Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsia partialis continua and motor epilepsy

JMA COWAN, JC ROTHWELL, RJS WISE,* CD MARSDEN

From the University Department of Neurology, Institute of Psychiatry, and King's College Hospital Medical School, and the MRC Cyclotron Unit,* Hammersmith Hospital, London, UK

SUMMARY A patient is described who had a combination of stimulus-sensitive cortical myoclonus, epilepsia partialis continua, and Jacksonian motor epilepsy. He eventually required surgery because of the severity of his seizures. Electrophysiological recordings made before and during surgery, and PET scans performed before surgery identified an abnormal area of cerebral cortex in the post-central parietal region. It is suggested that the stimulus-sensitive myoclonus arose because input into this region from peripheral sensory afferents produced an abnormal discharge which was fed forwards via cortico-cortical connections to the precentral motor cortex, to produce a reflex muscle jerk. The epilepsia partialis continua may have been caused by spontaneous discharges arising in the same region of parietal cortex. Both forms of jerking disappeared after resection of this part of the cortex.

Much has been written on the subject of human myoclonus, especially that arising in the cerebral cortex. Patients with cortical myoclonus frequently exhibit abnormally large cortical sensory evoked potentials (SEPs) and, with back-averaging techniques, it is often possible to detect a time-locked potential in the electroencephalogram (EEG) over the contralateral sensorimotor cortical area preceding spontaneous jerks. Such neurophysiological investigations usually have had to rely upon SEPs and EEGs recorded from scalp electrodes, which may not provide accurate localisation of the source of abnormality. This report concerns a patient with stimulus-sensitive cortical myoclonus, epilepsia partialis continua, and Jacksonian motor epilepsy, who eventually came to surgery because of the severity of his seizures. This provided a rare opportunity to correlate findings from direct cortical recording with those of scalp EEG, both before and after resection of an area of the post-central parietal cortex identified as the abnormal site. This area was shown to be metabolically abnormal, prior to surgery, by positron emission tomography, even though computed tomographic (CT) scans showed no abnormality other than atrophy. The opportunity was also taken to examine long-latency stretch and cutaneous reflexes after excision of this area of cortex.

Case report

PM, a right handed 20-year-old male was referred by Dr M Yealland, in February, 1982 because of intractable epilepsy. There was no family history of neurological illness. His birth and early history were normal. At the age of 9 years he began to suffer from seizures. These commenced with a strange feeling in the left upper limb which then started to shake. Gradually, the shaking spread to involve the rest of the limb and then the left side of the face, the left side of the trunk and the left lower limb. Each seizure lasted about one to two minutes. He did not lose consciousness, but was incontinent of urine during attacks in the first few years. The seizures continued at a frequency of about four to eight per week. At the time of presentation, each attack lasted for about one and a half minutes with the left upper and left lower limbs feeling “numb” for about an hour afterwards, during which time he would feel sleepy. There had been no response to treatment with phenytoin, carbamazepine, phenobarbitone, primidone, valproate or clonazepam, all given to the point of toxicity. The fits became more frequent on occasions and he had to be admitted to hospital once in 1980 and twice in both 1981 and 1982, because of repeated seizures, virtually amounting to status epilepticus.

In 1978, he began to notice increasing clumsiness when
Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsy partialis continua and motor epilepsy

Fig 1   A. View of the operative field. The anterior aspect of the wound is on the left, and the superior aspect is at the bottom of the photograph. The right frontal, central and parietal regions can be seen through the craniotomy. The first area to be resected is outlined by the black thread. Small letters placed on the pial surface of the brain indicate the approximate extent of the motor cortex, being the lowest threshold points from which contralateral movements could be elicited by electrical stimulation. B. Section of intra-operative corticogram taken with the patient conscious. The traces are unipolar records from ball electrodes on the pia referred to a large metal clip on the skull. Numbers at the end of each trace indicate the electrode positions in the diagram of figure 7. Position 7 is the point of maximal spontaneous spike activity.

using the left hand and forearm. In the middle of 1980, he developed a continual twitching of the left fingers and hand which was always present, except during sleep, or after a flurry of seizures. Because of both the clumsiness and the twitching, he had to give up work as a trainee gardener, in 1981. Six months before his first admission to this department, he developed a twitch of the left side of the mouth which was similar to, and was present simultaneously with that of the fingers. On admission, he complained of left sided weakness.

On examination, he was alert and orientated. His speech was slow, deliberate and slurred, but there was no dysphasia. There was continual twitching of the left side of the mouth and jerks of the fingers of the left hand. The left hand and forearm would jerk in response to a variety of different stimuli, including light taps to the finger pads, electrical shocks of the digital nerves, and tendon taps to the finger flexor and extensor muscles. Limb bulk, tone and power were normal. There was impairment of position and two-point discriminatory sensations in the left fingers and a slight decrease of light touch and pin-prick sensations in the whole of the left hand.

