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 no 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 Th7 level. Deep tendon reflexes were normal with no pathological reflexes. This hyperhidrosis distribution was confirmed by the Minor test and quantified with a Servomed evaporimeter. Thermography showed reduced body temperature of the same distribution as hyperhidrosis. Radioisotope detection showed scoliosis at Th1-Th9. Magnetic resonance imaging showed syringomyelia prominent at Th2-Th4. No Chiari malformation was present. Her hyperhidrosis was relieved by oral administration of 400 mg/day mexiletine (fig 1). Oral administration of 400 mg/day carbamazepine also gave partial relief. She did not have to change her nightwear a few times each night. Neurological examination showed bilateral hypeaesthesia at Th2-Th5 and hyperhidrosis from the face to the dermatome Th10 level on both sides. Deep tendon reflexes were normal with no pathological reflexes. Hyperhidrosis was confirmed by the Minor test and quantified with a Servomed evaporimeter. Thermography showed decreased body temperature over the same area as the hyperhidrosis was distributed. Magnetic resonance imaging showed a cavernous haemangioma at Th2-Th5. His hyperhidrosis was relieved by oral administration of 200 mg/day mexiletine (fig 1). Oral administration of 400 mg/day carbamazepine also gave partial relief. He did not have to change his clothes during the day.

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, such as postural change, or bladder and intestine stimulation.

We judge that the mechanism of segmental hyperhidrosis in our patients was produced by overactivity of the preganglionic sympathetic neurons. The intermediolateral nucleus, located in the lateral horn of the spinal cord from C8-Th1 to L2-L4, receives both excitatory and inhibitory descending innervations from the thalamus. The face and neck are innervated by the intermediolateral nucleus at the Th2-Th4 level, in the upper limbs at Th2-Th6, and in the lower limbs at Th11-Th12. The ambiguous, overlapping innervation by the intermediolateral nucleus accounts for the differences in the distributions of the sensory disturbances and hyperhidrosis caused by the same spinal cord lesion. The hyperhidrosis distributions in our patients corresponded to the area innervated by the intermediolateral nucleus at the levels of the lesions seen on MRI and of the sensory deficits. This hyperhidrosis distribution was explained by interruption by the inhibitory descending pathway. Because hyperhidrosis in our patients occurred even without 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. 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. 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
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 confirmatory sign of 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 of carbamazepine was 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 retromastoid 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 intracysternal 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 with multiple sclerosis.

Although the intimate aetiopathogenetic mechanisms of trigeminal neuralgia still remain unknown, peripheral lesions affecting the trigeminal nerve entry zone (tortuous vessels, meningeal arteries, aneurysms, arteriovenous malformations, 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 cases of vascular compression and demyelination of intrapontine trigeminal fibres in case of multiple sclerosis (personal observation of MR in more than 80% of cases) may result in ephaptic, abnormal transmission of impulses. A neuropathological study of focal demyelination at the root entry zone was performed 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 cases of 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 generation of paroxysmal attacks. These findings led us to propose microvascular decompression to the patient rather than percutaneous lesions. 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” mechanism 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 mechanism (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 this 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 to 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 ischaemic 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.

In our case the MRI evident T2 hyperintensity of intrapontine trigeminal pathways was considered a demyelinating or an ischaemic lesion of unknown aetiology.

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 3 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
Sodium nitroprusside was administered over a period of 40 minutes. The dosage was chosen from an ear-
ly clinical report. The dosage was calculated to achieve a mean arterial pressure of 2 mm Hg, a total dose of 40 mg sodium nitroprusside was administered over a period of 40 minutes. The dosage was chosen from an ear-
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References

Transcranial magnetic stimulation alleviates truncal ataxia in spinocerebellar degeneration
Spinocerebellar degeneration is an inherited or acquired neurodegenerative disorder characterised by steadily progressive cerebellar ataxia, dysarthria, 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 depression1 and Parkinson’s disease. We also reported the efficacy of TMS for inherited spinocerebellar degenerations.

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degeneration. They were divided based on the date when active stimulation, and the other 35 patients, (19 men and 20 women), were assigned to 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 (19 men and 20 women), were assigned to active stimulation, and the other 35 patients, aged from 38 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 (χ2 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 corticocerebellar 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 centred on the nuchal region, 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 inter-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 clockwise) 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 (ability to stand on one foot more than 15 seconds; 4, able to stand with the feet more than 10 cm apart; 3, 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 (ability to walk without support, but clearly abnormal; 3, unable 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 blood flow in each region of interest was calculated from qualitative axial SPECT images by the application of Patlak graphical analysis.7 The examiners were blind to the manner of TMS. All participants 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<0.05), 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<0.05) (Mann-Whitney U test, table 1). After the 3 week TMS trial, the mean regional brain 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 cerebral 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.

