Successful autologous stem cell transplantation in a patient with chronic inflammatory demyelinating polyneuropathy [letter]
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Mexiletine is effective on segmental hyperhidrosis: report of two cases

Segmental hyperhidrosis occurs in patients who have spinal cord lesions such as syringomyelia, vascular disorders, or tumours. Severe segmental hyperhidrosis not only causes discomfort, but may disturb daily life, as it did with our patients. There have been reports on the treatment of segmental hyperhidrosis. We report on two patients with segmental hyperhidrosis caused by syringomyelia and cavernous haemangioma of the spinal cord, in which oral administration of mexiletine ameliorated the symptoms.

A 41 year old woman experienced dysaesthesia of the left side of the trunk, which gradually progressed. At the age of 36 she developed dysaesthesia of the left upper limb as well. At that time, she had excessive sweating on the left side of her face, trunk, and upper limb, even when it was not hot, and had to change her nightwear a few times each night. Neurological examination detected dissociated sensory disturbance in the left half of the trunk in the Th2-Th4 area. Hyperhidrosis was present on the left side from the face to the trunk at the Th4 level in the spine. The room was maintained at 60±1% humidity and 36±1°C. Sweat rates were measured for both patients and five healthy men (mean age 53.6 years) and five women (mean age 42.3 years). Data for the patients were obtained before treatment (open circles) and after oral administration of 200 mg/day mexiletine (closed circles).

In addition, we tried mexiletine to eight patients (five women, age range 18–34 years) with primary hyperhidrosis, which is defined as excessive, uncontrollable sweating without any discernible cause. No improvement was noted after oral administration of 200 mg/day mexiletine.

This is the first report of the use of mexiletine to treat segmental hyperhidrosis. Three mechanisms have been hypothesised as possible causes of segmental hyperhidrosis brought about by spinal cord lesions: (1) disinhibition of preganglionic sympathetic neurons due to interruption of the inhibitory descending pathway, (2) overactivity of preganglionic sympathetic neurons due to spinal cord lesions, and (3) autonomic hyperreflexia due to cutaneous stimulation, we suspect persistent spontaneous overactivity of the preganglionic sympathetic neurons due to spinal cord lesions.

Oral administration of mexiletine was an excellent remedy for the segmental hyperhidrosis in our patients. Mexiletine is a sodium channel blocker reported to be effective for treating painful neuropathies.7 In terms of its mechanism, mexiletine is thought to inhibit the spontaneous activity of regenerating fibres in the spinal cord or to act on the spinal mediated nociceptive flexor reflex.8 We consider it compatible that in our patients mexiletine inhibited the spontaneous activity of the intermediolateral nucleus caused by spinal cord lesions. Our findings indicate that sodium channel blockers, such as mexiletine and carbamazepine, should be considered for the treatment of segmental hyperhidrosis caused by spinal cord lesions.

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References

Does the presence of a pontine trigeminal lesion represent an absolute contraindication for microvascular decompression in drug resistant trigeminal neuralgia?

Typical trigeminal neuralgia is characterised by recurring, paroxysmal, lancinating, shock-like pain within the distribution of one or

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more branches of the trigeminal nerve. Light tactile stimulation may trigger such an attack. Although the contribution of central and peripheral mechanisms to the aetiopathogenesis of trigeminal neuralgia still remains unclear, the concept of vascular compression of the trigeminal root as the main causal factor in idiopathic ‘tic douloureux’ has achieved widespread acceptance. Trigeminal neuralgia may also afflict patients with multiple sclerosis. In these cases, the demyelination of central trigeminal pathways is the accepted aetiology, and the presence of a T2 hyperintensity along the intrapontine course of trigeminal fibres is generally considered a contraindication to microvascular decompression. We recently saw a case of successful microvascular decompression in a patient without multiple sclerosis, despite an intrapontine trigeminal lesion.

This 66 year old previously healthy man presented with a 6 year history of intense, paroxysmal, electric shock-like pain in the territory of the second branch of the right trigeminal nerve. The pain was triggered by washing his face and shaving and it lasted for a few seconds. Painful attacks, initially rare, gradually increased in frequency and intensity and spread to the first trigeminal branch. When admitted to our hospital 1200 mg carbamazepine were ineffective in relieving the pain. Before admission phenotoin, baclofen, and lamotrigine had been tried without success. Neurological examination was negative and in particular there were no gross sensory deficits in his right trigeminal territory. Magnetic resonance imaging showed a T2 hyperintensity of intrapontine trigeminal fibres and nucleus (fig 1), without evidence of vascular conflicts with the trigeminal root entry zone. A controlateral, smaller, symmetric lesion was also evident. Multiple sclerosis and Lyme disease were ruled out by clinical history and appropriate investigations. The patient had no vascular risk factors. Holter monitoring, neck vessels colour echo Doppler, transthoracic echocardiographic examination, and brain MR angiography did not disclose alterations suggesting a possible ischaemic origin.

At operation, performed through a keyhole retrosigmoid craniectomy, the root entry zone of the nerve was found crossed by an “intratrigeminal” vein, which was electrocoagulated and divided. No other vascular contacts could be detected by careful exploration of the intracisternal tract of the nerve. The postoperative course was uneventful. Paroxysmal pain slowly faded away during subsequent weeks. Carbamazepine could be completely withdrawn after 2 months.

This case is intriguing for two reasons: firstly, because we obtained an apparently paradoxical therapeutic answer in a case where microvascular compression should have been generally contraindicated; secondly, because of the rarity of the T2 hyperintensity of intrapontine trigeminal fibres and nucleus in patients without multiple sclerosis.

Although the intimate aetiopathogenetic mechanisms of trigeminal neuralgia still remain unknown, peripheral lesions affecting the trigeminal nerve entry zone (tortuous vessels, meningiomas, aneurysms, artereovenous malfunctions, lipomas, epidermoid cysts, osteomas, etc) and multiple sclerosis are certainly involved in the mechanisms causing paroxysmal pain. Demyelination of trigeminal fibres at the level of trigeminal root entry zone in case of vascular cross compression and demyelination of intrapontine trigeminal fibres in case of multiple sclerosis (personal observation of MRI in more than 80% of cases) may result in ephaptic, abnormal transmission of impulses. A neurovascular conflict with focal demyelination at the root entry zone was surgically and pathologically confirmed in a patient with multiple sclerosis and trigeminal neuralgia by Lazar and Kirkpatrick. The anatomopathological evidence of demyelination of intrapontine trigeminal fibres in a patient with multiple sclerosis and trigeminal neuralgia was provided by Crooks and Miles. Recent studies hypothesised that vascular compression (and possible consequent demyelination) of the trigeminal root and demyelination of intrapontine trigeminal fibres due to multiple sclerosis can coexist and perhaps cooperate in the etiology of painful attacks. These findings led us to propose microvascular decompression to the patient rather than percutaneous lesive methods. At 2 year follow up the patient is still pain free, confirming the hypothesis that vascular compression and central demyelination can coexist and cooperate in provoking pain paroxysms. Hence, the classic distinction between the supposed “all central” and the “all peripheral” mechanisms for trigeminal neuralgia associated with multiple sclerosis and the “all peripheral” mechanism for the trigeminal neuralgia related to vascular compression should be overcome in favour of a unique (pathway with trigeminal neuralgia and multiple sclerosis are included), mixed central-peripheral mechanism in which abnormal impulses coming from demyelinated axons (multiple sclerosis, vascular compression, and any other possible cause of demyelination along the central and the peripheral course of gasserian ganglion fibres) modulate the nuclear activity.

An alternative view of our case might be the one advocating the hypothesis that pain relief might be due to surgical damage to the trigeminal root. This view has been strongly challenged by the results of recent studies on sensory effects of microvascular decompression. In our patient, too, no evidence of sensory deficits could be found postoperatively. Generally a minimum myelin damage, without any gross nerve hypofunction, is involved in the aetiopathogenesis of trigeminal neuralgia. In rare cases demyelination is so widespread along trigeminal fibres to be visualised by MRI and only patients with multiple sclerosis show the classic T2 hyperintensity along the intra-axial trigeminal pathways.

So the best of our knowledge this case is the third reported in which such a lesion was found in a patient without multiple sclerosis. In the two previous cases a pontine ischaemia was supposed: in one, multiple cerebral ischaemic lesions with widespread cortical atrophy were found; in the other the pontine lesion was isolated, but an MR angiogram showed a focal stenosis of the vertebral artery. A few patients with small pontine infarcts were reported with trigeminal symptoms, but without paroxysmal pain.

This case suggests that the presence of a pontine trigeminal lesion is not an absolute contraindication for microvascular decompression in cases of drug resistant typical trigeminal neuralgia.

References

Reversal of tissue hypoxia by a single intraventricular dose of sodium nitroprusside in a patient with severe medically refractory cerebral vasospasm after subarachnoid haemorrhage

A 29 year old man was referred to our department from a local hospital in November 2000, for treatment of acute subarachnoid haemorrhage. At examination, he presented with stupor and a Glasgow coma scale score of 7 and was intubated and artificially ventilated. Diagnostic angiography was performed the same day and demonstrated a ruptured anterior cerebral artery aneurysm. The complex configuration of the aneurysm precluded embolisation as a treatment option. Surgery was
Figure 1 Left frontal brain tissue oxygen pressure (p\text{tI}O_2), mean arterial pressure (MAP), and intracranial pressure (ICP) in a patient with cerebral vasospasm after subarachnoid haemorrhage. Low p\text{tI}O_2 values indicating critical brain hypoxia prompted emergency endovascular balloon dilatation of the vasospastic C1 segment of the left internal carotid artery, which only temporarily improved cerebral oxygenation. After intraventricular administration of a single dose of 40 mg sodium nitroprusside, cerebral oxygenation improved permanently.

References


Transcranial magnetic stimulation alleviates truncal ataxia in spinocerebellar degeneration

Spinocerebellar degeneration is an inherited or acquired neurodegenerative disorder characterised by slowly progressive cerebellar ataxia, dystarthis, and gait disturbance. These symptoms restrict daily activities. However, no satisfactory therapy has been established. Transcranial magnetic stimulation (TMS), originally introduced to the medical field to evaluate the function of the CNS, is recently becoming a therapeutic tool for neuropsychiatric disorders, such as major depression and Parkinson's disease. We also reported the efficacy of TMS for inherited spinocerebellar...
Table 1  Age, disease duration, and disease severity of patients enrolled in this study and the results of comparisons between active stimulation and sham stimulation, and between cerebellar-type SCD and OPCA type SCD receiving active stimulation

<table>
<thead>
<tr>
<th></th>
<th>Active</th>
<th>Sham</th>
<th>p Value†</th>
<th>Active cerebellar</th>
<th>Active OPCA</th>
<th>pValue‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>56.31 (1.96)</td>
<td>58.83 (1.47)</td>
<td>0.32</td>
<td>62.00 (2.03)</td>
<td>54.48 (2.80)</td>
<td>0.08</td>
</tr>
<tr>
<td>Disease duration (y)</td>
<td>7.73 (0.71)</td>
<td>6.67 (0.82)</td>
<td>0.33</td>
<td>8.28 (1.27)</td>
<td>5.52 (0.91)</td>
<td>0.10</td>
</tr>
<tr>
<td>Disease types (%) (cerebellar:OPCA)</td>
<td>17:22</td>
<td>16:19</td>
<td>0.85</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>10 m time (s):</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before TMS</td>
<td>14.37 (1.80)</td>
<td>13.78 (1.23)</td>
<td>0.79</td>
<td>13.74 (2.92)</td>
<td>16.24 (2.26)</td>
<td>0.23</td>
</tr>
<tr>
<td>After TMS</td>
<td>9.90 (0.73)</td>
<td>12.65 (1.20)*</td>
<td>&lt;0.05</td>
<td>8.11 (0.49)</td>
<td>12.16 (1.21)*</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>10 m steps</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before TMS</td>
<td>23.88 (2.19)</td>
<td>25.19 (1.71)</td>
<td>0.64</td>
<td>23.91 (2.22)</td>
<td>24.41 (3.62)</td>
<td>0.28</td>
</tr>
<tr>
<td>After TMS</td>
<td>19.47 (1.19)</td>
<td>23.49 (1.69)*</td>
<td>&lt;0.05</td>
<td>19.97 (1.89)</td>
<td>19.39 (1.55)</td>
<td>0.40</td>
</tr>
<tr>
<td>Tandem steps</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before TMS</td>
<td>1.24 (0.37)</td>
<td>2.26 (0.67)</td>
<td>0.19</td>
<td>2.39 (0.77)</td>
<td>1.05 (0.49)</td>
<td>0.20</td>
</tr>
<tr>
<td>After TMS</td>
<td>7.92 (2.12)</td>
<td>3.25 (1.38)</td>
<td>0.0005</td>
<td>10.72 (3.52)</td>
<td>4.81 (1.87)*</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Standing capacities:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before TMS</td>
<td>2.36 (0.18)</td>
<td>2.43 (0.21)</td>
<td>0.72</td>
<td>2.11 (0.34)</td>
<td>2.48 (0.18)</td>
<td>0.17</td>
</tr>
<tr>
<td>After TMS</td>
<td>1.64 (0.11)</td>
<td>2.11 (0.17)*</td>
<td>&lt;0.05</td>
<td>1.23 (0.16)</td>
<td>1.86 (0.13)</td>
<td>&lt;0.01</td>
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<tr>
<td>Walking capacities:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before TMS</td>
<td>2.21 (0.12)</td>
<td>2.11 (0.20)</td>
<td>0.64</td>
<td>2.00 (0.34)</td>
<td>2.19 (0.13)</td>
<td>0.41</td>
</tr>
<tr>
<td>After TMS</td>
<td>1.67 (0.13)</td>
<td>2.07 (0.20)</td>
<td>0.09</td>
<td>1.23 (0.16)</td>
<td>1.62 (0.18)</td>
<td>0.51</td>
</tr>
</tbody>
</table>

Values are shown as mean (SEM).

*p<0.05 compared with each value before TMS in sham stimulation group; †p Values shown in this table were analyzed between active and sham stimulation groups; ‡p Values shown in this table were analyzed between active cerebellar stimulation and cerebellar type stimulation group

degeneration.\(^3\) We now report here a placebo controlled trial of TMS over the cerebellum for patients with spinocerebellar degeneration.