Electroencephalograms showed gross deficiencies of fast rhythms over the right central and parietal areas. In addition, there was an excess of slow activity on the right, especially in the sylvian-central area. Multiple spike foci were seen on the right side, frequently in the central-parasagittal, mid-central, sylvian-central and mid frontal areas, and occasionally in the right temporal region. There were no left sided spikes. A normal response to photic stimulation was evident, bilaterally.

Skull radiographs were normal. CT scans with contrast showed atrophy of the right hemisphere with a prominent cisterna magna. Right carotid angiography was normal. Positron emission tomographic (PET) scans and detailed neurophysiological studies were performed and are detailed below.

On psychological testing, the short Wechsler Adult Intelligence Scale (WAIS) showed a verbal IQ of 84, a performance IQ of 79 and a full-scale IQ of 81. No firm conclusions could be made as to whether or not a focal brain disturbance might be responsible for the low scores.

Because of the inability of anti-convulsant drugs to control his disabling seizures, and their focal origin, it was decided to proceed to electrocorticography with a view to excision of the focus so defined. A two-stage operation was performed by Mr C Polkey. During the first stage, a right temporo-parietal craniotomy was performed under general anaesthesia. Three days later, the second stage was undertaken with the patient awake. Omnopon and atropine had been given the night before, as premedication. Electrocorticography (Dr M Driver), identified the abnormal discharges as arising mainly from the right parietal lobe (for details see results). Following electrocorticography, general anaesthesia was induced. A piece of parietal cortex, measuring 6 cm × 4 cm × 3 cm was removed (fig 1). Because of continued spiking seen on the EEG, a further 3 cm × 2 cm × 1 cm area was excised from the superior margin of the initial resection line. The tissue taken felt tough and abnormal. After the resections, there was still spontaneous spiking from a pre-central lead (position 4, fig 7). Because this area felt normal and had a normal reaction to cortical stimulation, it was decided not to resect it. A small piece of the excised tissue was placed in glutaraldehyde for electron microscopy and the rest was placed in formalin.

The specimens were examined by Dr B Dunkley. The grey-white junction was softened, granular and yellowish in two different areas. Several blocks were taken for light microscopy and all showed similar changes. The cortex was reduced to an outer zone of densely gliotic tissue of varying depth, containing varying numbers of neurons, and an inner zone of loose astrocytic fibrils and prominent capillary blood vessels, but no neurons. Many gemistocytic astrocytes were present in this inner zone and occasional nerve cell bodies were seen remaining in the “spongy” astrocytic mesh-
work. The white matter showed fairly good preservation of myelin staining, but here too there was astrocitic gliosis. The leptomeninges were thickened by fibrosis and were adherent to the surface of the cortex. The blood vessels of the meninges and of the cerebral parenchyma appeared to be normal. There was no evidence of inflammation. Electron microscopy did not reveal any viral particles.

The patient made a good recovery from the operation, having only one seizure on the first post-operative day, during which the head and eyes turned to the left. There were also a few short-lasting episodes of twitching of the left thumb, but this then disappeared. Carbamazepine and clonazepam were continued.

The operation, however, resulted in further deficit. There was a left inferior quadrantanopia, a mild weakness of the left upper and lower limbs when tested against resistance, diminution in pin and touch appreciation in the left hand and forearm and marked worsening of position sense in the left hand, wrist and elbow. It was now difficult to maintain a constant position with the left hand, even with visual feedback. Neuropsychological assessment revealed significantly lower scores on tests of visual constructive ability and on the WAIS than pre-operatively.

In the eighteen months since the operation, the spontaneous twitching of the left upper limb has returned on occasions, but is absent almost all of the time. The seizures have been as frequent as before, but have not involved as wide an area of the body, and there have been no further episodes of status epilepticus or severe fits. The left upper limb reflexes have become brisk. The post-operative reduction of pin-prick and touch sensations have recovered, but the visual and proprioceptive defects have remained, unaltered. The patient was able to return to the gardening job that he had had to leave because of the seizures.

Methods

Physiological testing

Neuropsychological studies included routine EEG polygraphy, back-averaging of the EEG activity preceding spontaneous jerks, somatosensory evoked potentials, muscle stretch reflexes and cutaneous reflexes.

Preliminary EMG recordings were made from up to eight muscles that were involved in the myoclonic jerks. From these records, the muscle with the most well-defined and regular EMG burst was chosen to trigger collection of data for back-averaging. Each single trial was collected separately and was later averaged. In this way, any record showing movement artifact or a poor trigger point could be rejected on visual examination.