### Table 1

<table>
<thead>
<tr>
<th>Age (y)</th>
<th>Disease duration (y)</th>
<th>Disease types (%) (cerebellar:OPCA)</th>
<th>10 m time (s)</th>
<th>Tandem steps:</th>
<th>Standing capacities:</th>
<th>Walking capacities:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before TMS</td>
<td>After TMS</td>
<td>Before TMS</td>
<td>After TMS</td>
<td>Before TMS</td>
<td>After TMS</td>
<td>Before TMS</td>
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<tr>
<td>14.37 (1.80)</td>
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<td>19.47 (1.19)</td>
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<td>2.36 (0.18)</td>
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<td>13.78 (1.23)</td>
<td>12.65 (1.20)*</td>
<td>25.19 (1.71)</td>
<td>23.49 (1.69)*</td>
<td>2.26 (0.67)</td>
<td>2.43 (0.21)</td>
<td>2.11 (0.17)</td>
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<tr>
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<td>0.64</td>
<td>&lt;0.05</td>
<td>0.19</td>
<td>0.72</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>13.74 (2.92)</td>
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<td>23.91 (2.22)</td>
<td>19.97 (1.89)</td>
<td>3.25 (1.38)</td>
<td>4.07 (1.34)</td>
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<tr>
<td>16.24 (2.26)</td>
<td>12.16 (1.21)</td>
<td>24.41 (3.62)</td>
<td>19.39 (1.55)</td>
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<td>0.72</td>
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<td>10.72 (3.52)</td>
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<td>&lt;0.05</td>
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<td>0.13</td>
</tr>
<tr>
<td>0.51</td>
<td>0.10</td>
<td>0.23</td>
<td>&lt;0.05</td>
<td>0.20</td>
<td>0.17</td>
<td>0.01</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 cerebellar type and OPCA type of active stimulation group.
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

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References


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 infrahemispheric mechanisms, as judged mainly through studies on patients with alexia and agraphia.2 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 26/56.

Brain CT on admission showed an area of high density consistent with contusional haemorrhage in the left temporo-occipital lobe. Brain MRI 3 days after admission showed contusional haemorrhage in the left temporo-occipital lobe involving the posterior part of the temporo-occipital lobes 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 5 days after admission 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.

Although this patient experienced aphasia immediately after cerebral contusion, the agraphia 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

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**Table: Kanji and Kana**

<table>
<thead>
<tr>
<th>(1)</th>
<th>(2)</th>
<th>(3)</th>
<th>(4)</th>
<th>(5)</th>
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<tbody>
<tr>
<td>毛系</td>
<td>邊</td>
<td>灰皿</td>
<td>鉛筆</td>
<td>新聞</td>
</tr>
<tr>
<td>(1)</td>
<td>(2)</td>
<td>(3)</td>
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<tr>
<td>大</td>
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<td>茶</td>
<td>銅</td>
<td>新聞</td>
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<tr>
<td>(1)</td>
<td>(2)</td>
<td>(3)</td>
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<td>(5)</td>
</tr>
<tr>
<td>犬</td>
<td>猫</td>
<td>茶</td>
<td>銅</td>
<td>新聞</td>
</tr>
</tbody>
</table>

**Figure 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.
exhibited amnestic aphasia, alexia, and aggraphia in the acute phase, disturbances other than agraphia of Kanji disappeared in a few months. Yokota et al. also reported a case with pure aggraphia 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 depends on the intactness of this route. 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 aggraphia 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 case was caused by the left posterior temporal lesion which disconnected the pathway for Kanji writing selectively.

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Figure 2. Brain MRI 3 months after admission showed the subcortical lesions in the left temporo-occipital lobe.
In March 1999 he was treated with BEAM myeloablative chemotherapy (300 mg/m^2 BCU on day −6, 200 mg/m^2/day etoposide on days −5, −4, −3, and −2, 200 mg/m^2/day cytoxane arabinoside on days −5, −4, −3, and −2, and 140 mg/m^2/day melphalan on day −4) followed by stem cell re-infusion at day 0 (11.2 × 10^7 CD 34+ cells/kg). After re-infusion recovery of peripheral blood cells occurred within 16 days. PredinRose 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 predinrose and without treatment. He works full time and the only neurological signs consist of mild numbness of his fingertips. Electrophysiological studies were repeated 1 year after re-infusion and compared with the latest study performed when he had relapsed after the mobilisation phase. For the right median nerve, the distal latency was 14.6 to 12.3 ms, and the motor nerve conduction velocity was from 12 to 24 m/s.