Seventy four patients with spinocerebellar degeneration gave written informed consent to participate in this study, which was approved by the ethics committee of Tohoku University. No patient had a history of seizures or any abnormalities established by EEG. They also had no orthopaedic problems. Thirty nine patients, aged from 27 to 76 years (25 men and 11 women), were assigned to active stimulation, and the other 35 patients, aged from 38 to 76 years (19 men and 20 women), were assigned to active stimulation, and the other 35 patients, aged from 27 to 76 years (25 men and 11 women), were assigned to sham stimulation. They were divided based on the date when they were admitted to our hospitals. The age, disease duration (unpaired t test), disease type (x² test), and disease severity (unpaired t test) were matched between the two groups. Disease types were divided into two groups: cerebellar type (sporadic and hereditary cortical cerebellar atrophy including spinocerebellar atrophy (SCA) 6) and olivopontocerebellar atrophy (OPCA) type (sporadic OPCA, SCA1, SCA3, etc). Transcranial magnetic stimulation over the cerebellum was administered at almost the same time in the evening once a day for 21 consecutive days. A Magstim 200 (Magstim, Wales, UK), a transcranial magnetic stimulator with a 14 cm circular coil, was used. The stimulus coil was placed tangentially (active stimulation) or vertically (sham stimulation) over the scalp and centered on the inion, 4 cm lateral to the right of the inion, and 4 cm lateral to the left of the inion. The site order of delivery was always the same, first the right, then the centre, and then the left repeatedly 10 times in the same order. Ten pulses (the pulse interval was about 6 seconds, which depended on the time that the Magstim 200 needed to be fully charged; five were counter clockwise and five were clock wise) were delivered on each region. The duration of each stimulus pulse was 0.1 ms. The stimulator output was adjusted to 100% of the maximum output capacity, which was about 2.5 times the motor threshold of the participants. We measured the motor threshold by stimulating the hand area of the left cortex using a 14 cm circular coil and recording the motor evoked potentials from the right first dorsal interosseous muscle while the hands were relaxed. The average threshold was 39.4 (SD 3.8)%. We evaluated their truncal ataxia according to the time required for a 10 m walk (10 m time), the number of steps for a 10 m walk (10 m steps), and the number of practicable steps on a walk with the feet in the tandem position (tandem steps). Standing capacities (0, able to stand on one foot more than 15 seconds; 1, able to stand with the feet in the tandem position; 2, able to stand with the feet together; 3, able to stand with the heels together; 4, able to stand with the feet less than 10 cm apart; 5, able to stand with feet more than 10 cm apart; 6, unable to stand without support) and walking capacities (0, almost normal but unable to run; 1, able to walk without support, but clearly abnormal; 2, able to walk without support but with considerable staggering; 4, able to walk with a handrail; 5, able to walk with considerable support; 6, unable to walk, even with accompanying person) were also evaluated. The evaluation was done by the patients' physicians, who did not participate in this study. We measured the regional brain blood flow by consecutive single photon emission computed tomography (SPECT) using technetium-99m ethyl cystinate dimer before and after the 3 week TMS trial in 28 patients in the active stimulation group. We put the regions of interest on the bilateral frontal lobes, temporal lobes, occipital lobes, putamina, cerebellar hemispheres and pontine base. Quantitative analysis of the SPECT images was performed with a computer program (Patlak graphical analysis). The examiners were blind to the manner of TMS. All patients completed the 3 week TMS trial without any adverse effects. One patient was fearful and needed a reduction in the stimulus intensity to 80% of the maximum output. Before the TMS trial, there were no differences in 10 m time, 10 m steps, tandem steps, standing capacities, and walking capacities in both groups (Mann-Whitney U test). Comparing between active and sham stimulation after 3 week TMS, active stimulation was significantly more effective in 10 m time (p<0.05), 10 m steps (p<0.05), tandem steps (p<0.005), and standing capacities (p<0.05) (Mann-Whitney U test, table 1). However, 10 m time (p<0.05), 10 m steps (p<0.05), and standing capacities (p<0.05) were significantly improved in the sham stimulation group (Wilcoxon signed rank test, *table 1). The cerebellar type of spinocerebellar degeneration was significantly more sensitive to TMS than the OPCA type in 10 m time (p<0.05), tandem steps (p<0.05), and standing capacities (p<0.01) (Mann-Whitney U test, table 1). After the 3 week TMS trial, the mean regional blood flow significantly increased in the cerebellum and pons from 52.52 (SEM 2.12) ml/100 g/min to 58.54 (SEM 1.89) ml/100 g/min (<p<0.005), and from 34.38 (SEM 2.21) ml/100 g/min to 39.68 (SEM 1.34) ml/100 g/min (<p<0.05), respectively (paired t test). The regional brain blood flow in the cerebellar cortices did not show any change.

In this study, we found a significant alleviation of truncal ataxia in patients with spinocerebellar degeneration treated by active TMS. Although placebo or training effects were demonstrated in 10 m time, 10 m steps, and standing capacities (*table 1), the effects of active stimulation were far beyond those of sham stimulation. With our method, active stimulation evoked the contraction of nuchal and shoulder muscles. However, sham stimulation produced the same noise as that of active stimulation, and some scalp sensation. Patients did not know the difference between active and sham stimulation because no patients had experienced active stimulation previously. Therefore, patients having sham stimulation did not notice that they were receiving inactive stimulation. Our results showed that the disease type was important for the effect of TMS. We matched not only the age, disease duration, and disease severity of both groups but also the type of disease. We considered this study to be placebo controlled. Therefore, TMS over the cerebellum actually alleviates truncal ataxia in patients with spinocerebellar degeneration.
Although we do not yet know the mechanism by which TMS works, we are very much interested in the increase in cerebellar blood flow after TMS. This finding suggests that TMS over the cerebellum may activate the decreased cerebellar function. This effect may be caused by direct stimulation to the cerebellum as we used the maximum strength of stimulation at 2.5 times the motor threshold. Another possibility is that sensory input from the contracted muscles by TMS might occur.

After the TMS trial we continued TMS in some patients. Patients receiving TMS once or twice a week kept their improved condition at least for 6 months, but patients receiving TMS every 2 weeks returned to their baseline in 2 weeks. Therefore, we think that the effects of TMS with our method are maintained for about 1 week. We conclude that TMS over the cerebellum is a promising treatment for spinocerebellar degeneration.

Acknowledgement

This work was financially supported by the Magnetic Health Science Foundation.

References


Table 1: Examples of dictation. This patient showed intact Kana writing and difficulties in Kanji writing: 1, no response; 2, partial completion; 3, partial substitution; 4, neographism; 5, correct.

<table>
<thead>
<tr>
<th>Kanji</th>
<th>Meaning</th>
<th>Pronunciation</th>
</tr>
</thead>
<tbody>
<tr>
<td>犬</td>
<td>dog</td>
<td>(imu)</td>
</tr>
<tr>
<td>毛</td>
<td>wool</td>
<td>(keito)</td>
</tr>
<tr>
<td>灰</td>
<td>trash</td>
<td>(haizara)</td>
</tr>
<tr>
<td>鉛</td>
<td>pencil</td>
<td>(enpitsu)</td>
</tr>
<tr>
<td>新聞</td>
<td>newspaper</td>
<td>(shinbun)</td>
</tr>
</tbody>
</table>

Agraphia in Kanji after a contusional haemorrhage in the left temporo-occipital lobe

The Japanese language has two different writing systems, Kana (Japanese syllabograms) and Kanji (Japanese ideograms), and Japanese sentences usually consist of combinations of both. Recently, there has been speculation that different pathways are involved in Kanji and Kana reading and writing. It has been suggested that the processing of Kanji and Kana involves different intrahemispheric mechanisms, as judged mainly through studies on patients with alexia and agraphia. Wernicke's area and its surrounding left middle temporal lobe may play the most important part in Kanji reading when visual information is transmitted by any pathway. We recently had the opportunity to examine a patient who had mild transient aphasia and persistent agraphia of Kanji after a discrete contusional haemorrhage in the left temporo-occipital lobe. A 66-year-old right-handed man (pensioner) with 12 years of school education was involved in a traffic accident and sustained a closed head injury. He was admitted to our hospital the next day. He had no history of neurological problems. His parents, brothers, and sisters are all right-handed. On examination, he was alert and cooperative. Neurological examination disclosed no motor or sensory disturbance.

Formal language assessment was undertaken 3 days after injury using the standard language test for aphasia (SLTA). His spontaneous speech was fluent. He had no difficulty initiating speech, articulated normally, and did not have logorrhoea. Echolalia was not seen and phonological structure was clear. Object naming was moderately impaired, with considerable paraphasia. Word fluency (animal naming) was 3/min. Repetition was excellent. Reading aloud was good; however, writing of both Kanji and Kana was impaired, and considerable paragraphia was evident.

In addition, he demonstrated constructional apraxia using figure copying of a three dimensional cube but not oral, ideomotor, or ideational apraxia. There were no signs of motor impersistence or unilateral spatial neglect. The score on Raven's coloured progressive matrices was 26/36.

Brain CT on admission showed an area of high density consistent with contusional haemorrhage in the left temporo-occipital lobe. Brain MRI 5 days after admission showed contusional haemorrhage in the left temporo-occipital lobe involving the posterior part of the temporal cortex and the adjacent white matter. There were no abnormalities in the left angular gyrus.

Aphasia was gradually resolved, his scores on SLTA 8 weeks after the onset becoming normal. Despite this, his writing ability remained clearly disturbed. We asked him to read and write on dictation 221 Kanji and 76 Kana, all of which are taught in the first 2 years of primary school in Japan. He was unable to write 83 of the 221 Kanji (37.6%), which he reported that he knew but could not recall. He tended to have difficulty with Kanji characters, which were more complex and had been learnt later (fig 1). Error types included partial substitution (two), partial completion (seven), neographism (29), and no response (45). On the other hand, he made no mistakes in reading Kanji or in reading and writing Kana letters. On the revised Wechsler adult intelligence scale (WAIS-R), his verbal intelligence quotient (IQ) was 105 and his performance IQ was 101, with an overall IQ of 103. Brain MRI 3 months after admission showed a subcortical lesion in the left temporo-occipital lobe (fig 2).

Although this patient experienced aphasia immediately after cerebral contusion, the aphasia resolved over a subsequent 8 week period. He was left with an isolated disturbance of Kanji writing. In the absence of any dementia, aphasia, or disturbance of consciousness, his condition was classified as one of pure agraphia. Soma et al described “pure agraphia of Kanji” in three patients, who had a lesion in the left posterior temporal area extending to the angular gyrus on CT. Although they
exhibited amnestic aphasia, alexia, and agraphia in the acute phase, disturbances other than agraphia of Kanji disappeared in a few months. Yokota et al. also reported a case with pure agraphia of Kanji, and suggested that the process of writing Kanji involves a different pathway from that which mediates Kanji reading in the left temporal lobe.

Iwata proposed a hypothetical neuronal mechanism in the writing of Kanji and Kana as follows: the Broca’s and motor association areas are the final coordination centres, but the angular gyrus plays an important part in sending the graphic information to the motor areas. However, spontaneous writing and responses to dictation are usually initiated by Wernicke’s area, which gives rise to two different pathways to the angular gyrus. One is the auditory somaesthetic association pathway leading directly from Wernicke’s area to the angular gyrus; Kana writing mainly follows this pathway. The second pathway is from Wernicke’s area to the occipital lobe by way of the posterior inferior temporal area. This is the pathway involved in selecting the correct Kanji graphemes according to the meaning of the word, and thus recalled visual engrams of letters are sent to the angular gyrus.

Our patient demonstrated that initial amnestic aphasia, and agraphia of Kanji were associated with a lesion in the left temporal lobe. Because the posterior temporal region is located in close proximity to the angular gyrus and Wernicke’s area, the pathological process in the first area affects the second two regions in its acute phase. We conclude that the persistent symptom of pure agraphia for Kanji in this study was caused by the left posterior temporal lesion which disconnected the pathway for Kanji writing selectively.

Successful autologous stem cell transplantation in a patient with chronic inflammatory demyelinating polyneuropathy

A patient with chronic inflammatory demyelinating polyneuropathy is reported on who never had spontaneous remissions for 10 years, but who is in remission for 2 years after an autologous stem cell transplantation (ASCT). Before ASCT he needed at least 20 mg prednisone/day and in addition intravenous immunoglobulin (IVig) treatment at regular intervals. An ASCT was considered in this patient because of serious side effects of immunosuppressive treatment.

Chronic inflammatory demyelinating polyneuropathy (CIDP) is characterised by weakness with sensory impairments in the arms and legs. The reflexes in the arms and legs disappear. The onset is insidious. There may be progressive deterioration or a course with remissions and exacerbations. Routine blood examination is usually normal. Protein in CSF is increased in most patients. Electrophysiological studies may show slow nerve conduction velocities, conduction blocks, or dispersion. Inflammatory demyelinating polyneuropathy is considered to be an autoimmune disease which is supported by the presence of inflammatory cells in sural nerve biopsies and the beneficial response to immunosuppressive treatment in most patients.

We describe a patient with CIDP who has had this disease for 10 years without spontaneous remissions. He responded to immunosuppressive therapy, but needed high doses and had severe side effects. Therefore, we decided to try to induce a long lasting remission by autologous stem cell transplantation (ASCT).

This patient was 38 years old when, in 1988, he began to have tingling and numbness in his fingers. The sensory symptoms progressed to his arms and legs, followed by weakness. Initially his symptoms were attributed to too much stress but at the end of 1990 when weakness had worsened to such an extent that he was no longer independent in his daily activities, it was decided that it was time for a neurological evaluation. At that time he had weakness of his arms and legs, MRC grade 4. There was atrophy of his intrinsic hand muscles and fasciculations were seen in the muscles of his arms. Except for diminished knee reflexes, there was areflexia. He had numbness of his arms and legs in a glove and stocking distribution. Electrophysiological studies showed slowed nerve conduction velocities of the median, ulnar, tibial posterior, and peroneal nerves in the range of 15–25 m/s with prolonged distal latencies and F wave responses. The distal compound muscle action potentials were small. Few muscles had signs of denervation. Blood examination was unremarkable and CSF total protein was 0.67 g/l without cells. Sural nerve biopsy was consistent with CIDP, showing inflammatory cells and demyelination.

In 1990 he was started on 60 mg/day prednisone. After 2 weeks improvement began. This improvement continued to normal strength leaving only slight numbness in his fingers. During tapering off of the prednisone dose he had mood disturbances at every change of dose. Repeatedly he had a relapse at doses lower than 20 mg/day. Subsequently we tried to replace prednisone by 150 mg/day azathioprine but after 2 years of treatment with azathioprine it was still not possible to decrease the dose of prednisone to less than 20 mg/day. Azathioprine was replaced by methotrexate (7.5 mg/week) for 6 months without a beneficial response. Thereafter he was treated with IVIg. After two cycles of 30g/day IVIg for 5 days he needed infusions every other week to maintain his improved condition. We tried to prolong these intervals by adding prednisone. With 20 mg/day prednisone he remained free of relapses for about 4 months. However, these intervals became gradually shorter; he developed arthralgies after administration of IVIg and had reported gastric pain. After 8 years of immunosuppressive treatment, without spontaneous remissions, we discussed the possibility of autologous stem cell transplantation (ASCT).

