The most convincing, because of the larger numbers (51 affected subjects), is the combined subgroup 37 months to 18 years with a p value for males close to the expected 0·25, for females close to 0·1, and for both sexes combined 0·17. This might indicate about 30% reduction in penetrance in this subgroup.

As Furukawa et al18 have suggested, sex influence may be associated with hormonal differences between males and females, the female gonadal hormones possibly playing the role of a 'female sparing factor'. Depending on the age at onset, this would be expressed:

(1) in earlier age, in a milder course of the disease in females, and in some dropping of the proportion of affected females between the third and eighth year of life;
(2) after the eighth year of life, in a significant decrease of the number of female cases; and

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The text continues with further discussion and analysis of the data presented in tables 3 and 4.
after the age of 13 years in an almost complete lack of female cases.

These facts might well be connected with some known milestones of female hormonal development, for instance the increase of FSH and oestrogen secretion around the eighth year of life as well as a sharp increase in some fractions of oestrogen just before the menarche, that is, around the thirteenth year of life.\textsuperscript{23}

The question of whether or not the established sex influence is related to the specific form of chronic proximal SMA of childhood and adolescence remains to be answered. From the study of Namba et al.,\textsuperscript{18} it seems that the more frequent occurrence of SMA in males than in females is also, and even to a greater degree, observed in adults. However, their data should be treated with caution since they were based on material composed of all genetic forms of SMA, including the sex linked form which is clearly different. According to other sources, no sex influence was detectable in the dominant\textsuperscript{24} and in the autosomal recessive proximal adult form of the disease.\textsuperscript{25} On the other hand there was a preponderance of males in the distal autosomal recessive form of SMA.\textsuperscript{28}

It is unlikely that the confirmation and explanation of the putative role played by hormones in the modulation of phenotypic expression in CPSMA could be achieved other than through experimental studies. Perhaps some known animal models of SMA\textsuperscript{27} \textsuperscript{28} could be of help.

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