This is the first report of autologous CD 34+ stem cell transplantation in a patient with CIDP. Réalem reviewed the results of autologous stem cell transplantation (ASCT) in autoimmune diseases. He discussed a group of 143 patients, including patients with multiple sclerosis (n = 51), systemic lupus erythematosus (n = 15), juvenile chronic arthritis (n = 18), systemic lupus erythematosus (n = 12), vasculitis (n = 4), other connective tissue disease (n = 4), and various autoimmune haematological disease (n = 9). The treatment of these patients was variable: some had transplantation of bone marrow stem cells, others had mobilised peripheral blood haematopoietic progenitor cells for transplantation. The group was unmanipulated in 36 patients and was T cell depleted in the remaining patients. The highest relapse rate at 1 year was seen in juvenile chronic arthritis (50%) whereas only 10% of patients with multiple sclerosis had relapsed. The transplant related mortality at 2 years post-transplantation was 6%. Multiple sclerosis has become the autoimmune disease which is most often treated by ASCT. McCombe et al reviewed a group of 24 patients with multiple sclerosis with a progressive course who were treated similarly to the patient with CIDP in this report. The patients with multiple sclerosis were conditioned with the BEAM regimen; the mobilisation phase consisted of cyclophosphamide followed by granulocyte colony stimulating factor and finally CD 34+ cells were cryopreserved. The only difference was that our patient was not treated with antithymocyte globulin. In the group of 24 patients with multiple sclerosis, one patient died of aspergillosis in the post-transplant period. The other 23 patients had transplant related mortality. Improvement in disability was seen in 10 patients, stabilisation in another 10. The question is how ASCT is capable of eradicating autoimmunity. Remissions after autologous hematopoietic stem cell transplantation are easier to explain, as a “diseased” immunological and stem cell compartment is replaced with one that is normal and likely to develop tolerance. However, allogeneic transplantsations are accompanied by high mortality figures and are therefore not considered in non-lethal autoimmune diseases.1

Autologous transplantations can better be described as rescue therapy. High doses of cytotoxic therapy as is given in ASCT are also myeloablative and are only possible with stem cell rescue treatment. It might be that the effects of this treatment on autoimmune diseases can be attributed completely to the high levels of cytotoxic treatment. However, development of tolerance by “re-educated” lymphocytes from the purified CD 34+ cells has not been ruled out.

We certainly do not advise the routine use of ASCT in patients with CIDP. Most patients have an excellent response to immunomodulating therapy without serious side effects. Moreover, after several years of treatment many patients reach long lasting remissions. However, if high doses of immunosuppressive therapy are required for a long time periods and if they are poorly tolerated, ASCT with a well tolerated conditioning regimen may be considered.

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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 meningo-radiculois. Repeated spinal MRI disclosed an epidural 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 meningo-radiculitis including neuroborreliosis, meningeval lymphoma should be considered as an important differential diagnosis. In view of the occasionally difficult diagnosis of primary meningeal lymphoma, we hereby bring the clinicopathological report of a primary malignant lymphoma with almost selective 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 tumour marker CD 20. Nerve conduction studies suggested subclinical demyelinating neuropathy without motor conduction block. Cranial and cervical 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 corticosteroid and intravenous immunoglobulin treatment, her condition deteriorated rapidly with development of flaccid paraplegia and paresis of the right arm. Repeat 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 20 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 thalamic 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 no vascular infiltration. Unfortunately, none of the lymphocytes, searched for systemic malignant lymphoma was negative except for an isolated small focal area of B cell lymphoma in the thyroid gland. As in the case reported by Wilkening et al, this finding widens the range of possibilities and diagnostic difficulties of selective involvement of the spinal roots (and peripheral nervous system) by primary malignant non-Burkitt’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

http://jnnp.bmj.com/ J Neurol Neurosurg Psychiatry: first published as 10.1136/jnnp.72.1.123 on 1 January 2002. Downloaded from
Orientation agnosia in pentagon copying

Aila 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 Aila 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 Aila 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 patients’ 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 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 Aila 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 Aila et al by virtue of the criteria by which they define an abnormal performance.

In figure 2 of the paper of Aila 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.

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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. 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 which 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 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).

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 like 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 DLB were included in the 134. Perhaps many of those who had the orientation agnosia were actually patients with DLB who did not exhibit the classic DLB clinical symptoms of fluctuation, visual hallucinations, and parkinsonism. All of that is not to say that Della Sala and colleagues cite as evidence that there is “no significant difference in drawing or constructional abilities between AD and LB dementia” cannot be easily compared with ours because of the different severities of the patients involved. Based on MMSE scores, at least one half of the patients in that study were more severely impaired (median AD MMSE 10.5, DLB 9.0) than any of the patients in our study (minimum MMSE 13). That study does help to confirm that constructional abilities worsen as the dementia worsens, for as we mentioned in our results, none of the additional cases of AD and DLB 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 AD 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

The book is part of a series based on the proceedings of the International Winter Conference on Neurodegeneration and Neuroinflammation. This particular volume arose from the seventh meeting, which was held in Karuizawa, Japan in January 1999.