In May 1998 peripheral blood stem cells were harvested by leukopheresis after mobilisation with cyclophosphamide (4 g/m²) and granulocyte colony stimulating factor (G-CSF; 5 μg/kg). CD 34+ cells were positively selected by immunomagnetic beads (Clinimax®, purity 98.7%) and cryopreserved. Before this procedure the patient had had IVIg every other week together with 20 mg/day prednisone. With this regular IVIg treatment he was in an optimal condition: he had no weakness and only mild sensory impairment of his fingers. After the mobilisation procedure, prednisone could be tapered off to 8 mg/day. Even without IVIg he remained in this improved condition for 5 months before he relapsed. Weakness in the arms and legs returned together with sensory impairment. Electrophysiological studies were repeated. We decided first that a neurological condition by IVIg treatment before myeloablative treatment was started. This treatment resulted in considerable improvement; only mild sensory impairment of the fingers remained.
In March 1999 he was treated with BEAM myeloablative chemotherapy (300 mg/m² BCNU on day –6, 200 mg/m²/day etoposide on days –5, –4, –3, and –2, 200 mg/m²/day cytosine arabinoside on days –5, –4, –3, and –2, and 140 mg/m²/day melphalan on day –1) followed by stem cell re-infusion at day 0 (11.2 x 10⁹ CD 34+ cells/kg). After re-infusion recovery of peripheral blood cells occurred within 16 days. Prednisone was tapered off but doses lower than 5 mg were accompanied by complaints of fatigue probably due to adrenal insufficiency. Two years after stem cell re-infusion, he is still free of relapses with 5 mg/day prednisone and without treatment. He works full time and the only neurological signs consist of mild spastic paraparesis of the lower limbs. Laboratory tests were negative except for her CSF showing increased protein content and pleocytosis up to 180 cells/mm³, mainly transformed but with no neoplastic lymphocytes, again none being immunopositive for the B cell markers CD 19 and CD 20. Extensive search for extranuclear malignancy including repeated cranial, spinal, and abdominal CT and MRI scans were all negative. The patient died 6 months after the onset of neurological symptoms. Necropsy disclosed diffuse B cell lymphoma restricted to the cauda equina with massive infiltration of the anterior lumbar spinal nerve roots and dorsal ganglia, and very mild neoplastic infiltrates in the sacral and femoral nerves associated with severe myelin loss. There was no involvement of the spinal cord, brain, meninges, or cranial nerves. The right brachial plexus showed severe demyelination without onion bulb formation and perivascular infiltrates exclusively composed of T lymphocytes. Search for systemic malignant lymphoma was negative except for an isolated small focal area of B cell lymphoma in the thoracic gland. As in the case reported by Wilkening et al., this finding widens the range of possibilities and differential diagnostic difficulties of selective involvement of the spinal roots (and peripheral nervous system) by primary malignant non-Hodgkin’s lymphoma, often also referred to as “neurolymphomatosis” making an extensive diagnostic investigation necessary.

References


CORRESPONDENCE

Unusual presentation of a primary spinal lymphoma

Wilkening et al. described the unusual history of a formerly healthy immunocompetent white woman initially presenting with an S1 syndrome with lymphocytic pleocytosis in the CSF, leading to the misdiagnosis of meningoradiculitis. Repeated spinal MRI disclosed an enigmatic mass lesion, and histological examination of the tumour biopsy confirmed the diagnosis of primary spinal lymphoma of Burkitt’s type. The authors emphasise that in every case with suspected meningoradiculitis including neuroorbitalis, meningoral lymphoma should be considered as an important differential diagnosis. In view of the occasionally difficult diagnosis of primary lymphoganglia we briefly present the clinicopathological report of a primary malignant lymphoma with almost complete involvement of the spinal roots: A woman aged 65 years without known systemic lymphoma presented with a painful focal right brachial plexus lesion and, 2 weeks later, developed progressive motor neuropathy with flaccid paraparesis of the lower limbs. Laboratory tests were negative except for her CSF showing increased protein content and 85 lymphocytes/mm³, not expressing the B cell markers CD 19 and CD 20. Nerve conduction studies suggested subchronic demyelinating neuropathy without motor conduction block. Cranial and cerebral CT and MRI were negative. Sural nerve biopsy disclosed a demyelinating neuropathy with acute myelin loss, some remyelination with incipient onion bulb formation, and wallerian degeneration, but without inflammatory or neoplastic infiltration. Muscle biopsy showed neurogenic muscle atrophy. Despite high dose corticosteroids and intravenous immunoglobulin treatment, her condition deteriorated rapidly with development of flaccid paraplegia and paréses of the right arm. Repeated CSF analysis showed increasing protein content and pleocytosis up to 180 cells/mm³, mainly transformed but with no neoplastic lymphocytes, again none being immunopositive for the B cell markers CD 19 and CD 20. Extensive search for extraneural malignancy including repeated cranial, spinal, and abdominal CT and MRI scans were all negative. The patient died 6 months after the onset of neurological symptoms. Necropsy disclosed diffuse B cell lymphoma restricted to the cauda equina with massive infiltration of the anterior lumbar spinal nerve roots and dorsal ganglia, and very mild neoplastic infiltrates in the sacral and femoral nerves associated with severe myelin loss. There was no involvement of the spinal cord, brain, meninges, or cranial nerves. The right brachial plexus showed severe demyelination without onion bulb formation and perivascular infiltrates exclusively composed of T lymphocytes. Search for systemic malignant lymphoma was negative except for an isolated small focal area of B cell lymphoma in the throracic gland. As in the case reported by Wilkening et al., this finding widens the range of possibilities and differential diagnostic difficulties of selective involvement of the spinal roots (and peripheral nervous system) by primary malignant non-Hodgkin’s lymphoma, often also referred to as “neurolymphomatosis” making an extensive diagnostic investigation necessary.

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References


Authors’ reply

We greatly appreciate the interesting comments of Jellinger and Grisold who describe...
another unusual case of primary spinal lymphoma. In their case, by contrast with that presented by us the spinal roots were selectively involved without the formation of a mass lesion during the disease. Moreover, CSF studies in their case showed increased protein content and pleocytosis but no neoplastic cells, which had been found in our patient. Together, the case presented by us and the case of Jellinger and Grisold underlines the fact that the range of primary cerebral and especially primary spinal lymphoma is very wide with different clinical presentation, location of manifestation, prognosis, and probably of aetiology. More cliniconeuropathological studies on primary spinal lymphoma are necessary to better understand this disease and to develop therapeutic strategies for its different types.

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Orientation agnosia in pentagon copying
Ala et al have recently claimed that patients with Alzheimer's disease (AD) were less likely to make errors on the pentagon copying component of the mini mental state examination (MMSE) than patients affected by Lewy body (LB) dementia. Using (standard and modified) MMSE scoring criteria they found normal copies in 16 of 27 patients with AD but in only two of 13 patients with LB. Although this difference was non-significant, they concluded that this feature might be useful in differential diagnosis. It is of some interest that the MMSE scoring criteria employed by Ala et al did not regard errors of rotation as abnormal. In the light of recently identified visuospatial deficits which affect the domain of orientation, it may well be that the criteria of Ala et al for a “normal” copy are inappropriate. This has implications for the claimed diagnostic power of their measure.

Patients with orientation agnosia are able to recognise objects, but have a selective visuospatial impairment, in that they have lost knowledge of object orientation. The most striking feature of this disorder is shown in the patient's drawing of objects, where they rotate their copy by 90 or 180 degrees relative to the original. This previously underinvestigated deficit is far from rare. In one study, seven of 63 (11%) patients with stroke grossly rotated their copy of the MMSE. All had a right hemisphere lesion and showed some evidence of left visuospatial neglect. In a second study 16 of 240 (7%) of a series unselected by pathology showed the same sign, but without clear lateralising features.

To better investigate this issue in degenerative disease, we reviewed the records of a consecutive series of 134 patients with AD. All had been given the MMSE, which includes the interlocking pentagon figure (aligned horizontally) as a copy task, and a battery of neuropsychological tests. Errors in copying the geometrical figure were systematically graded. In general agreement with the findings of Ala et al, 57 of 134 (43%) drew unrecognisable figures, adding or omitting details.

Three showed signs of closing in, and seven (3% of the sample) made clear rotation errors, misplacing one or both pentagons by 90, 120, or 180 degrees relative to the original. These seven patients did not differ from the overall sample in terms of general severity of dementia, as measured by the MMSE total score, and by daily activity scales. Eight patients (6%) presented with signs of visuospatial neglect, but patients with neglect and orientation agnosia did not overlap in this sample, suggesting that orientation agnosia should be considered an independent sign. Our data thus suggest that orientation agnosia is a significant visuospatial deficit in degenerative disease, which is not identified as such in the study of Ala et al by virtue of the criteria by which they define an abnormal performance.

In figure 2 of the paper of Ala et al, it is clear that some copies, scored as errorless, showed clear signs of orientation agnosia (AD cases 3, 9, 15, 18, 19, and 24). If these errors are considered, the claimed difference between AD and LB dementia disappears, confirming the results of previous studies which showed no significant difference in drawing or constructional abilities between AD and LB dementia. 5

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References

Authors' reply
We were very pleased to see the interest that Della Sala and colleagues have shown in our study. 1 Orientation agnosia is certainly a fascinating subject, and we applaud their research into its neuropsychology. We think, however, that their analysis of our study has several weaknesses.

Firstly, because we had not included the original interlocking pentagon models with the copies in our figures, they incorrectly inferred that six of our patients with AD made rotational errors. As mentioned in our results, many different printed models of the interlocking pentagons were used in our study. To clarify for your readers exactly what the patients were shown to copy and how they performed, Figure 1 presents the models and the patients' copies for AD cases 3, 9, 15, 18, 19, and 24. To allow the study of rotational changes, the exact orientation to the bottom of the page is preserved for each model and for each copy. Although all have been reduced for publication, the relative sizes of the models and the copies are also preserved. By our inspection none of the questioned cases exhibits any significant rotational error. We also studied the copies of the remaining patients with AD and the patients with DLB in our study, and we could not form any conclusions about rotational errors with either group.

It should also be noted that we specifically used the original published criteria 2 for grading the copies of the interlocking pentagons in the MMSE. These criteria state that "tremor and rotation are ignored" (table 1 of our article). 2 The diagnoses of our patients were neuropathologically proved. It would be interesting to know if the diagnoses of the 134 patients with AD that Della Sala and colleagues mention were also neuropathologically proved. If their patients were diagnosed clinically, we...
would not be surprised if some patients with DBL were included in the 134. Perhaps many of those who had the orientation agnosia were actually patients with DBL who did not exhibit the classic DBL clinical symptoms of fluctuation, visual hallucinations, and parkinsonism. Also, one of the patients in our study (minimum MMSE 13). That study does help con firm that constructional abilities worsen as the dementia worsens, for as we mentioned in our results, none of the additional cases of AD or DBL in our brain bank with initial MMSE scores less than 13 had acceptable pentagon copies.

Finally, albeit a minor detail at variance with Della Sala and colleagues, we did report that the group copied the pentagon significantly less accurately than the AD group (P=0.002 using the original criteria).

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References

BOOK REVIEWS

Advances in research on neurodegeneration. Volume 7

By Yoshikuni Mizuno, Donald B Calne, Reinhard Horowski, Werner Poewe, Peter Riederer, and Moussa Youdim (Pp 214, US$69.95). Published by Academic Press, San Diego, 2000. ISBN 0 12 298190 1

This book gives comprehensive and thorough up to date coverage of this topic. It starts from the premise that neuroprotective treatment of older people is a useful and productive endeavour, and then sets out to assist in making the process as efficient as possible, while being clearly respectful and humane. It guides the reader through every aspect of the process, from tasks which need to be addressed before the patient arrives, through the clinical interview and the formal testing, onto interpretation of test findings and, finally, to the rounding off with providing feedback and follow up. The practical aspects are supported by example letters, documents, forms, and reports, together with 11 useful case studies. In addition there are chapters reviewing in detail assessment procedures for memory and other cognitive functions, and four useful chapters examining the neuropsychological profiles of the most common (and a few rarer) conditions likely to be encountered in clinical practice, from Alzheimer’s disease through to normal pressure hydrocephalus. The important topic of depression, its nature, and its impact on neuropsychological performance, receives a chapter in its own right.

The author is to be congratulated on producing a book which travels well outside the North American context in which it was written; by largely concentrating on the Wechsler tests (and the most up to date versions of them) the detailed discussion of tests and their uses and properties seems familiar and accessible. The book is soundly based in current research and theory—for example, on Lewy body dementia and the vascular syndromes—but also conveys a strong sense of clinical experience and wisdom. The reader will not find here a defence of the contribution of neuropsychology to the management of cognitive disorders in older people, but a better account of the state of the art would be difficult to imagine.

Robert Woods

Thrombolytic therapy for stroke


The end of the 20th century saw a transition in the approach to acute stroke from therapeutics nihilism to enthusiasm. Stroke is now...
Neurobiology of spinal cord injury


"The future for regenerative therapies of the spinal cord is clearly very bright"—thus ends this volume in Humana's Contemporary Neuroscience series, which looks at different aspects of spinal cord injury with special attention to possible strategies for repair.

Mechanisms of neuronal damage after injury, locomotor function in the transspinal spinal cord, axon extension and elongation, as well as pharmacological and cell transplantation strategies to encourage axon regrowth and remyelination of remaining fibres, and current therapies for patients with spinal cord injury are all covered. Individual chapters were contributed by experts in the field, many of whom carried out seminal work on the topics they describe. This ensures that the text is on the whole, up to date, and written with authority, although sometimes a little too focused on subjects close to the authors' hearts. Consistency in style, layout, and detail between the different chapters is somewhat lacking, which doesn't make this a particularly enjoyable read.

Much of the book concentrates on the study of spinal cord injury models and the insights gained about mechanisms of cell death and loss of function as well as the usefulness of repair strategies, with two of the 11 chapters dedicated to cell transplants. The text devotes very little attention to the clinical relevance of experimental findings, and is not really concerned with experimental findings gained from human spinal cord injury. None of the current pharmacological therapies for patients and their possible mechanisms of action are discussed in the last chapter.

This would be a good introduction for those who are unfamiliar with mechanisms of neuronal injury and repair, and also provides a good update on recent developments in experimental strategies for repair. Those with some experience in this subject will probably find that this volume does not provide sufficient detail in most areas to be an essential addition to their bookshelf.