The EEG was recorded using 1 cm diameter silver/silver-chloride disc electrodes. Similar electrodes were placed over the cervical vertebrae, at Erb's point and on the medial aspect of the arm, 10 cm above the elbow over the median nerve, for somatosensory evoked potential recording, and 2–3 cm apart over the bellies of muscles studied for EMG activity. The signals were processed and pre-amplified with Devices 3160 pre-amplifiers and were amplified using Devices 3120 amplifiers. Filters were set 3 dB down at 80 Hz (high-pass) and 2.5 kHz (low-pass) for EMG and short-latency somatosensory evoked potential recording, and at 0.16 Hz (high-pass) and 2.5 kHz (low-pass) for EEG and long-latency somatosensory evoked potential recording. EMG signals were rectified or smoothed, depending on the requirements of individual studies. The resulting signals were stored on paper, on magnetic tape using a Racal tape recorder running at three and three quarter inches per second, or on floppy discs using a DEC PDP-12 computer, running programmes written by Mr HB Morton.

Somatosensory potentials were evoked either by mixed nerve stimulation of the median nerve at the wrist, or by cutaneous nerve stimulation using ring electrodes around the index finger. A Devices 3073 constant voltage stimulator was used. Stimuli were large enough to just cause visible movement of the fingers, when given at the wrist, or to give rise to a solid, but not uncomfortable tapping feeling, when delivered to the fingers (an intensity approximately twice sensory threshold). In both cases, stimuli were timed so as not to coincide with electrocardiographic activity. Because sensation was reduced in the left hand, in order for the stimulus to be appreciated, the shocks applied to the left fingers were of a greater voltage than those applied to the right. Short latency spinal and cortical potentials were evoked by stimuli given at approximately 3 Hz and recorded with a sampling frequency of 5 kHz per channel. Long latency somatosensory evoked potentials were evoked at random intervals every 2–3.5 seconds and were recorded with a sampling frequency of 1 kHz per channel. There is some difficulty in labelling the components of the somatosensory evoked potential in patients with grossly abnormal potentials. In this paper, as before, 2 the major components were designated by polarity and sequence (N1, P1, N2 etc), to avoid equating any of the myoclonic potentials with those seen in normal individuals (N20, P25/P30, N35).

Stretch reflexes were evoked in the flexor pollicis longus muscle by rapid extension of the interphalangeal joint of the thumb with the other related joints being securely clamped. The patient was asked to exert a small, constant torque of 0.06 Nm by flexing the pad of the thumb against a torque motor. Every 2–3.5 seconds, the torque was suddenly increased by a factor of two for 200 ms, or by a factor of four for 50 ms (in different studies), and the resulting EMG was recorded from the agonist muscles.

Cutaneous reflexes were studied using ring electrode stimulation of the ring finger at three times sensory threshold. Stimuli were given at 2 Hz, while the patient exerted a constant contraction of his first dorsal interosseous muscle by fanning out the fingers of the hand, according to the technique of Jenner and Stevens. 3 The averaged, rectified, EMG response of the first dorsal interosseous muscle to 256 stimuli was obtained.

A ring block of the thumb was produced with 4 ml of 1% plain lignocaine.

Intra-operative electrocorticography was performed using metal ball electrodes on the pial surface. The signals were fed into a conventional EEG machine and selected signals were recorded on magnetic tape for later analysis, using an output from the EEG machine taken after the pre-amplification stage as the input for the tape recorder.

PET scanning

The patient was studied twice pre-operatively with positron emission tomography and the oxygen-15 steady state inhalation technique, to determine regional cerebral blood
Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsy partialis continua and motor epilepsy

flow (rCBF), oxygen consumption (rCMRO2) and their relationship—the fractional extraction of oxygen (rOER). There was an interval of one week between the studies, and on the second occasion, consecutive use was made of the oxygen-15 and fluorine-18 labelled 2-fluoro-2-deoxy-D-glucose (18F deoxyglucose) as tracers, in an attempt to determine regional glucose metabolism (rCMRGlu), and the relationship between rCMRO2 and rCMRGlu—the metabolic ratio (rMR). rCMGlu was calculated using a simplified Sokoloff operational tracer equation. A lumped constant was incorporated into the tracer equation to define the different affinities of deoxyglucose (tracer) and oxygen (natural substrate) for the blood-brain barrier carrier molecule and intracellular hexokinase, and a value for this constant described for normal brain was used in this study. Regional cerebral blood volume (rCBV) was also measured on both occasions, using carbon-11 monoxide as a tracer; rCBV provides an index of local vasodilatation, and the data was also used to correct the overestimation of rOER and rCMRO2 that is inherent in the simplified operational equation for the oxygen-15 steady-state technique as originally described.