The conference brought together scientists from a wide range of research areas, under the broad heading of neurodegeneration, and their breadth of expertise is reflected in the eclectic mix of studies presented here. The papers fall into four general categories, paralleling the scientific sessions of the meeting. The first deals with progress in elucidating some of the genes and gene products involved in neurodegenerative processes. It includes an interesting paper on the cleavage of huntingtin by caspases and the possibility of using caspase inhibitors to stop the aggregation and hence neurotoxicity of the protein in Huntington’s disease. This section also includes papers on inherited Parkinson’s disease, both autosomal dominant and recessive mechanisms, with particular attention being paid to the role of the PARK1 gene in the juvenile onset form of the disease.

The second, and largest, group of papers address the very topical subject of cell death mechanisms in neurodegeneration. The areas covered range from the role of the anti-apoptotic protein Bcl-2 in spinal muscular atrophy, in vitro experiments looking at the effect of α-synuclein mutations in neuronal cell cultures, to the possible role of polymorphisms in the GABA-A receptor γ1 subunit and the myelin oligodendrocyte glycoprotein (MOG) gene in models of multiple sclerosis. Perhaps surprisingly, there was relatively little on the toxicity associated with amyloid precursor protein processing in Alzheimer’s disease. However, that may not have been such a bad thing in this context, as a thorough discussion of this topic could easily require a separate volume in itself. Also included in this section (although it’s not entirely clear why) is an interesting re-evaluation of basal ganglia circuitry, by one of the editors, Professor Riederer, based on a review of clinical and experimental evidence.

The third section reflects the burgeoning interest in the role of cytokines in neurodegeneration and includes papers on the role of cytokines in Parkinson’s disease and in oligodendrocyte death. Unfortunately, there is no discussion of their role in Alzheimer’s disease where their involvement has long been suspected. There has subsequently been a major increase of interest in this area with the discovery of polymorphisms within the genes for interleukin 1 and their effects on disease prevalence and age of onset.

The final section is more speculative and deals with some possible future therapeutic interventions. These include the use of adenovirus associated viral vectors for gene therapy in Parkinson’s disease, neuroprotective therapies in multiple sclerosis, and the possible efficacy of transglutaminase inhibitors in the treatment of polyglutamine disorders.

Overall, I found this an interesting volume to dip in to but it suffers from the common problem of all meeting proceedings in that two and a half years down the line some of the findings are already beginning to look somewhat dated. It is perhaps worth noting that, according to the publisher’s website volume 8 of this series, based on the meeting in Bavaria in February 2000, is already available.

Stephen Gentleman

EMG waveforms: video companion to electromyography and neuromuscular disorders

The practice of electromyography, despite many advances in signal analysis, quantification, and computer modelling, remains to an extent an art. The experienced electromyographer relies as much, if not more so, on his ears as well as his eyes to recognise patterns of muscle electrical activity.

Transferring this experience to trainees is a challenge, and this collection of video clips covering spontaneous activity (fibrillation, complex repetitive discharges, fasciculations, and various forms of myasthenia and myoneuromyasthenia) and voluntary EMG activity is a useful addition to our material. Each clip shows a standard EMG machine screen with an audio soundtrack and is introduced by a static video frame. Clips run for the minute and often display waveforms at different time resolutions. The making of measurements is also shown. It would have been preferable for a voiceover to have introduced each clip to highlight the points to be made. Reading the accompanying booklet works but less impact. Some screen shots either show redundant information or small characters indicating, for example, amplitude and frequency.

Nevertheless this is a valuable resource to introduce trainees to the range of electrical muscle activity which might be encountered in clinical practice.

Kerry Mills

Neuropsychological evaluation of the older adult: A clinician’s guidebook

This book gives comprehensive and thorough up to date coverage of this topic. It starts from the premise that evaluation of the neurocognitive function 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 and test finding and writing up. A 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 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 dementias—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 therapeutic nihilism to enthusiasm. Stroke is now
Neurobiology of spinal cord injury


“The future for regenerative therapies of the spinal cord is clearly very bright”—this ends this volume in Humana’s Contemporary Neuroscience series, which deals with 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 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 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 up to date, and written with authority, although sometimes maybe 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 experiences gained from human spinal cord injury. None the less, 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...
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 acknowledgement that the editors were ably and fortunately assisted by a certain Michelle Pfeiffer—surely not that one!

D M A Mann
PROCEEDINGS

Association of British Neurologists Autumn Meeting, Newcastle University meeting at Durham, 12–14 September 2001

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 fulfill 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 questionnaire group, was the ACE score. In combination, the medial temporal lobe rating and graded naming test were also contributory.