Given that many of the authors have spent much of their careers on possible strategies for spinal cord repair, it is perhaps not surprising that this book exudes a general optimism in this regard. Certainly, progress has been made in several different areas, which may one day bring substantial benefit to patients, either alone or combined—details of each, and how they may compliment each other—are probably the main things that the reader will take away from this.

Rike Zietlow

Cognitive neurorehabilitation


As the French term "re-education" implies, learning is a key process in rehabilitation. Nothing might seem more challenging than to effect useful learning in people with cognitive impairments. Until recently, a cloud of pessimism hung over cognitive deficits, but the later sections of this useful book provide

Martin Brown

www.jnnp.com
authoritative evidence that interventions can produce positive changes in people with impairments in memory, in attention, perhaps in executive function, and in language processing.

Is it rehabilitation? Most of the convincing effects described here are at impairment rather than disability level and therefore are not necessarily positive rehabilitation outcomes. But the book takes account of social factors, as well as problematic psychological categories such as motivation, which provide a pragmatic context within which future interventions might operate. It seems likely that more will be achieved in this new field.

Is it cognitive? One source of pessimism about therapy for cognitive impairment has been the dead hand of verbal learning, the “old paradigm” for cognitive science. It is hard, and often futile, to teach word lists to people with amnesia. A much more achievable goal is to alter their behaviour. A sort of neobehaviourism is creeping into cognitive science, inspired by artificial learning (neural) networks. The most relevant aspect of a chapter on the potential benefits of constraining an unaffected limb in hemiplegia is the account of “learned non-use” as a possible barrier to inducing movement in the plegic limb. Might the same mechanism lead a person with aphasia to use less speech, or someone with amnesia to become more restricted? Learned non-use can be described in terms of operant conditioning, as can other therapeutically fruitful concepts such as errorless learning. A good deal of cognitive rehabilitation is not really cognitive. The restitution of cognition (in its classical sense, as the handling of information in the conscious domain) still eludes us. But this book provides some signposts along the way.

Christopher D Ward

Neuropathology and genetics of dementia: advances in experimental medicine and biology, volume 487


This book represents the Proceedings of the XVIIIth International Winter Meeting of the Swiss Society of Neuropathology, held in St Moritz in March 2000. This is written as a series of short articles and longer reviews, rather than the usual unstructured cluster of synopses of experimental findings presented orally at such scientific gatherings. Thematically, the review articles focus on the role of abnormal aggregated or misfolded proteins in the brains of persons with diverse neurodegenerative diseases. Accordingly, there are chapters detailing the pathology and genetics of tauopathies, amyloidopathies and synucleinopathies, coupled with other chapters covering ubiquitin and polyglutamine disorders, inclusion body myositis and neuronal ceroid lipofuscinoses. The possible role of mitochondrial dysfunction in such disorders is discussed. Overall, the chapters are of a uniformly high standard and of sufficient breadth and depth to be useful and interesting to the specialist and casual reader alike. Rapid publication has ensured that most contributions are still well within their shelf life, although the camera ready format in which the volume has been prepared means the inclusion of occasional typographical and referencing errors, and a rather poor reproduction quality to the figures, particularly the half tones. The last is a particular shortcoming, given the weight of text devoted to neuropathological description. None the less, there is much useful information in this book, set out in a readable and up to date manner. It can be recommended particularly to junior researchers and physicians, who will find it a comprehensive and easily accessible reference. Lastly, it is noted in the acknowledgements that the editors were ably and fortuitously assisted by a certain Michelle Pfeiffer—surely not that one!

D M A Mann
A COMPARISON OF NEUROPSYCHOLOGICAL ASSESSMENT AND MRI IMAGING SCORES IN THE EARLY DIAGNOSIS OF ALZHEIMER’S DISEASE

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The aim of this study is to address whether neuropsychological measures or imaging scores best predict which patients with questionable dementia progress to fulfilling criteria for Alzheimer’s disease (AD). MRI images of 50 subjects (18 with early AD, mean MMSE 21.4, and 32 with memory complaints, mean MMSE 27.3) were examined using the temporal lobe rating scale. Subjects underwent a standard battery of neuropsychological tests at baseline, and were clinically followed up for a mean of 19 months. Patients who converted to AD (converters n=11), relative to non-converters, were impaired at baseline assessment on general cognitive tests (Addenbrookes cognitive examination (ACE) and ADAS cog), measures of episodic memory, category fluency, and the graded naming test. The mean scores of the temporal lobe rating scale demonstrated that converters had more hippocampal and parahippocampal atrophy than non-converters. Discriminant analysis demonstrated that the best test for distinguishing the converters from the questionably group, was the ACE score. In combination, the medial temporal lobe rating and graded naming test were also contributory.

MRI imaging measures may be valuable if used in conjunction with neuropsychology, but progression to dementia is best predicted by performance on neuropsychological tests.

EARLY COGNITIVE DECLINE IN CREUTZFELDT-JAKOB DISEASE OCCURRING IN RECIPIENTS OF PITUARY-DERIVED HUMAN GROWTH HORMONE

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Most cases of Creutzfeldt-Jakob disease (CJD) in recipients of human cadaveric growth hormone, present with a cerebellar syndrome. Dementia is thought to occur late and as a minor feature of the illness. However, neuropsychology data published on these cases is largely qualitative and anecdotal. The first published case, of a 22 year old woman, does include neuropsychology performed 7 months after the onset of a cerebellar syndrome. This showed evidence of intellectual decline, with a verbal IQ of 80. Subsequent reports hint at the possibility that cognitive problems may be present in the initial stages of the illness.

In order to assess early cognition, we report detailed assessment at referral (mean 4.5 months from the onset of symptoms, range 4–6 months) in five patients with histologically proved human growth hormone derived CJD. All presented with a cerebellar syndrome and only one had noticed mild memory problems. On formal testing however, four had demonstrable, mild intellectual decline, as measured on the WAIS-R. One case showed selective visual memory impairment and one had noticed mild memory problems. On formal testing however, four had demonstrable, mild intellectual decline, as measured on the WAIS-R. One case showed selective visual memory impairment and the other had noticed mild memory problems. On formal testing however, four had demonstrable, mild intellectual decline, as measured on the WAIS-R. One case showed selective visual memory impairment and the patient who converted to AD (converters n=11), relative to non-converters, was impaired at baseline assessment on general cognitive tests (Addenbrookes cognitive examination (ACE) and ADAS cog), measures of episodic memory, category fluency, and the graded naming test. The mean scores of the temporal lobe rating scale demonstrated that converters had more hippocampal and parahippocampal atrophy than non-converters. Discriminant analysis demonstrated that the best test for distinguishing the converters from the questionably group, was the ACE score. In combination, the medial temporal lobe rating and graded naming test were also contributory.

MRI imaging measures may be valuable if used in conjunction with neuropsychology, but progression to dementia is best predicted by performance on neuropsychological tests.

LATERALITY OF UNEXPLAINED MOTOR AND SENSORY SYMPTOMS: A SYSTEMATIC REVIEW 1965–2000

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Background: Unilateral unexplained motor and sensory symptoms present commonly to neurologists. The literature has suggested that left sided symptoms are far more common than right.

Methods: A systematic review of the literature since 1965 to test this hypothesis. All studies describing one or more patients with unexplained, hysterical, psychogenic, or non-organic sensory or motor symptoms (excluding pain, visual, and auditory symptoms) were included.

Results: 124 studies with 1045 patients were analysed. The overall proportion of left sided weakness and sensory symptoms was 57% (CI 53%–60%). This figure was accentuated in studies which specifically set out to study this question (64% on the left). However, when laterality was reported incidentally there was no significant effect (51% on the left). Unexplained movement disorders were left sided in 38% (CI 31–38%). Female sex was related to left sided symptoms. Handedness and date of study did not influence laterality.

Interpretation: The data do not provide strong evidence for lateralisation of unexplained motor and sensory symptoms except for right sided movement disorder and possibly left sided symptoms in women. Retrospective studies specifically looking at lateralisation may have produced misleading positive results through bias. Consideration of the biology of these phenomena should not place undue weight on laterality.

HAEMOGLOBIN CONCENTRATION AND STROKE RISK IN 13 000 PATIENTS WITH ESTABLISHED CEREBROVASCULAR DISEASE

SC Howard, DA Power, PM Rothwell on behalf of Cerebrovascular Cohort Studies Collaboration Radcliffe Infirmary, Oxford, UK

Small studies and case reports have suggested that polycythaemia may increase the risk of ischaemic stroke. There is physiological evidence to support this, but there have been no adequately powered cohort studies. We therefore pooled detailed individual patient data from cohort studies and clinical trials in which patients presenting with a TIA or non-disabling ischaemic stroke had haemoglobin measured at baseline and had been followed up.

Using data on 13 009 patients with 60 000 patient-years of follow up from eight independent studies, the relation between haemoglobin and risk of stroke was assessed by Cox regression adjusting for age, sex, other baseline factors associated with haemoglobin, and stratifying by study population.

Stroke risk had a "U" shaped relation across eight predefined categories of haemoglobin concentration (p=0.03). This was most marked for fatal ischaemic stroke (p=0.001), haemorrhagic stroke (p=0.02), and recurrent ischaemic stroke in patients presenting with a lacunar event (p=0.005). The respective hazard ratios for haemoglobin above the normal range were 3.57 (95% CI=1.87–6.84), 2.97 (95% CI=1.07–8.20), and 1.49 (95% CI=0.79–2.81).

Haemoglobin concentration is an independent predictor of ischaemic and haemorrhagic stroke in patients with established cerebrovascular disease. Correction of mild polycythaemia or anaemia may reduce the risk of stroke.

A COMPARISON OF CT VERSUS DIFFUSION WEIGHTED IMAGING IN HYPERACUTE STROKE USING A SYSTEMATIC QUANTITATIVE SCORE (ASPECTS)

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Background: Tissue plasminogen activator (TPA) when administered <3 hours of symptom onset for acute ischaemic stroke produces improved functional and neurological outcome. Recent advances in neuroimaging with magnetic resonance (MR) diffusion weighted imaging (DWI) have raised hopes regarding the accurate identification of ischaemic tissue with improved safe administration of the treatment. CT is as good as DWI when the scans are read by experienced observers.

Methods: All patients had a CT scan (within 6 hours of stroke onset) prior to MRI (<7 hours of onset). Non-contrast enhanced CT scans were performed on four generation scanners with 5 mm slice thickness parallel to the infraorbital meatal line, and appropriate window width (80 HU) and centre level review settings (40 HU). MRI imaging was performed on a 3Telsa machine and included DWI (single shot EPI TR=7000 ms, TE=99 ms, b=1000 mm s⁻¹, 19 5 mm slices/5 mm gap). A quantitative scoring system (ASPECTS) was applied to CT and DWI at baseline and follow up (<36 hours) by five
independent observers (three stroke neuroradiologists and two neuroradiologists) with only knowledge of the stroke side. The reliability of each modality was assessed for each observer with intra-class correlation coefficients. Bland Altman plots were used to measure the agreement between CT and DWI using ASPECTS.

Results: 80 patients (63% male; mean age 68.4±13.2) had CT and DWI imaging. The median NIHSS was 9. The mean onset to CT (136 minutes) compared with onset to MRI (240 minutes) was 0.001. The median baseline CT and DWI ASPECTS values were equal at 8. Intraclass correlation coefficients confirmed excellent reliability between observers for CT (0.81 [95% CI 0.76–0.86]) and DWI (0.76 [95% CI 0.72–0.84]) at baseline. CT ASPECTS predicted DWI ASPECTS (linear regression p=0.001) at baseline. The difference of means between DWI and CT at baseline was 0.3 ASPECTS, allowing for the mean difference in time for the scans.

Conclusions: CT appears comparable with DWI for assessing patients with acute stroke and when viewed by experienced observers using ASPECTS, and when sound CT imaging quality is achieved. The CT scan remains the imaging modality of choice in hyperacute ischaemia using ASPECTS, and when sound CT imaging quality is achieved.

**HISTORY OF STROKE IN FIRST DEGREE RELATIVES OF PATIENTS WITH TIA AND ISCHAEMIC STROKE IN RELATION TO CLINICAL SUBTYPE AND INTERMEDIATE PHENOTYPES**

UGR Schulz, CP Warlow, PH Rothwell Radcliffe Infirmary, Oxford, West- ern General Hospital, Edinburgh, UK

Some rare mendelian stroke syndromes have been characterised, but the genetic epidemiology of the common forms of ischaemic stroke and TIA is uncertain. Previous studies have not considered specific clinical and aetiological subtypes or assessed whether familial susceptibility is related to intermediate phenotypes.

We studied family history of stroke in 545 ischaemic strokes (Oxfordshire Community Stroke Project) and 469 Tias (hospital based series). We extracted related history of stroke in first degree relatives (FDRs) to assess baseline clinical characteristics, subtype of stroke, and risk of recurrence.

Information on family history was available in 537 strokes and 467 Tias.

Stroke had occurred in at least one FDR in 101 (18.9%, 95%CI 16–22) strokes and in 98 (21%, 95%CI 17–25) Tias. In both cohorts, a history of hypertension was strongly associated with family history (p<0.01) and became more frequent with increasing numbers of affected FDRs (p<0.01). However, there was no relation between family history and hyperlipidaemia, diabetes, stroke subtype, or risk of recurrence.

We found no evidence of a major independent genetic component in Tia and ischaemic stroke. However, the strong link between family history and stroke and hypertension suggests that genetic susceptibility to hypertension may explain a substantial proportion of the familial clustering of cerebral ischaemic events.

**A LATERALITY SHIFT OF BROCA’S AREA IN THE RECOVERY OF SPEECH PRODUCTION AFTER STROKE**

SC Blank, RJS Wise Hammersmith Hospital, London, UK

The production of both propositional speech (PrSp) and non-propositional speech (NPSp) in normal subjects activates the left fronto- operculum (Broca’s area). We used PET to investigate speech production in five aphasic stroke patients with lesions that included the left frontal operculum.

All patients initially presented with severe non-fluent aphasia. Patients had survived 16 months—10 years after stroke (median 4 years), when recovery of aphasia was deemed to be maximal. All subjects could count aloud fluently, but the extent of recovery of PrSp varied considerably, from occasional single word output to sentences. Subjects were scanned under three experimental conditions; counting aloud (NPSp), PrSp in response to autobiographical questions, and a non-speech state (XSp) when they attended to occasional environmental sounds. In normal subjects, PrSp and NPSp contrasted with XSp revealed activation within Broca’s area and the left anterior insula. In patients, activation was seen in the right fronto opercular cortex. Thus, in patients with extensive damage to left premotor areas for articulation, homologous regions within the right hemisphere are recruited for the motor act of speech. This is direct evidence of plasticity within the neural systems engaged in speech production, and confirms Moberg’s hypothesis about laterality shift of function in the frontal operculum.