Results

PHYSIOLOGY

Pre-operative

The EMG activity responsible for the spontaneous myoclonic jerks was found to be most prominent in the left first dorsal interosseous muscle, but also involved the left thenar, left finger flexor and left finger extensor muscles (fig. 2). The duration of each burst was brief, lasting from 30 to 60 ms. Two jerks sometimes occurred in rapid succession, separated by an interval of less than 30 ms. The frequency of the myoclonus ranged from less than 1 Hz to approximately 5 Hz. At first sight, the activation of the individual muscle groups in each jerk appeared to be approximately synchronous. However, closer examination showed that there was a slight “jitter” of about 4 ms between the onset of activity in the first dorsal interosseous and the other muscles. Because of this, the intrinsic hand muscles were sometimes activated before the forearm muscles and sometimes after. Back-averaging the EEG activity preceding spontaneous jerks revealed a positive-negative wave, the positive peak of which preceded the onset of EMG activity in the first dorsal interosseous muscle by about 20 ms. This positive-negative wave was largest over the somatotor hand area of the right hemisphere where it measured 17 μV (fig. 3).

Myoclonic jerks could also be provoked by electrical stimuli of the cutaneous nerve of the forefinger, or by mixed nerve stimulation of the median nerve at the wrist, in the left arm. The same stimuli also evoked large potentials over the right hemisphere. Figure 4 shows the EMG and EEG responses to electrical stimuli applied to the forefinger. Stimulation of the left forefinger produced a reflex jerk in the left

Fig 3. Back averaged (of 128 sweeps) EEG activity preceding spontaneous myoclonic jerks in the left first dorsal interosseous muscle (1st DI). A large positive-negative wave precedes the onset of EMG activity by about 20 ms. Electrode spacing 2 cm. Electrode 3 is 7 cm lateral to the vertex. Linked mastoid reference.

Fig 2. Sections from surface EMG records showing detail of three separate spontaneous myoclonic jerks (A, B, C) in the left hand. Jerks are most prominent in the first dorsal interosseous muscle (1st DI), but also occur synchronously in thenar (TH), forearm flexor (FF) and forearm extensor (FE) muscles. A rapid double jerk is illustrated in A.
forearm flexor and hand muscles, with a latency of 50 ms. This was preceded in the SEP, by a large P1–N2 response (45 μV), maximal over the contralateral central region. The P1 peak had a latency of 30 ms which was some 20 ms prior to the reflex EMG response in the hand. In contrast, stimulation of the right hand produced no reflex muscle jerk and cerebral potentials over the contralateral hemisphere were barely visible when amplified by the same amount as the signals on the other side.

The short-latency SEPs, following stimulation of the median nerve at the wrist are shown in fig 5. The gains have been increased in order to demonstrate the early responses. The peripheral (axilla and Erb’s point) and spinal potentials were normal following stimulation of either side. However, the large P1–N2 component over the right hemisphere is not so evident in this figure, because low frequency components have been filtered in order to display the short-latency components, and the reference electrode was placed at Fz rather than using linked mastoids. The N1 response occurred at the same latency as the N20 response on the normal side and was slightly smaller.

Muscle stretches also elicited myoclonic jerks, in the left forearm and hand. When the terminal phalanx of the thumb was extended, stretch reflexes were evoked in the flexor pollicis longus (FPL) muscle and a large reflex jerk was seen in the first dorsal interosseous muscle (fig 6). The response in the first dorsal interosseous muscle lasted for 65 ms, and occurred 7.5 ms after the normal responses in the FPL muscle. Responses of such a size in the first dorsal interosseous are not seen following thumb stretch in normal individuals and it is presumed that the response in PM was a reflexly-evoked myoclonic jerk. It was accompanied by a large, contralateral, somatosensory evoked potential.

In order to determine whether cutaneous or muscle afferents were responsible for the reflex jerk, the left thumb was anaesthetised with a lignocaine ring block. This abolished all cutaneous sensation in the thumb and also abolished the giant SEP in the cortex following electrical stimulation of the thumb (fig 6). As in normal individuals, the stretch reflex response in the flexor pollicis longus was reduced by anaesthesia. Despite this, the myoclonic response in the first dorsal interosseous muscle produced by thumb stretch was relatively unaffected, as was the somatosensory response that such stimuli evoked. Hence muscle afferents were involved in the myoclonic responses, although a small contribution from cutaneous afferents innervating the skin over the stretched flexor pollicis longus muscle cannot be excluded.

b Intraoperative

The pre-operative studies suggested that this patient’s myoclonus was associated with abnormal discharges in the contralateral sensorimotor cortex. Intraoperative recording allowed the site of the abnormality to be defined with far greater precision. Spontaneous spike discharges were seen (fig 1b), especially at position 7, where they were largest and most frequent, but also at position 15 and, to a lesser extent at positions 4 and 5. Intravenous sodium thiopentone was given. This caused position 6 to become the site of most frequent spiking, as well as producing the expected increase of background rhythms in all channels. Electrocorticography revealed little abnormality of the precentral cortex. The motor strip was mapped out using 50 Hz alternating current stimulation and was judged to be normal.