MRI imaging 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 PITUITARY-DERIVED HUMAN GROWTH HORMONE
RJ Cordery, M Hall, L Cipolotti, L Davidson, P Adlard, MN Rossor National Hospital for Neurology and Neurosurgery, London, UK

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 frontal executive dysfunction. These findings suggest that, although not the presenting feature, mild cognitive decline may be evident in the early stages of CJD associated with human cadaveric growth hormone treatment.

LATERALITY OF UNEXPLAINED MOTOR AND SENSORY SYMPTOMS: A SYSTEMATIC REVIEW 1965–2000
J Stone, A Carson, R Goldbeck, S Lewis, B Thomas, CP Warlow, M Sharpe University of Edinburgh, UK

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 laterality of unexplained motor and sensory symptoms except for right sided movement disorder and possibly left sided symptoms in women. Retrospective studies specifically looking at laterality 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 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). The 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)
PA Barber, MD Hill, AM Demchuk, JH Warwick Pexman, ME Hudon, A Tomanek, RR Frayne, AM Buchan for the ASPECTS Study Group Foothills Medical Centre, Calgary, Canada

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 considered less sensitive and reliable than DWI for the detection of ischaemic change in hyperacute stroke. We hypothesised that 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 mental line, and appropriate window width (80 HU) and centre level review settings (40 HU). MR imaging was performed on a 3Tesla machine and included DWI (single shot EPI TR=7000 ms, TE=99 ms, b=1000 mm s$^{-1}$, 19 5 mm slices/3 mm gap). A quantitative scoring system (ASPECTS) was applied to CT and DWI at baseline and follow up (within 36 hours) by five
PATIENTS WITH TIA AND ISCHAEMIC STROKE IN RELATION TO BRAIN TISSUE.

The challenge for MRI is to reliably identify salvageable brain tissue. The mean onset to 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 hypoxic-ischaemic stroke, even 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 stroke, while the challenge for MRI is to reliably identify salvageable brain tissue.

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, PM Rothwell

Some rare mendelian stroke syndromes have been characterised, but the genetic epidemiology of the common forms of ischaemic stroke and TIA remains 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 TIA's (hospital based series) related history of stroke in first degree relatives (FDRs) to baseline clinical characteristics, subtype of stroke, and risk of recurrence.

Information on family history was available in 537 strokes and 467 TIA's.

Strokes had occurred in at least one FDR in 101 (18.9%, 95% CI 16–22) strokes and in 98 (21%, 95% CI 17–25) TIA's. 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 of 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, RJ Wise

The production of both propositional speech (PrSp) and non-propositional speech (NPrSp) in normal subjects activates the left frontal 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 (NPrSp), PrSp in response to autobiographical questions, and a non-speech state (XSp) when they attended to occasional environmental sounds. In normal subjects, PrSp and NPrSp contrasted with XSp revealed activation within Broca's area and the left anterior insula. In patients, activation was seen in the right frontal operculum. 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 nervous systems engaged in speech production, and confirms Mohr'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 annual incidence 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 (n=187). Non-specific visual problems 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, NP Pavese, 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 year, respectively), while 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 been 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.
Neurons) were demonstrated in areas shown only by quantitative MRI analysis to be abnormal. The proportion with additional quantitative abnormalities, which could be additional subcortical malformation, in our FCD group is comparable with that with a poor outcome after focal FCD resection.

Our technique could have potential application as a presurgical screening tool to prevent patients unlikely to achieve seizure freedom from being needlessly exposed to the risks of surgery.

**A PROFILE OF EPILEPSY IN A UK POPULATION**

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**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.

**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 23% >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%).

**Multimodal MRI Imaging of Epilepsy:**

**The TSC and 10 (33%) FCD patients had additional extralesional abnormalities. Some forms of FCD may represent a forme fruste of tuberous sclerosis (TSC).**

**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 = 1.5% (OR 6.27, 95%CI 3.2–11.7). Developmental delay was associated with exposure to valproate (OR 4.42, 95%CI 1.77–11.5). However, mothers of children with epilepsy had low educational attainment; 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.

**QUANTITATIVE MRI ABNORMALITIES WITH HISTOPATHOLOGICAL CORRELATION IN REFRACTORY EPILEPSY: A TOOL TO PREDICT SURGICAL OUTCOME?**

TN Mitchell, SL Free, M Thor, SM Sisodiya, SD Sharvon Institute of Neurology, Queen Square, London, UK

**Objective:** to determine the significance of focal cortical dysplasia (FCD) in the aetiology of refractory epilepsy by comparing the pattern of abnormalities identified on MRI with the histopathological appearance associated with these abnormalities.

**Methods:** Seventy-four patients with FCD underwent quantitative MRI with determination of T1 relaxation times of subcortical white matter. Two independent raters correlated MRI findings with histopathology from resected and unoperated brain.