**CLINICAL FEATURES AND DISEASE COURSE OF PSP IN THE UNITED KINGDOM**

U Nath, Y Ben-Shlomo, RG Thomson, N Wood, AJ Lees, DJ Burn Royal Victoria Infirmary, Newcastle upon Tyne, UK

Objective: To conduct a clinical survey of PSP and also to estimate its current prevalence rate in the UK.

Methods: 187 cases of PSP were identified in the context of a United Kingdom based prevalence study. Record-based confirmation of diagnosis according to the NINDS-SPSP criteria were performed in all cases and structural clinical assessment was also performed in a third of these cases. Information relating to subsequent death of patients was compared with national death certificate data collected all patients recorded as having PSP according to the ninth version of the International Classification of Diseases (ICD9).

Results: The median age of the clinical cohort was 72 years (range 40–92). Specific clinical features such as diplopia and eyelid apraxia occurred in 39% and 17% of all patients but data on these items was not routinely recorded. Information from those patients undergoing standardised assessment (n=49) revealed frequencies of 61% and 43% for these features respectively. Tremor was present in 21% of all cases and information from clinically examined cases suggested that this was usually postural. Early onset of diplopia and cases without tremor appeared to be associated with a shorter disease survival. PSP was mentioned in 57% of death certificates of cases identified by record review.

Conclusion: PSP is frequently misdiagnosed. Visual symptoms are prominent and often disabling. Clinical factors such as early onset of diplopia are significantly associated with poorer survival. Mortality rate for PSP is significantly underestimated.

**PROGRESSION OF NIGROSTRIATAL DYSFUNCTION IN A PARKIN KINDRED: AN 18F-DOPA PET AND CLINICAL STUDY**

NL Khan, DJ Brooks, N Pavee, NW Wood, AJ Lees, P Piccini Institute of Neurology, Queen Square, London; Imperial College School of Medicine, Hammersmith Hospital, London; Royal Free Hospital and University College Medical School, London, UK

We have used 18F-dopa PET to serially study a family with young onset parkinsonism with mutations in the parkin gene whose clinical picture is characterised by the presence of severe resting leg tremor. Four patients have been studied twice, 10 years apart, to assess disease progression. Additionally, we have studied five asymptomatic family members; four of whom carry a single parkin mutation and one individual with a normal genotype. Two of the carriers and the individual with the normal genotype had a repeat 18F-dopa scan. In the parkin patients the rate of loss of 18F-dopa uptake was similar in caudate and putamen (1.3% and 1.2% per annum respectively). The rate of disease progression is slower than that in idiopathic PD. The group of asymptomatic carriers also showed significant dopaminergic dysfunction in striatum bilaterally and in ventral midbrain and three of them developed subtle extrapyramidal signs. However in the two carriers scanned twice, 18F-dopa dysfunction did not progress with time, at least over a 7 year period.

The slow rate of disease progression in parkin patients may explain the near normal longevity of some young onset PD patients. The development of mild signs of basal ganglia dysfunction in three carriers with abnormal striatal 18F-dopa uptake suggests that these individuals are “manifesting heterozygotes” and may eventually develop parkin disease.

**UNILATERAL TRANSPLANTATION OF HUMAN PRIMARY FETAL TISSUE IN FOUR PATIENTS WITH HUNTINGTON’S DISEASE: NEST-UK SAFETY REPORT**

AE Rosser on behalf of the NEST-UK consortium Universities of Cardiff, Cambridge, Aberdeen, Belfast, London, and Manchester, UK

Huntington’s disease (HD) is an inherited CAG repeat disorder typically manifesting in mid-life and leading to death within approximately 20 years. The principle pathology is focal degeneration of the major projection neurons of the striatum, making cell replacement by neural transplantation a potential treatment strategy. There is substantial experimental evidence that neural transplants can yield functional recovery in animal models of HD and several clinical trials have commenced. To date, only a few HD patients have received operations and the approach remains experimental. We report here the first phase of a Medical Research Council (MRC) supported clinical safety trial of neural transplantation in four patients with mild-moderate HD who have received unilateral grafts of human fetal striatal tissue. This is a multicentre UK-based trial affiliated to the organisation NECTAR.
(European Network for CNS transplantation and repair). At the time of writing all significant adverse events relate to the immunotherapy. There were no acute surgical events; no evidence of tissue contamination or overgrowth; and no evidence that the procedure has produced acceleration of the clinical course. Neuronal transplants would be expected to take at least 12 months to establish and to start to ameliorate function, thus the efficacy of these procedures will be reported at a later date.

FAST MICROGRAPHIA AND PALLIDAL PATHOLOGY
N. P. Quinn Institute of Neurology, Queen Square, London, UK

Micrographia is well recognised in Parkinson’s disease (PD). Typically, the initial size and speed of the patient’s handwriting is good, but rapidly becomes progressively smaller and slower, and may ultimately grind to a halt. There exists another, quite distinct, form of micrographia which I call “fast micrographia”, in which handwriting is microscopic from the outset, does not (indeed cannot) become progressively smaller, is accomplished at (astonishingly) normal speed, and does not fatigue. However, this micrographia can be so marked as to resemble a straight line, so that even the patients are often unable to read what they have written. In contrast with the micrographia of PD, many patients with fast micrographia have completely normal alternating finger movements.

Fast micrographia is most often observed in patients with progressive supranuclear palsy, who sometimes volunteer “microscopic” writing as an expediting complaint. However, it can also be seen in patients with acquired bilateral lesions of the globus pallidus. Illustrative videotapes will be shown.

Fast micrographia should be considered a clinical feature suggestive of underlying pallidal pathology.

PROSPECTIVE STUDY OF DEVELOPMENT OF INFANTS BORN TO MOTHERS WITH EPILEPSY
F. H. James, S. Fairgrieve, S. A. Lynch, M. J. Jackson University of Newcastle upon Tyne, Newcastle, UK

Objective: to establish if infants born to mothers with epilepsy are at increased risk of developmental delay.

Methods: Our prospective study of pregnant women with epilepsy identified 292 babies whose mothers gave consent to follow up with developmental assessment using Bayley’s scales.

Results: We have examined 149 babies at 2–3 years to date: 59 were exposed to valproate, 61 to carbamazepine (50 in each group were on monotherapy). 31 children (21%) have delayed development and 13 (8.7%) have severe delay when the population prevalence of severe delay = 1.3% (OR 6.27, 95%CI 3.2–11.1). Developmental delays correlated with exposure to valproate (OR 4.42, 95%CI 1.77–11.5). However, mothers of children with epilepsy had low educational achievement; only 47% achieved five GCSEs (grade A–G) compared with 85% of the population. Only 26% of mothers of delayed children obtained five GCSEs.

Conclusion: Children born to mothers with epilepsy in this study have increased risk of developmental delay at 2–3 years. The risk is increased with exposure to valproate and with maternal educational underachievement. Examination of the rest of the cohort followed by multivariate analysis will be undertaken in an attempt to identify the cause of developmental delay in these infants.

A PROFILE OF EPILEPSY IN A UK POPULATION

Objective: To describe the characteristics of epilepsy in a representative UK sample, including demographic features; onset age; duration; severity; antiepileptic drug (AED) use and the impact of epilepsy on life and work.

Methods: A large, geographically comprehensive survey using a postal questionnaire distributed by general practitioners to 3455 unselected patients receiving AEDs for epilepsy, in all UK regions.

Results: There were 1652 replies. The mean age was 44 years; 47% males. The mean onset age, 25 years, and the mean duration, 20 years, were comparable with epidemiological studies. In the preceding year, 52% of patients had no seizures; 8% a single seizure, 17% 2–9 seizures, and 3% 10 seizures; 64% had mild epilepsy and 32% severe. There was a marked and significant decrement of seizure frequency with increasing age. The most common AEDs were carbamazepine (37%), valproate (36%), phenytoin (29%). Mono- therapy was used in 68% of patients. Patients taking multiple AEDs reported significantly higher adverse effects. The major impacts on life were work and school difficulties, driving, and psychological, with variation according to age and severity.

Conclusions: Seizures remain uncontrolled in half of people with epilepsy in the UK with significant psychosocial impact. Our findings help address the deficiency of information on the characteristics of epilepsy in the elderly, in whom the condition is an increasing health problem.

THE MANAGEMENT OF ADULTS WITH EPILEPSY
D. Smith, I. Crook, M. Williams on behalf of the North West Clinical Neurosciences Partnership Project Walton Centre for Neurology and Neurosurgery, Liverpool, UK

Suspected and established epilepsy are common reasons for referral to hospital clinics. Optimal management, ideally, involves cooperation between an epilepsy clinic and primary care. Because these facilities are not widely available these patients can be exposed to major diagnostic, incomplete treatment and suboptimal therapy and inadequate counselling. There is a need for expansion or reorganisation of services and concise guidelines could facilitate the care of these patients.

The North West Clinical Neuroscience Partnership project has developed a framework for service provision for adults with possible or definite epilepsy. A multidisciplinary team made contributions, based on reviews of available evidence, to the guidelines document. This document contains recommendations on (a) a structure for service provision, (b) diagnosis of epilepsy, and (c) the management and follow up of newly diagnosed and refractory epilepsy and includes specific advice regarding choice of treatment and the management of epilepsy in women, people with learning disability and the elderly.

This framework could facilitate best practice/shared care in either established or evolving services. Specific aspects of care are easily amenable to prospective audit. Adherence to guidelines is a transparent method of demonstrating the accountability of a service assuring commissioners/patients that a system capable of delivering quality care is in existence.

CLINICAL FEATURES OF PARANEOPLASTIC NEUROLOGICAL DISORDERS: A RETROSPECTIVE STUDY

There is paucity of information about paraneoplastic neurological disorders (PND), a heterogeneous group of immune mediated conditions associated with a variety of different tumours. We report the preliminary results of a retrospective observational study investigating the clinical features of 57 patients with PND, ascertained principally via the British Neurological Surveillance Unit. The average (SD) age of onset was 63 (11) years and there was a female preponderance
The most often reported syndromes were sensory neuro(enceph)alopathy (33%) and cerebellar degeneration, either pure (21%) or in association with brainstem/limbic encephalitis/neuropathy (23%). SCLC was the most frequent underlying tumour type (28%), followed by breast (16%), ovarian (9%) and other lung (7%) carcinoma. No tumour was identified in 19%. Forty six per cent of patients received immunomodulatory therapy and 67% of patients had their tumour treated. Only a minority of patients responded to these treatments and often only briefly. Progressive deterioration was the rule and 40% of patients with adequate follow up were dead at 12 months. We conclude that the diagnosis of PND leads to the detection of a tumour in 80% of cases but treatments are poor. More research is needed to improve the prognosis of these difficult conditions.

Audit of 2 week urgent referrals for CNS/cranial tumours
AJ Wills, DSNA Pengiran Tengah Queens Medical Centre, Nottingham, Derbyshire Royal Infirmary, Derby, UK

Objective: To audit patients referred from general practitioners (GPs) with the 2 week system of closer communication between primary and secondary care would be a more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be a more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources. Perhaps a system of closer communication between primary and secondary care would be more effective use of time and resources.

Methods: GP referral letters were reviewed and compared with Department of Health (DoH) guidelines. Patient case notes were examined to determine the actual neurological diagnosis. A list of patients with cancer that were not identified by the system was compiled separately.

Results: Forty five patients were referred during a 9 month period, with 41 sets of case notes available. 13 referrals did not follow the DoH guidelines. Only four patients actually had CNS tumours (two astrocytomas, one primary cerebral metastasis, two gliomas, one meningioma). The remaining patients were referred to determine the actual neurological diagnosis. A list of patients with cancer that were not identified by the system was compiled separately.

Conclusion: It can be concluded that intranasal sumatriptan 20 mg is effective in the acute treatment of cluster headache when compared with placebo.

A double blind placebo controlled study of intranasal sumatriptan in acute cluster headache
PJ Goodeby, JA van Vliet, A Bahra, V Martin, SK Aurora, NT Mathew, MD Maddison, I Hart, K Mills, A Vincent, J Newsom-Davis, Neurology, Queen Square, London, UK

Cluster headache attacks reach a peak of pain intensity quickly, and are short lasting; therefore acute treatments must be rapidly acting. This study sought to determine if intranasal sumatriptan is an effective treatment of acute cluster headache.

Patients with episodic and chronic cluster headache, by International Headache Society criteria, were recruited and after explanation and obtaining informed consent, were randomised to a double blind placebo controlled two period crossover study. Patients were instructed to treat two acute cluster headaches of at least moderate pain severity, with at least a 12 hour break, using intranasal sumatriptan 20 mg or matching placebo. Patients recorded the time of onset of the attack, time of treatment, headache severity on a five point scale (0, no pain; 1, mild; 2, moderate; 3, severe; 4, very severe) at 5, 10, 15, 20 and 30 minutes after treatment. The primary end point was the headache severity defined as: very severe, severe or moderate pain becomes mild or nil, at 30 minutes. Secondary measures included pain free rates, treatment of associated symptoms and adverse events.

The primary end point was analysed using a mixed effects approach with MLwiN (www.ioe.ac.uk/mulitlevel) with p<0.05 as the level of significance for testing.

In total, 118 patients were recruited; 97 males and 21 females. Of these 86 provided efficacy data on attack one and 80 on attack two. Twenty five provided no efficacy data because their bout ended, seven were lost to follow up. Six cycled out of a cluster period between attack one and attack two. Modelling the treatment outcome as a binomial where response was determined by treatment, using the patient as the level 2 variable, and considering period effect, sex, site and other headache type, the other variables of interest included were: intranasal sumatriptan 20 mg was superior to placebo at 30 minutes on the headache response end point (p=0.01). For the first attack the placebo response rate was 11/37 (30%) and the sumatriptan 20 mg rate 29/50 (58%); for the second attack the rates were 15/45 (33%) compared to 36/60 (60%), respectively. There were no serious adverse events.

It can be concluded that intranasal sumatriptan 20 mg is effective in the acute treatment of cluster headache when compared with placebo.