While conscious, electric shocks were given to the left index and middle fingers, at an intensity of twice sensory threshold. The resulting somatosensory evoked potentials were very similar to those seen with scalp recording (fig 7). The major deflection at electrode 6 had an onset, latency and polarity equivalent to the P1–N2 component recorded from the scalp. No N1 component was evident, probably because of a relatively low setting of the high frequency filters used on the EEG machine preamplifiers. The smaller potentials recorded at the other electrodes probably are due to volume conduction of the electrical activity at or near electrode 6.

The patient had no spontaneous myoclonus during
Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsia partialis continua and motor epilepsy

Fig 5 Short-latency SEPs following stimulation of the median nerve at the wrist. A: pre-operative, B: three weeks post-operative. Records are the average of 999 sweeps. Traces from ipsilateral axilla and Erb's point are bipolar surface recordings. Traces from an electrode over the fifth cervical spine (C5) and from contralateral sensory hand area (scalp) (7 cm lateral on a line joining the external auditory meatus to a point 2 cm posterior to the vertex) are unipolar, referred to an electrode at Fz. Post-operatively, all cortical potentials disappear after stimulation of the left side, although the early potentials in subcortical tracings are unchanged. (The later waves in the C5 pre-operative recording are conducted from the brain via the FZ reference.) The difference in size between the axilla potentials on each side probably was due to slightly different electrode placings on each arm. The difference in pre-operative N1 amplitudes is within the normal range. Stimulus intensity was at motor threshold on each side. The giant P1–N2 component, following stimulation of the left side is not seen as clearly here as in the previous figure, because low frequency components of the EEG signal have been filtered out to display the short latency events more clearly. The 10 μV calibration refers to the top trace and the 5 μV calibration refers to the bottom three traces, in each section.
Fig 6  Effect of thumb anaesthesia produced by a local anaesthetic ring block of the proximal phalanx on the EEG and EMG responses to cutaneous or stretch stimuli applied to the left (abnormal) thumb. Records on the left show the response to electrical stimulation (STIM) of the digital nerves with ring electrodes. In the intact (control) state this evokes a large SEP over the contralateral hand area (EEG: unipolar, referred to linked mastoids) and reflex muscle jerks in flexor pollicis longus (FPL) and the first dorsal interosseus (1st DI) muscles. With the thumb anaesthetised, all these responses disappear. Records on the right show the responses to forcible extension of the distal phalanx of the thumb by a torque motor. In the control state, traces show a large SEP, and reflex responses in FPL and 1st DI. The position record shows the excursion of the thumb joint. In the 1st DI, the responses to stretch are barely changed after thumb anaesthesia, whilst they are considerably reduced in FPL. Traces are averages of 128 trials.

the operation, perhaps due to the premedication, and so back-averaging could not be performed. Neither was there any reflex myoclonic response in the left hand to electrical stimulation of the fingers.

c  Post-operative
Studies were repeated on several occasions during the next year. No spontaneous myoclonus and no reflex myoclonus could be found post-operatively in response to tapping, muscle stretch, or electrical shocks to the left hand, all of which had produced myoclonus pre-operatively. The N1, P1, N2 phases of the somatosensory evoked potential were found to be absent over the right hemisphere, when the stimulus was given to the left hand. Spinal potentials were preserved (fig 5). The contralateral somatosensory evoked potential following right hand stimulation was unchanged compared with records made before the operation.

The opportunity was also taken to investigate the effect of the removal of a well defined region of the parietal cortex on two groups of long-latency reflexes, postulated by some authors to involve transcortical pathways. When the right flexor pollicis longus muscle was stretched, normal spinal and long-latency reflexes were seen in the EMG (fig 8A). However, when the same stimulus was applied to the corresponding left sided muscle, no long-latency response could be seen. The short-latency response on the left was now twice its pre-operative size (fig 6) and was twice the size of the normal short-latency response in the right arm.