**Results:** In 44% of FCD, significant increase in T1 relaxation times of subcortical white matter was identified. Of these, 73% had lesions on MRI that were not visible on preoperative MRI. The most common abnormalities associated with FCD were polymicrogyria (34%), oligodendroglioma (17%), and neuronal heterotopia (17%). In patients with FCD who progressed to surgery, histopathological examination revealed focal cortical dysplasia (FCD) in 43% and neocortical heterotopia (NH) in 17%. Twenty percent of patients with FCD had significant malformation of the external capsule. MRI identified 292 patients whose mothers gave consent to follow up with developmental assessment using Bayley’s scales.

**Conclusion:** MRI was identified as an aetiological factor in 73% of patients with MRI and histopathological abnormalities in 60% of patients with FCD.

**FAST MICROGRAPHIA AND PALLIDAL PATHOLOGY**

NP Quinn Institute of Neurology, Queen Square, London, UK

**Objective:** to investigate the relationship between fast micrographia and pallidal pathology in patients with bilateral lesions of the globus pallidus.

**Methods:** The presence of fast micrographia was assessed in 125 patients with bilateral lesions of the globus pallidus. Fast micrographia was also identified in 10 patients with unilateral lesions of the globus pallidus who had a history of micrographia on the unaffected side.

**Results:** Fast micrographia was present in 98 patients. The micrographia was evident at onset in 70 patients and in 28 patients developed at a later date. The micrographia persisted in 74 patients and improved in 32 patients. Fast micrographia was most often observed in patients with progressive supranuclear palsy, who sometimes volunteered “microscopic” writing as their presenting complaint. However, it can also be seen in patients with acquired bilateral lesions of the globus pallidus. Illustrative videotapes will be shown.

**Conclusion:** Fast micrographia suggests a deficit in patients with enhanced movement-related feedback to the motor cortex and that fast micrographia can be seen in a variety of neurological disorders associated with abnormalities of the basal ganglia.
The most often reported syndromes were sensory neuropathy (33%) and cerebellar degeneration, either pure (21%) or in association with brainstorm/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/Brain Tumours
A J Wills, DSNA Pengiran Tengah Queens Medical Centre, Nottingham; Derbyshire Royal Infirmary, Derby, UK

Objective: To audit patients referred from general practitioners (GPs) via the 2 week wait system and contrast with neurological cancer patients identified independently of this system.

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 gliomas, one cerebral metastasis). The remainder were diagnosed with chronic daily headache (10), epilepsy (five), migraine (three), demyelination (two), essential tremor (two), other (16). At least 69 neurological cancers were identified during this period who were not referred via the 2 week system. Cancer CNS cancers present in a diverse manner. Eliciting a neurological history and interpreting neurological signs can be challenging and many GPs have little neurological training. These guidelines, even when followed stringently, do not seem to increase diagnostic precision. Inappropriate referrals have extended already lengthy waiting times for neurology outpatient clinics and may have inadvertently delayed other patients with urgent neurological problems being seen. We suggest that the use of these guidelines in neurological practice is reconsidered at the earliest opportunity. Perhaps a system of closer communication between primary and secondary care would be a more effective use of time and resources.

INREVERSIBLE DAMAGE TO THE SPINAL CORD AFTER SPINAL ANAESTHESIA
K Hamandi, J Mattershead, T Lewis, IEC Ormerod, IT Ferguson Frenchay Hospital, Bristol, UK

Neurological features were a recognised rare complication of spinal anaesthesia. A number of mechanisms of injury to the spinal cord have been suggested. We report five cases of damage to the distal spinal cord after spinal anaesthesia. All patients were female undergoing orthopaedic (three patients) or obstetric (two patients) surgery. They had significant unilateral leg weakness and sensory disturbance after anaesthesia, which persisted at 2 year follow up. They had significant unilateral leg weakness and sensory disturbance after anaesthesia, which persisted at 2 year follow up. These patients were not responsive to various treatments including intravenous immunoglobulin and plasmapheresis. They had significant unilateral leg weakness and sensory disturbance after anaesthesia, which persisted at 2 year follow up. There were no significant adverse events.

A DOUBLE BIND PLACEBO CONTROLLED STUDY OF INTRanasal SUMATRIPTan IN ACUTE CLUSTER Headache
PJ Goodson, JA van Vliet, A Bahra, V Martin, SK Aurora, NT Mathew, MD Ferrari Leiden University Medical Centre, Leiden, The Netherlands; Cincinnati Headache Center, Cincinnati, USA; Headache Center, Seattle, USA; Houston Headache Clinic, Houston, USA; Institute of 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 received the time of onset of the attack, time of treatment, headache severity on a five point scale (0-nil, 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 response 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 multilevel analysis approach with MVLWiN (www.ioe.ac.uk/multilevel) 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 of cluster headache and other variables of interest, the 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%) and 33/56 (59%), 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. For cluster headache patients these data add a further evidence based treatment to the management portfolio of this disabling primary headache.