A study of the acquired neuromyotonia phenotype of peripheral nerve hyperexcitability
P Maddison, I Hart, K Mills, A Vincent, J Newsom-Davis Radcliffe Infirmary, Oxford, University of Liverpool, Liverpool, The Walton Centre for Neurology and Neurosurgery, Liverpool; King’s College Hospital, London, UK

Acquired neuromyotonia (NMT) and cramp fasciculation syndrome (CFS) represent the severe and mild manifestations of generalised peripheral nerve hyperexcitability (PNH) respectively. Anti-voltage gated potassium channel (VGKC) antibodies have been implicated in autoimmune PNH disease pathogenesis. We studied 44 consecutive...
patients with NMT seen in two centres to determine the significant clinical, immunological, and electrophysiological features of this phenotype.

The mean age at symptomatic onset was 47 years (range 9–74); over half (64%) were male. Associated autoimmune diseases and other autoantibodies were found in 22 (50%); an immunoprecipitation assay detected raised titres of anti-VGKC antibodies in 16 out of 40 (40%) patients.

All patients had at least two of the three principal symptoms of muscle twitching, cramps, and muscle stiffness. Other often observed features were increased sweating in 25 (57%) muscle weakness in 16 (34%), and pseudomyotonia in 16 (34%).

Spontaneous, abnormal activity detected electromyographically comprised characteristic doublet, triplet, and multiplet motor unit (or partial motor unit) discharges, in addition to fasciculations and fibrillations. The most common abnormality on EMG, in all but three patients, were doublet discharges, with intraburst frequencies often exceeding 100 Hz (mean 168 Hz, range 40–280 Hz). Multiplets (runs of >3 discharges) were found in the remainder.

NMT is a phenotype of autoimmune PNH that seems to be distinct from CFS only in terms of the severity of symptoms, signs and EMG features observed, with both sharing immunological associations.

HETEROPLASTIC MITOCHONDRIAL DNA MUTATIONS: DETERMINING THE MECHANISM OF INHERITANCE
PF Chinnery, DC Samuels, DT Brown, DM Turnbull

Mitochondrial DNA (mtDNA) mutations cause progressive disabling neurological disease in at least 1 in 8000 of the UK population. Many mtDNA mutations are heteroplasmic, with a mixture of mutant and wild type mtDNA present within the same individual. A higher percentage level of mutant mtDNA (mutation load) is associated with more severe disease. Females harbouring heteroplasmatic mitochondrial DNA point mutations transmit a variable amount of mutant mtDNA to their offspring, making it extremely difficult to predict the outcome of pregnancy. In mice, the variation in mutation load between offspring is determined by random drift and a restriction in the number of mtDNA molecules early in embryogenesis (the mitochondrial "genetic bottleneck"). It remains to be established whether the same mechanism operates in humans. We studied the mechanism of inheritance of the most common transmitted heteroplasmic mtDNA defect: the A3243G ("MELAS") mutation. By measuring mutation load in 82 individual human primary oocytes we show that random drift is the principal mechanism behind the transmission of heteroplasm in human pedigrees with mtDNA disease. We also provide the first direct evidence of a mitochondrial genetic bottleneck in humans, enabling the first direct measurement of bottleneck size. These findings will have important implications for genetic counselling.

RESULTS OF A UNITED KINGDOM-WIDE CLINICAL AND MOLECULAR GENETIC STUDY OF MYOTONIA CONGENITA
NP Davies, LH Eunson, MG Hanna

Myotonia congenita is the commonest skeletal muscle channelopathy and is caused by mutations in the skeletal muscle chloride channel gene (CLCN1) on chromosome 7q35. The aims of the study were to look at genotype-phenotype correlation in this group, and to analyse the clinical and genetic features of myotonia congenita in the UK.

In order to check that no mutations had been formed SSCP analysis of all 23 exons of the CLCN1 gene in index cases from 42 families was performed. In all 23 exons doublet discharges, with intraburst frequencies often exceeding 100 Hz (mean 168 Hz, range 40–280 Hz). Multiplets (runs of >3 discharges) were found in the remainder.

NMT is a phenotype of autoimmune PNH that seems to be distinct from CFS only in terms of the severity of symptoms, signs and EMG features observed, with both sharing immunological associations.

A CYSTEINE RESIDUE CLOSE TO THE C-TERMINUS OF THE E SUBUNIT IS ESSENTIAL FOR SURFACE ACETYLCHOLINE RECEPTOR (ACHR) EXPRESSION: IMPLICATIONS FOR ACHR DEFICIENCY SYNDROME

ACHR deficiency is a recessively inherited form of congenital myasthenia in which adult ACHR (α;β) are severely reduced or absent at the neuromuscular junction. We have identified two new mutations at the extreme C terminus of the subunit in ACHR deficiency patients and have investigated the mechanism by which such distal mutations may affect expression.

Naturally occurring and artificial mutations were introduced into the ε subunit cDNA by in vitro mutagenesis. Wild type or mutant ε subunit cDNAs were transfected into HEK 293 cells with cDNAs for the β1, β2, and ε ACHR subunits. Surface expression of ACHR was quantified by radioimmunoprecipitation of 

125I-α-bungarotoxin labelled receptor using an ε specific antiserum. To determine the subcellular location of ε subunits, fluorescently labelled ε subunits were visualised using confocal microscopy. Lack of a cytosine residue (Cys 470), four amino acids from the C terminus of the ε subunit, effectively abolished surface expression of the intact receptor in HEK 293 cells. Preliminary confocal data suggests that the mutant ε subunits accumulate in the endoplasmic reticulum.

These novel results reveal a previously unidentified signal sequence involved in the assembly and/or transport of ACHR. Additionally, these results provide a mechanism by which frameshift or non-sense mutations in the C terminus region of the ε subunit lead to ACHR deficiency.

SERUM AUTOANTIBODIES TO OLIGODENDROCYTE AND NEURONAL CELL SURFACE ANTIGENS IN MULTIPLE SCLEROSIS PATIENTS
O Lily, J Palace, A Vincent

Recent evidence suggests that autoantibodies may be important in the pathogenesis of multiple sclerosis (MS), but the antigens to which they bind remain unknown. We incubated cultured human CNS cell lines with sera from MS patients and other inflammatory neurological disease controls (OIND). Surface binding was measured using flow cytometry and considered positive if greater than the mean +2 standard deviations of 12 healthy controls.

Antibody binding to two control cell lines was similar using MS or non-MS sera. By contrast, 70% (16/23) of secondary progressive (SP) MS sera demonstrated positive binding to SK-N-SH neuroblastoma cells compared with 29% (10/35) of relapsing remitting (RR) patients (p<0.0001) and 0/16 OIND controls. Also, 40% of MS sera showed positive binding to HOG oligodendrogliaoma cells with no significant difference between RR and SP patients. Differentiation of the oligodendroglial cells resulted in the loss of positive antibody binding seen using MS sera, suggesting that the relevant antigens are present on oligodendrocyte precursor cells rather than mature oligodendrocytes.

Sera with antineuronal antibodies generally also had anti-oligodendroglial antibodies, although antibodies against one cell line could not be absorbed by the other. These results support reports of heterogeneity in MS pathology, with autoantibody mediated processes important in a subgroup of patients.
**Conclusion:** In the UK only a small proportion of MND patients are referred for NIV, with marked heterogeneity of practice. This may reflect concerns that NIV may prolong suffering, and emphasises the need for a controlled trial assessing its impact on quality of life.

**AN MRI STUDY OF COMPLICATED EARLY CHILDHOOD CONVULSION**

RA Grunewald, T Farrow, CDC Ritchie, P Vaughan, J Mundy Royal Hallamshire Hospital, Rye Gate Children's Centre, Sheffield, UK

**Objectives:** About half of adults with hippocampal sclerosis have a history of complicated early childhood convulsion (CECC). The objective of the study was to assess the extent of structural brain abnormality present soon after a first CECC.

**Methods:** Children under the age of 5 years underwent brain MRI within 14 days of their first complicated febrile or non-febrile ECC. None had previously experienced an epileptic seizure. Hippocampal volumes and T2 relaxation times were measured. The control group consisted of children undergoing brain MRI for reasons other than epilepsy.

**Results:** Nine of the 17 experimental subjects had significant hippocampal volume asymmetry (3 standard deviations from the mean of the control group), although in only three of these was the asymmetry apparent on visual inspection of the MRI. Three experimental subjects had extrahippocampal neuropathology. None of the 10 control subjects had significant hippocampal volume asymmetry (p<0.001). T2 relaxometry did not suggest that oedema contributed to the asymmetry.

**Conclusions:** There is a high prevalence of structural brain abnormalities and hippocampal asymmetry in children within 2 weeks of the first complicated early childhood convulsion, unrelated to oedema. This suggests that in some cases the complicated EEC is the result of pre-existing brain abnormalities.

**A DOUBLE BLIND SINGLE DOSE CROSS OVER STUDY OF THE EFFECTS OF PRAMIPEXOLE, PERGOLIDE, AND PLACEBO ON TREMOR AND UPDRS (III) IN PARKINSON'S DISEASE**

P Navan, LJ Findley, R Pearce, J Jeffs, PG Bain Imperial College School of Medicine, London; Havering Hospital NHS Trust, Essex; Hammersmith Hospital, London UK

**Design:** 10 tremulous PD patients were pretreated with domperidone and taken off anti-parkinsonian medication. Each patient received single doses of 0.5 mg pramipexole, 0.5 mg pergolide, and placebo, but in random order. Rest tremor and UPDRS (part III) were measured at baseline and 30 minute intervals for 4 hours. The results were examined using ANOVA by area under the curve (AUC) and peak effect (Bonferroni corrected; significance: 5% level).

**Results:** (1) AUC analysis showed a significant difference between the three treatments on rest tremor (p=0.045) (pramipexole differed from placebo at 10% level) but not UPDRS (III). (2) Peak effect analysis demonstrated a difference in treatment effects on UPDRS (III) (p=0.04) (pergolide better than placebo, p=0.048) and rest tremor (p=0.005) (no significant difference between active drugs, both significantly better than placebo). Pergolide caused a significantly greater incidence of nausea (p=0.005) and vomiting (p=0.014) than pramipexole. No significant differences between the 3 treatments on incidence of drowsiness or sleep (McNemar’s test).

**Conclusion:** 0.5 mg pramipexole or pergolide can improve UPDRS (III) and rest tremor in PD. Pergolide had the strongest effect on peak UPDRS (III), whereas pramipexole had the greatest effect on rest tremor (AUC). Pramipexole is significantly better tolerated than pergolide.

**PALLIDAL STIMULATION FOR UNUSUAL MOVEMENT DISORDERS: 1 YEAR OUTCOME OF FOUR PATIENTS**

SG Parkin, RP Gregory, TZ Aziz The Radcliffe Infirmary, Oxford, UK

Pallidal stimulation has become an established mode of treatment in Parkinson's disease when medical therapy has failed. There is historical evidence to suggest that surgery to the basal ganglia can benefit a wide range of movement disorders. The relative safety of deep brain stimulation has allowed us to re-explore its benefit to other, less common conditions. We report the outcome of pallidal stimulation in a consecutive series of patients with idiopathic generalised dystonia, familial myoclonic dystonia, idiopathic progressive chorea, and tardive dyskinesia with cataplexy. All had failed an exhaustive trial of medical therapy. Highly significant benefits were seen in various movement disorder rating scales and quality of life measures. These were maintained at 1 year. The obvious clinical benefits, rarity and heterogeneity of conditions of this type, precludes the use of randomised clinical trials. Detailed follow up of individual patients is important to provide evidence for the application of pallidal DBS in such conditions.

**CRITERIA FOR INITIATING NON-INVASIVE VENTILATION IN MOTOR NEURON DISEASE**

SC Bourke, PJ Shaw, R Bullock, GJ Gibson University of Newcastle, Newcastle; University of Sheffield, Sheffield; Newcastle General Hospital, Newcastle, UK

**Objective:** We studied the criteria which best predict benefit from quality of life (QOL) and compliance with non-invasive ventilation (NIV) in motor neuron disease (MND).

**Methods:** 23 subjects were recruited. QOL (SF-36, chronic respiratory disease questionnaire, sleep apnoea quality of life index (SAQII), vital capacity, and maximum respiratory pressures were assessed two monthly and polysomnography four monthly. Subjects were offered a trial of NIV with any of: (1) orthopnoea, (2) unrefreshing sleep or day-time sleepiness, (3) Daytime PaCO2 >45 mm Hg, (4) So2 <90% for ≥5% of the night, (5) apnoea hypopnoea index >10.

**Results:** The improvement in the SAQII symptoms domain (most responsive index) after 1 month NIV (effect size: mean difference/SD), and compliance are shown. Results for subjects in whom only one criterion for NIV was met are in brackets.

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<td>AH1 &gt;10</td>
<td>7 [4] 3 [1]</td>
<td>4.2 [2.0]</td>
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<table>
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<tr>
<th>Orthopnoea index</th>
<th>SaO2</th>
<th>PaCO2</th>
<th>AH1 &gt;10</th>
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</thead>
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<tr>
<td>Symptoms Orthopnoea 1</td>
<td>SaO2</td>
<td>PaCO2</td>
<td>AH1 &gt;10</td>
</tr>
<tr>
<td>Orthopnoea 2</td>
<td>SaO2</td>
<td>PaCO2</td>
<td>AH1 &gt;10</td>
</tr>
<tr>
<td>Orthopnoea 3</td>
<td>SaO2</td>
<td>PaCO2</td>
<td>AH1 &gt;10</td>
</tr>
</tbody>
</table>

10/16 subjects starting NIV elected to continue. Among these, orthopnoea was the commonest indication (9/10). Moderate/severe bulbar weakness was associated with lower compliance (4.22 v 8.47 h/day) and less improvement in QOL (effect size: 1.85 v 3.28).

**Conclusions—Orthopnoea was the best predictor of compliance with NIV in higher symptom MND. Moderate/severe bulbar involvement was associated with poorer (although still clinically useful) compliance and response.

**RESPONSIVENESS OF QUALITY OF LIFE INSTRUMENTS AND DOMAINS TO NON-INVASIVE VENTILATION IN MOTOR NEURON DISEASE**

SC Bourke, PJ Shaw, R Bullock, GJ Gibson University of Newcastle, Newcastle; University of Sheffield, Sheffield; Newcastle General Hospital, Newcastle, UK

**Background:** As a prelude to a randomised controlled trial of non-invasive ventilation (NIV) in motor neuron disease (MND), we studied which quality of life (QOL) instruments are most responsive. We report the results of an efficacy analysis of the impact of NIV on QOL in 10 subjects with MND.

**Methods:** QOL and functional status were assessed at 0, 1, and 3 months after initiation of NIV, then 2 monthly intervals using SF-36, general wellbeing schedule (GWbS), chronic respiratory disease questionnaire (CRDQ), sleep apnoea quality of life index (SAQII), Epworth sleepiness scale (ESS), and ALS functional rating scale (ALS-FRS). The maximum improvement in each index was assessed by effect size (mean difference/SD).