Cutaneous reflexes were studied in the first dorsal interosseus muscle, following stimulation of the index finger. The responses on the right were normal and consisted of a small short-latency E1 phase followed by the long-latency I1 and E2 responses. However, on the left, only the E1 response could be seen, the I1 and E2 responses being absent (fig 8B). It should be recalled that such stimulation had produced a gross myoclonic jerk on the left side before operation.
PET SCANNING

Positron emission tomographic scans were performed at a number of transaxial planes above the orbitomeatal (OM) line. It was evident, on inspection of the functional data, that there was high rCBF throughout the right parietal lobe, but low rCMRO2, resulting in low fractional extraction of oxygen—rOER—from the arterial blood. The region of high flow also had high rCBV, which reflected the local vasodilatation. The values of rCBF, rCMRO2, rOER and rCBV from left and right parietal regions, 7 cm above the OM line, for both studies are summarised in the table; they are compared with normal values from six normal volunteers (four males, two females, mean age 35 ± 6 years, range 22–37 years).

It is evident that the right parietal region had much lower rCMRO2 at the times of the second study. For technical reasons, it was not possible to record the EEG during the PET studies, but it is tempting to conclude that the right parietal cortex was electrically less active during the second study; clinically, occasional jerks of the digits of the left hand occurred during both studies.

The first study included a transaxial plane at the level of the cerebellar hemispheres. rCMRO2 and rCBF of the cerebellum (1.7 ml/100 ml/min and 25 ml/100 ml/min, respectively) were less than half the normal values determined at the MRC Cyclotron Unit. There was no functional asymmetry between the two cerebellar hemispheres.
Table  rCBF, rCMRO2, rOER and rCBV from right and left parietal regions, at 7 cm above the OM line, from the two studies on the patient, contrasted with values from six normal volunteers

<table>
<thead>
<tr>
<th></th>
<th>rCBF</th>
<th>rCMRO2</th>
<th>rOER</th>
<th>rCBV</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>L</td>
<td>R</td>
<td>L</td>
<td>R</td>
</tr>
<tr>
<td>Study 1</td>
<td>41</td>
<td>76†</td>
<td>3·1</td>
<td>2·2*</td>
</tr>
<tr>
<td>Study 2</td>
<td>37</td>
<td>53*</td>
<td>2·7</td>
<td>0·9†</td>
</tr>
<tr>
<td>Normals (n = 6)</td>
<td>44</td>
<td>45</td>
<td>3·5</td>
<td>3·6</td>
</tr>
<tr>
<td>±SD</td>
<td>±4</td>
<td>±4</td>
<td>±0·5</td>
<td>±0·5</td>
</tr>
</tbody>
</table>

rCBF ml/100ml/min; rCMRO2 ml/100ml/min; rCBV ml/100ml.
*2–3 standard deviations from normal.
†3 or more standard deviations from normal.

Fig 9 Functional transaxial images from the oxygen-15 and 18F deoxyglucose study on the patient. The lighter the region of an image, the greater the value of a particular variable within that region. The plane depicted was 7 cm above the OM line. The low rCBF, rCMRO2 and rCMRGlu of the skull and scalp appear as a faint halo around the brain images of these variables. The images of the ratios appear larger because the low extracerebral signal creates patchy light- and/or dark-edge artifact when divided. rCBF in the anterior part of the right parietal lobe was relatively higher than rCMRO2 and rCMRGlu, reflected in the low rOER and rGER, respectively. rCMRGlu was preserved relative to rCMRO2 and therefore the rCMRO2:rCMRGlu ratio (rMR) was low in the right parietal lobe.
Electrophysiological and positron emission studies in a patient with cortical myoclonus, *epilepsia partialis continua* and motor epilepsy

Figure 9 shows grey-scaled functional images obtained from the second study at 7cm above the OM line. As well as CMRO2, CBF and OER, transaxial images of CMRGlut, fractional extraction of glucose (GER) and the CMRO2:CMRGlut ratio are depicted. The low rCMRO2, high rCBF and low rOER in the right anterior parietal region are evident. As calculated, rCMRGlut was a little higher in the right parietal region than the left (6-6 ml/100 ml/min and 6-3 ml/100 ml/min, respectively), but rGER and rMR were lower. rMR in the right parietal lobe was one third that in the mirror region.

**Discussion**

The case reported here is a rare example of a patient with cutaneous and stretch reflex myoclonus of the left arm; regular, repetitive, continual, spontaneous myoclonus (*epilepsia partialis continua*); single partial seizures; and repeated partial seizures which required to be treated as status epilepticus.