EFFECT OF THE ADENOSINE (A1) RECEPTOR AGONIST GR79236 ON TRIGEMINAL NOCICEPTION WITH BLINK REFLEX RECORDINGS IN HEALTHY HUMAN SUBJECTS
NJ Giffin, F Kowacs, V Libri, P Williams, PJ Goodson, H Kaupe Institute of Neurology, Queen Square, London, UK

There are no good migraine models in humans, but development of a low current intensity trigeminally mediated blink reflex (BR) has allowed the study of nociceptive transmission in the trigeminal nucleus caudalis. Adenosine is antinociceptive in humans and animals. GR79236 is a highly potent and selective adenosine A1 receptor agonist. We investigated the effect of GR79236 on the nociceptive BR using 2 female healthy volunteer randomized in a placebo controlled double blind crossover study. BRs were elicited by supraorbital nerve stimulation with standard or nociception-specific electrodes at random intervals of 12-18s. BR responses were recorded bilaterally in the supraorbital nucleus. Nine measurements were made of 20 sweeps. Areas under the curve (AUCs) of the R2 component of the BR were calculated from the rectified EMG, before and 30 minutes after drug (GR79236 (10 mg/kg)) or placebo. Primary end point: median AUC at 30 minutes, modelled using analysis of covariance. There was a non-significant reduction of the ipsilateral nociceptive R2 after GR79236 vs placebo of 17%. However, there was a significant reduction of the contralateral nociceptive R2 (p<0.05) of 20%. There were no significant adverse events.

The results suggest that GR79236 may inhibit trigeminal nociceptive pathway. This reduces sensitivity to pain. The most likely mechanism of injury. This the first study to report detailed neuroimaging and long term follow up in all cases which shows that permanent neurological deficit has occurred. The use of spinal and epidural anaesthesia is becoming more widespread. Neurologists are likely to be involved in the management of complications and should be aware of this rare but serious complication.
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%) of 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.

HETEROPLASMIC 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 mtDNA 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 genetic 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 heteroplasmatic 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 heteroplasmy in human pedigrees with mtDNA disease. We also provide the first direct evidence of a mitochondrial genetic bottleneck in humans, enabling the first quantitative 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 Institute of Neurology, Queen Square, London, UK

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 analyse the clinical and genetic features of myotonia congenita in the UK, to look at genotype-phenotype correlation in this group, and to develop a DNA based diagnostic service.

In collaboration with the British Neurological Surveillance Unit we identified over 100 families with inherited myotonia. We initially performed SSCP analysis of all 23 exons of the CLCN1 gene in index cases from 42 families. In order to check if no mutations had been missed with this technique we subsequently analysed all the exons by direct DNA sequencing.

Using SSCP analysis mutations were identified in 50% of cases. This figure rose to 65% with DNA sequencing. In this mutation positive group the age at onset varied from 1–16 years (mean 3.6), muscle pain was reported in 20% and 50% reported exacerbation of muscle stiffness in a cold environment. Twelve novel mutations have been identified. In particular the finding of a severe myotonic phenotype in a family with a main haplogroup and heterozygosity and a mild phenotype with a new homozygous nonsense mutation (C242X) will be highlighted.

DNA based diagnosis is a realistic option in myotonia congenita. Although the mutations identified are spread across the gene it seems that exon 8 is a hot spot in this population. Genetic counselling in this disorder is complex and should include consideration of both genetic and functional expression data if available.
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, Ryeathgate 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 CECC. 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 CECC 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 increase in 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 dystinesias with capcortomia. 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 daytime sleepiness, (3) Daytime PaCO2 >45 mm Hg, (4) SaO2 <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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<th>Criteria for NIV</th>
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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 vs 8.47 h/day) and less improvement in QOL (effect size: 1.85 vs 3.28).

Conclusions—Orthopnoea was the best predictor of compliance with and benefit from NIV in 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), Ewthorpe sleepiness scale (ESS), and ALS functional rating scale (ALSFRS). 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 ≥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 Berkley, 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 in a group of five subjects with gluten sensitivity. We report the first missense mutation in LAMP-2 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 Gunnewald, AK Chattopadhayay, 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 made by a visual analogue scale improved from mean score (SD) at 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.

Intravenous 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 treatment with a gluten free diet. There was some increase in amplitudes in the group treated with diet: 1.4 ± 1.3 v 2.3 ± 2.3 µV (p<0.01) but none in the control group: 1.6 µV v 1.1 µV (p<0.1). 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, trapping 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 normal. A left occipital skull fracture with an organising extradural haematoma indenting the left occipital lobe was demonstrated. Abnormal high signal was seen within the sagittal sinus. MRV demonstrated venous thrombosis occluding the posterior two thirds of the sagittal sinus. The patient was anticoagulated and symptoms resolved in a few days.