**Results:** Effect sizes for all responsive instruments are shown. The GWbS and SF-36 energy vitality and health change showed non-significant improvements. The largest improvements were in domains assessing sleep-related symptoms, which were maintained for a mean of 320 days (Kaplan-Meier). As expected the SF-36 physical function domain and ALSFRS declined with disease progression.

**Conclusion:** NIV improves QOL in selected patients with MND. QOL domains assessing sleep-related symptoms were most responsive, and the improvement was maintained at or above baseline for ≥320 days despite disease progression.
THE EFFECT OF CANNABIS BASED MEDICINAL EXTRACT ON LOWER URINARY TRACT DYSFUNCTION IN ADVANCED MULTIPLE SCLEROSIS: PRELIMINARY RESULTS

CM Brady, R DasGupta, CJ Wiseman, CM Dalton, KJ Berksley, CJ Fowler
National Hospital for Neurology and Neurosurgery, Queen Square, London, UK

Aims: The primary aim of this open label pilot study is to evaluate the safety, tolerability, and efficacy of two preparations of sublingual cannabis based medicinal extract (CBME) in patients with advanced multiple sclerosis (MS) and refractory lower urinary tract symptoms (LUTS).

Methods: Patients with advanced MS (Kurtzke score >6.5) and refractory LUTS (detrusor hyperreflexia demonstrated by cystometry) are eligible for recruitment into this ongoing study. Frequency-volume incontinence charts and pad testing are completed during the run in (3 weeks) and treatment (16 weeks) periods. For the first 8 weeks of treatment, patients receive CBME containing equal amounts of cannabidiol (CBD) and tetrahydrocannabinol (THC) whereas a THC only preparation is prescribed for weeks 9–16. Patients self titrate the CBME until urinary symptoms are relieved or unacceptable side effects occur. To assess the acute effects of CBME, cystometry is performed with CBME in the department.

Results: We report results of the first eight patients (2M:6F, age 31–64). A paired t test was used.

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Domains</th>
<th>Effect size</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>ESS</td>
<td>0.85</td>
<td>0.019</td>
<td></td>
</tr>
<tr>
<td>SF 36</td>
<td>0.93</td>
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<tr>
<td>Mental health</td>
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<tr>
<td>CRDQ</td>
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<td>0.018–0.044</td>
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<tr>
<td>SAQI</td>
<td>0.96</td>
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<tr>
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<td></td>
</tr>
<tr>
<td>Score</td>
<td>1.11</td>
<td>0.030</td>
<td></td>
</tr>
</tbody>
</table>

The average cystometric capacity increased from 265 ml at baseline to 439 ml at 8 weeks, and to 353 ml at 16 weeks after dosing with CBME in the department.

Conclusions: These preliminary results indicate that CBME may prove to be an effective additional treatment for refractory urinary symptoms in a selected group of patients with advanced MS.

INTRINSIC AND MISSENSE MUTATIONS WITHIN THE LAMP-2 GENE IN DANON DISEASE (X LINKED VACUOLAR CARDIOMYOPATHY AND MYOPATHY)

NP Davies, C Beeley, PM Elliott, J Holton, B Lake, DN London, P Lee, H Mundy, B Winchester, MG Hanna Institute of Neurology, Queen Square, London; Institute of Child Health, London, UK

Danon disease is an X linked disorder characterised by cardiomyopathy, vacuolar myopathy, and variable mental retardation. It is now known to be due to mutations within a gene encoding the lysosomal associated membrane protein LAMP-2 on Xq24.1.

We report three families with Danon disease. Affected males in each family presented with profound cardiac symptoms but had mild proximal myopathy. Mental retardation was apparent in only three out of five affected males. Proximal myopathy and mental retardation were not apparent in affected females. Age at death was 15–24 years for men and 29–40 years for women. Muscle biopsies revealed the characteristic vacuolar myopathy. We also showed intense deposition of the complement membrane attack complex (MAC) on vacuolated fibres. This immunoreactivity has been previously reported to be specific for another vacuolar myopathy (X linked myopathy with excess autophagy, XMEA). Genetic analysis revealed three new mutations in the LAMP-2 gene. We report the first missense mutation in LAMP-2 (W321R). In addition, two new intronic mutations were detected.

In summary, we confirm the clinical and pathological characteristics of Danon disease and suggest that deposition of MAC on vacuolated fibres may be a feature. We identified three previously unreported mutations in the LAMP-2 gene. The W321R mutation represents a new mutational mechanism, as all other mutations disrupt splicing or lead to premature truncation of the protein. In view of the sparse skeletal muscle signs in this disorder, we would recommend a skeletal muscle biopsy in any young patient with an apparently isolated cardiomyopathy.

GLUTEN ATAXIA AND GLUTEN NEUROPATHY: THE EFFECT OF A GLUTEN-FREE DIET

M Hadjivassiliou, RA Grunewald, AK Chattopadhyay, RH Kandler, JA Jarrett, GAB Davies-Jones The Royal Hallamshire Hospital, Sheffield, UK

We studied the effect of gluten free diet on patients with ataxia and/or peripheral neuropathy associated with gluten sensitivity.

Sixteen patients with gluten ataxia (sporadic ataxia with circulating anti-gliadin antibodies) were assessed before and after 12 months treatment with a gluten free diet. Subjective assessment of clinical state was scored on an analogue scale improved from mean score (SD) at baseline of 7.7 (1.5) to 4.9 (0.7). When compared with a heterogenous control group of eight ataxic patients without gluten sensitivity or who declined treatment with a gluten free diet, the improvement was significant (repeated measures ANOVA p=0.002). Improvement was also seen in time standing with feet together without corrective movement (baseline 19.1 seconds (22.7), 12 months 31.2 seconds (26.4), p=0.05 Student’s t test). There was a trend towards improvement in computerised measures of latency and accuracy of arm coordination.

Sural sensory action potentials were measured in a group of 13 patients with gluten neuropathy before and 12 months after starting a gluten free diet. Baseline and 12 month repeat measurements were also made in a control group of five subjects with gluten neuropathy who declined the diet. There was some increase in amplitudes in the group treated with diet: 1.4 µV (1.3) v 2.3 µV (2.1) but none in the control group: 1.6 µV v 1.1 µV (1). The change in amplitude between the two groups was not significant (p=0.067 Mann-Whitney).

These preliminary data suggest that a gluten free diet may be of benefit in some patients with neurological manifestations of gluten sensitivity.

HUMAN COMPLICATION OF THE FOOT AND MOUTH CRISIS

KAC Harkness, TG Staunton Norfolk and Norwich Hospital, Norwich, UK

We report an unusual complication of the foot and mouth crisis. After our patient, a marksman, had examined a culled bull for signs of life, the animal’s head moved, trappimg his feet and causing him to lose balance. He fell backwards onto the horn of another dead animal, sustaining a penetrating injury to the occiput. He did not lose consciousness but complained of loss of vision in the right hemifield, 30 minutes after the initial injury. In the neurosurgical unit CT demonstrated a left comminuted depressed occipital skull fracture with some impingement of the left occipital-parietal lobe. The patient was managed conservatively with intravenous antibiotics and discharged 1 week later.

Two weeks later the patient developed occipitofrontal headache with photophobia and presented to our neurology unit. He was ataxic, no meningism, no papilloedema, a resolving right field defect was noted. MRI and MRA images were performed. The vertebral arteries were normalised from mean score (SD) at baseline of 0.49 (0.17) to 0.7 (0.26). When compared with a heterogenous control group: 1.6 µV v 1.1 µV (1.2). The change in amplitude between the two groups was not significant (p=0.067 Mann-Whitney).

A UNIQUE MODEL OF LISSENCEPHALY

L Kinton, BA Barth, C Howart, JS Duncan, M Gardiner, NW Wood, S Wilson Institute of Neurology, Queen Square, London; University College, London, UK

Disorders of neuronal migration are important causes of drug resistant epilepsy. Mutations in two genes LIS1 and DCX have been shown to cause one such condition, agyria/pachygyria/band heterotopia complex.
The zebrafish, Danio rerio, is used as a model organism for studying development of vertebrate systems including the CNS. Advantages of this model organism include rapid external development, transparency, and most recently morpholino antisense oligonucleotide technology for creating targeted gene knockdown. Conservation of genetic pathways across vertebrates make it a valid model organism.

The zebrafish orthologue of the LIS1 gene has been cloned and it shows a high degree of sequence conservation with human LIS1 (81% nucleotide, 93% amino acid identity). The expression pattern in the developing zebrafish is analogous to mammalian expression indicating high levels in the developing CNS.

The novel technique of morpholino antisense oligonucleotide injection has been performed against the lis1 gene. Graded reduction in lis1 protein translation and hence activity in the zebrafish embryo results (knockdown), providing an in vivo model of lissencephaly. These embryos show abnormalities in the developing brain, most notably the telencephalon and hindbrain. This is the first time this technique has been used to model a neurological disorder important in epilepsy pathogenesis.

USE OF INTERNET MEDICAL WEBSITES AND OF NHS DIRECT OUTPATIENTS BEFORE CONSULTATION

A J Larner The Walton Centre for Neurology and Neurosurgery, Liverpool, UK

Objectives/methods: Two sources of medical information have recently become available to the general public: medical websites on the Internet, and the NHS Direct telephone helpline. This study measured the use of these resources before consultation by consecutive new referrals to general neurology outpatient clinics.

Setting: Two district general hospitals in north west England. Results: Of 198 patients seen over 3 months, 66 (33%) had home access to the Internet. Of these 66, 15 (22% = 7.6% of all patients) had accessed websites containing medical information; only one patient volunteered this information.

Of these 15, nine (60%) accessed information which was adjudged inappropriate in the light of the final diagnosis. This was due to: incorrect self diagnosis, based on self directed searches (5/9); incorrect diagnosis by another doctor (2/9); other, pre-existing correct diagnoses not relevant to current symptomatology (2/9).

In addition to accessing inappropriate information found that searches were time consuming and generally provided nothing beyond what they already knew. Only one patient found website information “very helpful”, but this was inappropriate to the final diagnosis.

Only 4/198 patients (2%) had telephoned NHS Direct before consultation; none volunteered this information. In only one did the call relate to neurological symptoms.

Conclusions: In this study around a quarter of patients with home access to the Internet used medical websites before neurological consultation; use of NHS Direct was negligible. Information accessed from Internet sites was adjudged inappropriate in 60% of cases. Internet searches, although seldom volunteered, may influence patients’ health beliefs and expectations, and hence the subsequent consultation. Such considerations may become increasingly relevant as personal Internet access expands.

AN IMPROVED METHOD OF QUANTIFYING ACTIVITY LEVELS IN NEUROLOGICAL PATIENTS

C Lawthom, RMW van Deerzen, CM Wiles University Hospital Wales, UK

Current assessments of mobility rely on brief observations of walking and questionnaire based scales that are surrogate measures of everyday activity. Accelerometry techniques now allow unobtrusive activity measurement over many days. Activity levels in two patients and two matched healthy controls were measured using the Stepwatch Activity Monitor, a device worn on the ankle to record cadence over 3 to 8 days. These data provide further support to our contention that gluten ataxia is immune mediated.

Gluten sensitivity is a state of heightened immunological responsiveness triggered by the ingestion of gluten in genetically susceptible individuals. Organ specific manifestations include enteropathy (coeliac disease) a dermatopathy (dermatitis herpetiformis) and CNS involvement (gluten ataxia). Gluten ataxia accounts for up to a third of idiopathic sporadic ataxias. It is defined by the presence of antigliadin antibodies in otherwise unexplained sporadic idiopathic ataxia. The mechanism of neurological damage remains obscure.

Postmortem brain tissue of two patients with gluten ataxia showed T cell infiltration and upregulation of chemokine expression. CSF from 10 patients with gluten ataxia showed upregulation of the chemokine IP-10 when compared with controls. Serum from patients with gluten ataxia stained human cerebellar Purkinje cells at dilutions up to 1:800 with strong staining seen up to 1:400. At 1:400 staining was not seen in the control groups (five patients with other causes of ataxia, five normal controls, five patients with coeliac disease). Commercial antigliadin antibodies stained human Purkinje cells in a similar manner. Persistent staining after adsorption with crude gliadin was only observed when using sera from patients with gluten ataxia. These data provide further support to our contention that gluten ataxia is immune mediated.

**Activity index**

<table>
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<tr>
<th>Activity level</th>
<th>Patient 1</th>
<th>Patient 2</th>
<th>Control 1</th>
<th>Control 2</th>
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</thead>
<tbody>
<tr>
<td>Mean activity (h/day)</td>
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<td>0.87</td>
<td>5.80</td>
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<tr>
<td>Range of activity (h/day)</td>
<td>1.3–4.1</td>
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<td>3.6–7.1</td>
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<td>33.8</td>
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<td>Right steps/d</td>
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<td>4245</td>
<td>3988</td>
</tr>
<tr>
<td>Mean distance/d (m)</td>
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<td>857–2820</td>
<td>92–2415</td>
<td>4152–2604</td>
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</tr>
</tbody>
</table>

Gluten ataxia is immune mediated

D Mahad, K Oates, C Williams, R Grunewald, SJ Howell, MN Woodroofe, M Hajjavioli Royal Hallamshire Hospital, Sheffield, UK

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EVIDENCE FOR DEMENTIA IN SPAST-HEREDITARY SPASTIC PARAPARESIS

P McMonagle, P Byrne, S Webb, N Parfrey, M Hutchinson St Vincent’s University Hospital, Dublin, Ireland

Objective: Cognitive impairment is recognised in SPAST-hereditary spastic paraparesis (HSP). It is unclear if this represents a dementia. Our aim was to examine SPAST-HSP pedigrees longitudinally for evidence of progressive cognitive decline.

Methods: Five families with SPAST-HSP were identified, 4 with detectable mutations and one based on linkage. Family members were tested for cognitive impairment at first visit and again after a minimum of 1 year. The Cambridge Cognitive Assessment (CAMCOG) has a maximum score of 107, scores <80/107 indicate cognitive impairment. Non-parametric tests of significance were used.

Results: 24 patients over age 20 from five families had paraparesis. Repeat CAMCOG were available for 22. The age range was 2.9 years. The age range was 54.5 years. CAMCOG declined over that time from 86.6 to 82.4/107 (p=0.018). In those with cognitive impairment at baseline (n=9) CAMCOG declined further from 74 to 64.4/107 (p=0.012). The average age for this group was 66.4 years. CAMCOG for controls (n=5,9 years, n=9) showed little change (91.8 to 90.2/107, NS).