There were a number of neurophysiological similarities between the *epilepsia partialis continua* and the cortical reflex myoclonus which suggest that both forms of jerking were manifestations of the same underlying deficit. Thus, the muscles involved, the duration of the EMG jerks and the presence of a large, contralateral, positive-negative potential in the EEG some 20 ms prior to the EMG burst were similar whether the jerks occurred spontaneously or whether they were evoked by somatosensory stimuli. Indeed, both forms of jerking disappeared after resection of the same area of the cortex. As in other cases, the N1 phase of the SEP was normal on the affected side of this patient, whilst later waves (P1–N2) were enlarged. The N1 phase probably corresponds to the N20 potential seen in normal subjects, which represents arrival of the afferent volley at the cortex. If this is so, then the large later wave may represent abnormal cortical processing of normal afferent inputs.

Although indicating the diversity of pathology that can give rise to the condition, previous reports of *epilepsia partialis continua* have tended to emphasise that an abnormality of the motor cortex is responsible for the disorder (see Thomas *et al* for references). Thomas *et al* discussed whether or not cortical or subcortical mechanisms are responsible for *epilepsia partialis continua*. They presented 32 cases of their own, including eight that had been examined after death and stated, “Autopsy findings in eight cases showed consistent involvement of the motor cortex or closely adjacent areas.” They also marshalled other evidence in order to show that it was a motor cortical abnormality that had been responsible for the jerking, in their patients, no matter where the pathologic lesion had been found. However, it has been argued before that areas of cortex other than the motor strip may be abnormal in patients with cortical myoclonus or *epilepsia partialis continua*.

Although the large, contralateral spike, which precedes cortical myoclonic jerks in the majority of patients, is most easily explained as the surface recording of a large paroxysmal depolarisation shift in motor cortical output cells, discharge of which is directly responsible for the myoclonic jerks, this may not always be true. Such potentials may be localised some centimetres anterior or posterior to the motor cortex, and the size of the giant SEPs may be dissociated from the size of the myoclonic jerk. Intraoperative monitoring revealed the state of affairs in the present patient. The abnormal area of cortex lay posterior to the central sulcus and it was from this region that large SEPs could be recorded. The same region also gave rise to spontaneous spike discharges. However, the site of maximal spontaneous spike recording (electrode no 7, fig 1) was different from the site which giant somatosensory potentials could be evoked (electrode no 6). Both sites, however, were in the right parietal lobe and were within a few centimetres of each other. Perhaps the whole region was electrically unstable and this instability caused one area to respond in an exaggerated, but limited manner when receiving stimuli from its usual input pathway, whereas the other area was even more excitable and so tended to discharge either spontaneously, or else in response to otherwise undetectable stimuli. The resulting abnormal parietal cortical activity, via cortico-cortical connections between pre- and postcentral areas of cortex, may well have “driven” a physiologically normal motor cortex to discharge, and so produce visible muscle jerks. In this way, both reflex and spontaneous myoclonus could be explained. Furthermore, spread of the spontaneous discharges within the abnormal parietal lobe could have fed forwards to a progressively larger area of the motor cortex, and so caused the patient’s Jacksonian seizures. However, after surgery this patient’s reflex myoclonus and *epilepsia partialis continua* disappeared, but his Jacksonian seizures persisted. Perhaps the latter continued because the area of persistent spiking in precentral cortex, demonstrated at electrocorticography, was not removed.

Surgery for *epilepsia partialis continua* has been reported previously. However, it has been frontal cortex that has been removed and this, of course, has caused the myoclonus to cease, if for no other reason than that the area resected included the cortical area responsible for the production of the clinically observed abnormality, whether or not that area was in itself abnormal. Interestingly, Thomas *et al* found surgery to be unsuccessful in controlling the *epilepsia partialis continua* in several of their patients and per-
haps this is further evidence that the true seizure source lay outside the area resected (the frontal lobe).

The reflex myoclonus in this patient was of special interest in view of the recent debate about the existence of transcortical long-loop reflexes in man. It has been claimed by those who originally described such reflexes\textsuperscript{14, 15} that, in the upper limb at least, the long-latency phase depends on a transcortical mechanism for its production. Others have disagreed with this view and have suggested that the spinal apparatus could be solely responsible for the production of both the early and late phases, whether due to its own intrinsic delays (to explain the later response),\textsuperscript{16, 17} or due to a second, later signal reaching the cord via slower peripheral nerve mechanisms, such as group II afferents,\textsuperscript{18} or even to repetitive burst discharges in muscle spindle Ia afferents following stretch.\textsuperscript{19} The long-latency responses to both stretch and cutaneous stimuli were abolished by resection of only the sensory cortex and sub-cortical white matter and this strongly supports the argument that such long-latency responses depend on a trans-cortical pathway and, furthermore, that this pathway must traverse the parietal lobe. Short-latency stretch and cutaneous reflexes, which are spinal in origin, increased in size post-operatively, whereas the long-latency reflexes disappeared. This gives further weight to the argument that the long-latency responses depend upon a different pathway.