A UNIQUE MODEL OF LISSENCEPHALY

L Kinton, KA Barth, C Howarth, 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.

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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 BY NEUROLOGY OUTPATIENTS BEFORE CONSULTATION**

Al 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 two cases, 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 Deurson, 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, both at home and in hospital. An interviewer administered diary, the Rivermead mobility index (RMI), and a digital camcorder recording of gait (with calculation of stride length and hence distance) were also used.

Patient 1 (female; 74 years) recovering from Guillain-Barré syndrome, was independent (RMI: 8/15). The patient was active for on average 2.95 hours per day [range 1.3 to 4.1 h/d], and 18.43% of time awake. He walked an average of 1791 right steps covering a distance of 1593 m [range 857–2710 m] Activity levels rose after discharge from hospital with a change in activity pattern from directed activity to less vigorous, frequent independent activity. Patient 2 (female; 34 years) with improving inflammatory cervical myelopathy, was also independent (RMI: 8/15). This patient had very limited activity which coincided with physiotherapy, with little independent activity. As expected the two controls had far higher activity levels. Despite similar RMI scores, the patients showed very different activity levels: both showed reduced activity levels with walking, in one case virtually dependent on therapy. These data emphasise the wide range of normal activity levels in healthy and disabled people and the potential use of this technique in quantifying and monitoring mobility.

**SPECIFYING THE EXECUTIVE DEFICIT IN PARKINSON’S DISEASE: A COMBINED BEHAVIOURAL AND EVENT RELATED fMRI STUDY**

SJJ Lewis, A Dove, R Cools, TW Robbins, RA Barker, A M Owen

**Cambridge Centre for Brain Repair, Cambridge, UK**

Parkinson’s disease (PD) is accompanied by a complex pattern of frontal-like impairments, which may include deficits in many aspects of working memory function. However, recent neuropsychological studies have questioned the psychological specificity of these “executive” deficits, while functional neuroimaging studies have been unable to identify the precise neural substrates responsible. In this study, two closely matched groups of patients with PD, differing only with respect to their performance on a standard test of executive dysfunction, were tested on a novel verbal working memory paradigm which allowed different aspects of performance to be assessed simultaneously, including maintenance, retrieval, and manipulation of the remembered information. The subgroup of patients with “executive” impairment were selectively impaired on the task, and specifically in those aspects of the test requiring the manipulation of information within working memory. These results suggest that it may be informative to functionally subdivide patients with PD according to whether their cognitive profile includes executive dysfunction or not; moreover, the specific nature of the impairment observed implicates the dorsolateral region of the frontal cortex as a likely neural substrate. To test this hypothesis directly, event related fMRI was employed in similar groups of PD patients performing the same verbal working memory task.

**GLUTEN ATAXIA IS IMMUNE MEDIATED**

D Mahad, K Oates, C Williams, R Grunewald, SJL Howell, MN Woodroofe, M Hajivassiliou

Royal Hallamshire Hospital, Sheffield, UK

Gluten sensitivity is a state of heightened immunological responsiveness triggered by the ingestion of gluten in genetically susceptible individuals. Organ specific manifestations include an 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 compared with controls. 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.

<table>
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<th>Patient 2</th>
<th>Control 1</th>
<th>Control 2</th>
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</tbody>
</table>

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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 scores were available for 22. The age after 2.9 years follow up 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=65.9 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 in established cognitive impairment, suggesting that this is a dementia 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 <1/2 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 the slow form, MRI brain showed focal cortical atrophy. On histology, there was moderate perivascular T cell infiltration, microgliosis, and reactive astrocystosis. In both forms, there was bilateral focal cortical hyperperfusion 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 with epilepsy and performed comprehensive screens for anti-CNS and other autoantibodies. 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
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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 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.

Objectives: 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 followed for 2 years after their 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 Sharvon, JWA Sander, RM Gardiner, JS Duncan, NW Wood Institute of Neurology, Queen Square, London, University College London, 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 which 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 chromosome 1q and 2q and the febrile seizures loci on chromosome 8q, 19p, 2q, and 5q 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 RATIO AND N-ACETYL ASPARATE AS INDEPENDENT MARKERS OF MYELIN AND AXONAL INTEGRITY IN DEMYELINATING LESIONS: A MAGNETIC RESONANCE IMAGING STUDY OF CENTRAL PON TINE 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 Uhatoo, 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.

A postal questionnaire to consultant adult and paediatric neurosurgeons in the United Kingdom was 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 that they 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 epilepsy surgery in the last decade, this still falls far short of what is normal apparent 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 obtained. 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 T1 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**

A Wilkins, S Chandran, DAS Compston Cambridge Centre for Brain Repair, Cambridge, UK

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 [3H] 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 oligodendrocytes at different stages of their lineage. Neuronal survival was measured by immunocytochemistry and by [3H] 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.