Conclusions: We have demonstrated longitudinal cognitive decline in those with SPAST-HSP and further decline in those with established cognitive impairment, suggesting that this is a dementing process.

TWO CLINICAL PRESENTATIONS OF ADULT ONSET RASMUSSEN’S SYNDROME SHARE COMMON IMMUNOLOGICAL AND PATHOLOGICAL FEATURES

RS Nicholas, AC Scott, IK Hart The Walton Centre for Neurology and Neurosurgery, Liverpool, UK

Childhood Rasmussen’s syndrome (RS) is a focal cortical inflammation causing epilepsy and progressive neurological deficits. We studied six patients to define the clinical and pathological features of adult RS and performed comprehensive screens for anti-CNS and other autoantibodies. All patients presented with epilepsy (average age 29 years). There were two patterns of clinical onset and progression—rapid neurological deficit <12 months after seizure onset; four patients) and slow (neurological deficit >42 months after seizure onset). The rapid form presented with focal increased signal on T2 MRI that later developed into ipsilateral hemisphere atrophy. Histology was similar to childhood RS. In low form, MRI brain showed focal cortical atrophy. On histology, there was moderate perivascular T cell infiltration, microgliosis, and reactive astrocytosis. In both forms, there was bilateral focal cortical hypoperfusion on SPECT scans. All patients had multiple serum and CSF immune abnormalities and had similar HLA haplotypes. We propose that adult RS has two clinical phenotypes—rapid and slow—that share similar brain, CSF, and serum autoimmune abnormalities. Our findings may help the early diagnosis of adult RS and identify candidates for immunomodulatory therapy before they develop extensive, irreversible neuron loss.

EPILEPSY SURGERY IN PATIENTS WITH ADDITIONAL PSYCHOGENIC SEIZURES

M Reuber, G Fernandez, M Kurthen, J Schramm, CE Elger University of Bonn, Bonn, Germany

Objective: To assess whether surgery to reduce or control epileptic seizures is safe and effective in patients known to have additional psychogenic seizures.

Methods: We reviewed our computerised database of 1342 patients evaluated for epilepsy surgery and identified 13 patients with both epileptic and psychogenic seizures on whom postoperative data were available. Outcome data were gathered from patient records and completed by telephone interviews. Mean postoperative follow up was 56 months. (HSP). It is unclear if this represents a dementia. Our aim was to examine SPAST-HSP pedigrees longitudinally for evidence of progressive cognitive decline.

Results: Epilepsy surgery led to relevant improvements in 11 of 13 patients. Seven patients became completely seizure free, two patients become free of epileptic seizures but continued to have infrequent psychogenic seizures, one patient reported an over 80% improvement of epileptic seizure frequency and an abolishment of psychogenic attacks, in one patient non-disabling epileptic seizures persisted at lower frequency but psychogenic seizures stopped. In two of 11 patients, epilepsy surgery failed to produce notable improvements. Although epileptic seizures improved, their psychogenic seizure disorder remained highly disabling. In both of these patients pathological illness behaviour had been noted preoperatively.

Conclusion: A diagnosis of additional psychogenic seizures should not be considered an absolute contraindication to epilepsy surgery although patients should undergo careful preoperative psychiatric evaluation.

EXPLORING THE CONNECTIVITY OF BAND HETEROTOPIA USING DIFFUSION TENSOR IMAGING

FJ Rugger-Gunn, M Guye, SH Eriksson, G Parker, GJ Barker, C Wheeler-Kingshott, JS Duncan University College London, London, UK

Introduction: Diffusion tensor imaging (DTI) is an MRI method that evaluates the diffusion of water molecules. DTI tractography allows mapping of white matter tracts. Band heterotopia (BHT) is a malformation of cortical development in which there are bands of ectopic grey matter in the subcortical white matter. Previous functional imaging has shown activations in BHT. We have used tractography in five patients with BHT to determine whether connecting tracts could be identified in vivo.

Methods: Five patients with epilepsy and bilateral BHT were scanned with DTI. Anisotropy (diffusion directionality) maps were calculated and evaluated using fast marching tractography. This produced maps of the likelihood of connection which were used to determine the pathways of white matter tracts through the areas of BHT, and compared with control data.

Results: White matter tracts were identified passing from and through the BHT to the cortex in each patient.

Conclusions: Using tractography we have shown that white matter tracts traverse and connect BHT to overlying cortex. This shows the structural basis of the functional connectivity implied by activation studies and the absence of focal neurological deficit. This result demonstrates the potential of the method to indicate structural connectivity in vivo.

THE PROGNOSTIC VALUE OF INTRATHecal IMMune RESPONSE AGAINST AXONAL COMPONENTS IN PATIENTS WITH MULTIPLE SCLEROSIS

OA Seidi, YK Semera, NA Gregson, MK Shariel Guy’s Hospital, London, UK

Background: We have recently reported a significant correlation between clinical disability and intrathecal release of antibodies against axonal cytoskeleton in patients with primary and secondary-progressive MS.

Objective: To determine the prognostic value of intrathecal antibodies to axonal proteins for future development of disability in MS patients.

Methods: In a prospective study, we measured intrathecal levels of antibodies against actin, tubulin and neurofilament light-subunit from 51 patients with relapsing-remitting, secondary, or primary-progressive MS. Patients were clinically monitored for 2 years after this lumbar puncture (LP). We included 52 patients with other neurological disorders and eight healthy subjects, as controls. We also measured antibodies against myelin basic protein (MBP) as a control antigen.

Results: Intrathecal levels of antibodies against all axonal proteins, but not anti-MBP, correlated with the expanded disability status scale (EDSS) score at the time of LP. Moreover, MS patients with high intrathecal levels of antibodies against axonal proteins accumulated more neurological disability at the end of the 2 year period compared with patients with low levels. By contrast, antibodies against MBP did not correlate with EDSS score at the end of the study.

Conclusion: The determination of intrathecal antibodies against axonal proteins may be clinically useful in MS patients.

MAPPING THE DISEASE LOCUS IN A LARGE KINDRED WITH GENERALISED EPILEPSY

A Siddiqui, M Davis, PH Dixon, M Johnson, M Koepp, SD Sherov, JWA Sander, RM Gardiner, JS Duncan, NW Wood Institute of Neurology, Queen Square, London; University College London, UK

A large British family with epilepsy underwent careful clinical evaluation using a structured seizure questionnaire. There was similarity to the GEFS plus [generalised epilepsy and febrile seizures plus] phenotype in that affected individuals manifested a variety of childhood onset epilepsy phenotypes with or without preceding febrile seizures. A simulation analysis estimated a maximum lod score of 4.17 at theta=0 and 3.84 at theta=0.05. The known loci for GEFS plus on chromosome 1q and 2q were investigated initially and excluded with lod scores ranging from −2.01 to −infinity.

Genome wide linkage analysis was undertaken using a medium density linkage mapping set. This contained 400 fluorescent dye labelled microsatellite markers that defined a 10 cM resolution index map.
This work identified two main regions of linkage on chromosome 2 and 5. These areas were investigated by genotyping further markers and constructing haplotypes. Two point lod scores are 3.01 at theta=0 and 2.41 at theta=0 respectively, and multipoint lod scores of 3.01 and 2.85 respectively. Searches are being carried out for candidate genes using the Draft Human Genome Browser.

MAGNETISATION TRANSFER RATION AND N-ACETYL ASPARATE AS INDEPENDENT MARKERS OF MYELIN AND AXONAL INTEGRITY IN DERMELINATING LESIONS: A MAGNETIC RESONANCE IMAGING STUDY OF CENTRAL PONTINE MYELINOLYSIS

NC Silver, DG MacManus, CA Davie, DH Miller Institute of Neurology, University College London, UK

Magnetisation transfer ratio (MTR) measurement seems promising as a putative marker of demyelination in the CNS. Small MTR reductions may also occur as a result of axonal loss, oedema, or inflammation. The individual contribution of tissue components to MTR values in healthy tissue is unknown. Central pontine myelinolysis (CPM), a condition characterised by severe demyelination and occasional minimal axonal loss in the absence of inflammation or oedema, provides a useful in vivo model to assess the effect of myelin disruption on MTR.

Three patients with CPM were studied with magnetisation transfer imaging and proton magnetic resonance spectroscopy (MRS) on one or more occasions during their illness. Moderate to severe MTR reductions were observed in acute lesions and resolution of MTR paralleled clinical recovery. Marked MTR reductions were observed despite general preservation of N-acetyl aspartate (NAA; an in vivo MRS marker of axonal integrity). These observations support the hypotheses that (a) myelin forms the predominant contribution to MTR values in healthy white matter, (b) demyelination itself can result in severe MTR reduction, (c) profound MTR reduction may be seen in the absence of significant neuronal damage, (d) MTR and NAA provide relatively independent markers of myelin and axonal pathology, and (e) MTR and NAA may provide useful diagnostic and prognostic information in CPM.

A PROSPECTIVE STUDY OF EPILEPSY SURGERY IN THE UK

JK Solomon, A McEvoy, SD Lhatoo, ND Kitchen, SD Sharvon Institute of Neurology, Queen Square, London, UK

The treatment of epilepsy has been revolutionised by improvements in epilepsy neurosurgery. We wished to determine the number of consultant neurosurgeons performing epilepsy surgery in the United Kingdom and to evaluate prospectively the total number of epilepsy procedures performed.

All practising consultant adult and paediatric neurosurgeons in the United Kingdom were sent a postal questionnaire asking if they had performed epilepsy surgery in the last year. Each identified surgeon was then asked to record the type and number of surgical procedures they had performed in a questionnaire mailed to them at the end of every month for 6 consecutive months commencing February 2000.

Thirty two consultant neurosurgeons in current practice who performed epilepsy surgery were identified. Out of a total of 192 monthly questionnaires there was a response rate of 95%. A total of 289 surgical procedures for epilepsy were performed, suggesting an annual figure of 578 procedures.

Although there has been an almost twofold increase in curative epilepsy surgery in the last decade, this still falls far short of what is required from epidemiological estimates. The need for more surgical assessment units as well as trained epilepsy surgeons is emphasised.

DERMATITIS HERPETIFORMIS, COELIAC DISEASE, AND NEUROLOGICAL DYSFUNCTION

DSNA Pengiran Tengah, DJ Unsworth, L Fry, G Giovannoni, A Church, B Turner, GKT Holmes, AJ Wills Queens Medical Centre, Nottingham; University College London; Institute of Neurology, London; Derbyshire Royal Infirmary, Derby, UK

Objective: To investigate the prevalence of neurological abnormalities observed in acute lesions and resolution of MTR paralleled clinical recovery. Marked MTR reductions were observed despite general preservation of N-acetyl aspartate (NAA; an in vivo MRS marker of axonal integrity). These observations support the hypotheses that (a) myelin forms the predominant contribution to MTR values in healthy white matter, (b) demyelination itself can result in severe MTR reduction, (c) profound MTR reduction may be seen in the absence of significant neuronal damage, (d) MTR and NAA provide relatively independent markers of myelin and axonal pathology, and (e) MTR and NAA may provide useful diagnostic and prognostic information in CPM.

Conclusions: The DH cohort was assessed for anti neuronal antibodies in detail. The presence of a novel spinal antibody in over 50% of the patients. The lifetime prevalence of essential tremor and migraine in these two cohorts was unremarkable. The patient with chorea had been on long term phenytoin, which is a reported association.

Conclusion: No cases of gluten ataxia were detected. The neurological abnormalities found in these patients are likely to be either chance or spurious associations contrasting with previous research. However, the unexpected discovery of this novel spinal antibody in DH patients requires further investigation, particularly in the light of previous studies suggesting an increased prevalence of sensory ataxia in patients with established gluten sensitivity.

WHITE MATTER T1 RELAXATION TIMES IN MULTIPLE SCLEROSIS ACQUIRED USING THREE DIMENSIONAL MAGNETIC RESONANCE IMAGING

L Vaithianathan, CR Terch, PS Morgan, LD Blumhardt Queen’s Medical Centre, Nottingham, UK

T1 relaxation time (T1) mapping provides a quantitative method for assessing disease burden in multiple sclerosis (MS).

previous studies have been limited to sampling small regions of normal appearing white matter or discrete lesions.

We compare total white matter T1 values in controls and MS patients from different clinical subgroups using histogram analysis. Relations between T1, disability, and T2 lesion volume are explored.

29 patients with MS (11 relapsing-remitting, RR; 11 secondary progressive; SP; seven primary progressive) and 11 healthy controls were recruited. High resolution T1 maps were acquired using a three dimensional fast low angle shot sequence. Dual echo T2 weighted images were also acquired. Disability was assessed using the expanded disability status scale (EDSS). Normalised white matter histograms were generated after extracting skull, cerebrospinal fluid and grey matter. T2 lesion volume was determined using a semiautomated technique.

Histogram standard deviation (SD) was significantly greater in each patient subgroup than in controls (p<0.0001), and in patients with SPMS compared with RRMS (p<0.02). In the overall patient cohort, histogram SD correlated significantly with T2 lesion volume (r=0.79, p<0.0001), and disability (r=0.53, p=0.004).

White matter histograms evaluate global tissue damage in MS, and may provide a sensitive, objective measure for monitoring disease progression.

OLIGODENDROCYTES SECRETE FACTORS WHICH DECREASE NEURONAL APOPTOSIS IN VITRO VIA PI3-KINASE/AKT SIGNALLING PATHWAYS

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There is increasing data both from experimental models and from pathological and clinical studies of multiple sclerosis showing that oligodendrocytes provide trophic support for neurons and withdrawal of that support may contribute to neuronal dysfunction. Understanding the nature of this support and underlying intracellular mechanisms may lead to development of therapeutic strategies to prevent neuronal/axonal abnormalities occurring in multiple sclerosis. We have used an in vitro model to study neuronal-oligodendrocyte interactions.

Rat embryonic cortical neurons were cultured in media conditioned by oligodendrocytes at different stages of their lineage. Neuronal survival was measured by immunocytochemistry and by ‘H GABA uptake. Apoptosis was measured by TUNEL staining. Protein kinase analysis was studied by western blotting and immunocytochemistry.

Neurons showed a marked increase in survival when cultured in medium conditioned by oligodendrocyte precursor cells and differentiated oligodendrocytes, and a decrease in apoptosis, compared to non-conditioned media. This effect was blocked by inhibitors of PI kinase but not by MAP kinase inhibitors. Furthermore oligodendrocyte conditioned media caused an increase in Akt (protein kinase B) phosphorylation in neurons consistent with an anti-apoptotic effect.

This study provides evidence that soluble factors from oligodendrocytes support neuronal survival and decrease apoptosis via PI kinase/Akt signalling pathways.