The PET scan findings were unexpected in that although the low CMRO\textsubscript{2} in the right hemisphere corresponded to the atrophy seen on the X-ray CT scan, the right parietal rCBF was disproportionately high relative to the rCMRO\textsubscript{2}. Not only was this reflected in the low rOER values, but was also higher than normal, despite the presence of atrophy. The tracer technique employed measured capillary blood flow and, therefore, this high flow cannot be explained by arteriovenous shunts. Furthermore, there was no evidence of any capillary angioma on histopathological examination of the resected cortical specimen. It would seem likely that the abnormal electrical activity had provoked vasodilatation and a high rCBF. In this context, it is of interest that studies of global CBF and rCMRO\textsubscript{2} during electroconvulsive therapy in man have demonstrated a disproportionate rise in CBF relative to the CMRO\textsubscript{2}, and animal experiments have also shown an increase in venous PO\textsubscript{2} during seizures.\textsuperscript{20} The low CBF and CMRO\textsubscript{2} of the cerebellum may have been due to several factors, including drug therapy (the patient was on carbamazepine and clonazepam at the time of study), permanent dysfunction secondary to previous drug toxicity, and cerebellar atrophy. Low cerebellar metabolism has been observed previously in a group of patients with temporal lobe epilepsy.

The combined oxygen and glucose study suggests that there was non-oxidative metabolism of glucose in the right parietal region, which, in the presence of an adequate oxygen supply, implies aerobic glycolysis. Excessive lactate production, without tissue hypoxia has been demonstrated during induced seizures in animals,\textsuperscript{20} and the same observation in man is, perhaps, not unexpected. However, there are reservations about the quantitative use of \textsuperscript{18}F deoxyglucose as a tracer in the study of epilepsy. Deoxyglucose has a greater affinity for the blood-brain carrier molecules than glucose, but has less affinity for hexokinase. If the rate limiting step for hexose phosphorylation shifts from the hexokinase-catalysed reaction to transport of substrate from plasma to tissue, the value of the lumped constant, which describes the differential rate of phosphorylation of deoxyglucose and glucose, will increase.\textsuperscript{21} Status epilepticus has been demonstrated to be associated with a reduction in tissue glucose,\textsuperscript{20} implying that under these pathological conditions, blood-brain carrier transport decreases. Therefore, the use of a "normal" lumped constant in this study for the calculation of rCMRGlu in all regions may have resulted in an overestimation of the degree of non-oxidative metabolism of glucose in the right parietal lobe. Recently, a method of quantifying the value of the lumped constant regionally, using an additional tracer study, has been suggested,\textsuperscript{22} and this may permit more certain estimates of rCMRGlu in seizure disorders.

Nevertheless, despite uncertainties about \textsuperscript{18}F deoxyglucose tracer modelling, this is the first study to demonstrate uncoupling between regional oxygen supply and demand, and a possible regional disturbance of normal oxidative metabolism of glucose, in focal epilepsy in man.

The histology was most unusual. Neuronal loss and gliosis have been reported in the brains of those who have suffered from epilepsy. These changes have been ascribed to the effects of the seizures,\textsuperscript{23} but there are significant differences between these changes and those seen in the patient reported here. In the former cases, the neocortical changes consisted of neuronal loss mainly confined to the intermediate layers, whereas in the case presented here, the destruction was most marked in the inner layers. Also, the dense gliosis seen in the outer zone of the cortex of this patient contrasts with the relatively mild (Chaslin's) gliosis found subpially in the brains of chronic epileptics. Before operation, it was suspected that the patient had a form of chronic encephalitis,\textsuperscript{24} because of the history of normal early development, the progressive course, the restriction of clinical and EEG change to the right hemisphere and the suggestion on the PET scan of abnormal metabolism in the right parietal lobe which might have hinted at viral infec-
Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsia partialis continua and motor epilepsy

However, no inflammatory cells were seen in the resected specimen. Thus, the cause remains unknown.

We thank PM for his interest and co-operation during the course of these investigations. The specialised equipment was designed and built by Mr HC Bertoya and Mr R Miller. This work was supported by the Medical Research Council. JCR was a Royal Society University Research Fellow.

References

Electrophysiological and positron emission studies in a patient with cortical myoclonus, epilepsy partialis continua and motor epilepsy.
J M Cowan, J C Rothwell, R J Wise and C D Marsden

J Neurol Neurosurg Psychiatry 1986 49: 796-807
doi: 10.1136/jnnp.49.7.796