Parkinsonism following bilateral lesions of the globus pallidus: performance on a variety of motor tasks shows similarities with Parkinson’s disease

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There is considerable interest in the pathophysiology of motor dysfunction in different basal ganglia diseases, especially in idiopathic Parkinson’s disease. Preparation and execution of voluntary movements assessed using different reaction time paradigms,1,2 or through recording of movement related cortical potentials (MRCP)3,4 have been extensively studied in PD. The overall conclusion is that the preparation, initiation, and the execution of voluntary movements are slow or abnormal in patients compared with healthy control subjects.5 It is also known that bradykinesia in PD becomes more prominent by repetitive, simultaneous, or sequential movements.7 Despite these interesting results, it has been very difficult to examine the role of individual nuclei in the pathophysiology of motor dysfunction in PD, as the underlying alterations in different pathways and nuclei are complex in this disease.8 9 Although the gross morphology of the globus pallidus (GP) is normal,10 it is widely accepted that the abnormal functional output of GP is one of the major contributors to motor dysfunction in PD.3,4 The renaissance in the surgical treatment of PD has provided a new tool with which to investigate basal ganglia function. In addition to pallidal surgery, isolated pallidal lesions offer another opportunity to study the significance of GP in different motor functions. However, such lesions are rare,11 and we found only 12 reported cases of isolated bilateral pallidal lesions from the literature. To date, there is only one case report of bilateral lesions of the globus pallidus (GP), with an additional lesion of the right substantia nigra, in which numerous electrophysiological tests were performed to study different aspects of motor function.12 The results showed delayed and slow movements of the upper limbs during different reaction time paradigms and while performing a fast goal directed arm movement task.12

We now describe a case with acquired isolated bilateral lesions of the GP, on whom a variety of behavioural and electrophysiological measures were obtained to investigate different aspects of motor function. These aspects of motor function in our case, and the role of GP, are discussed by comparing the present results to previously published results of healthy control subjects and patients with PD.

CASE DESCRIPTION

Our patient was a civil servant working with electronics. He had a past history of alcohol but no disulfiram, methanol, or narcotic drug abuse. He was admitted to hospital at the age of 26 years in a disorientated state after a severe drinking binge. Shortly afterwards, he lapsed into an unconscious state and had several generalised seizures. Investigations at that stage indicated a severe respiratory acidosis. However, he recovered quickly and was discharged from the hospital.

About three months after this episode he noted problems with his writing which had become small and untidy. Over the next three years these symptoms gradually progressed.
On examination, three years later, he had a mildly abnormal gait with dystonic external rotation of the left foot. He could walk heel to toe normally and hop on either leg, but his arm swing was slightly reduced on the right. His speech was hypophonic and he manifested palilalia, and festination and freezing of speech. His face was hypomimic and he had blepharoclonus. Pursuit and voluntary saccadic eye movements were normal. The glabellar tap was positive but there were no other primitive reflexes and the jaw jerk was absent. Rapid tongue movements were normal. Voluntary palatal movements and the gag reflex were normal. Postural reflexes were impaired. Although his axial tone was increased, there was no limb rigidity. He had no tremor, and the speed and amplitude of repeated finger and hand movements and tapping tasks were normal with no fatigue. However, alternate supination and pronation of the arms revealed a dramatic deficit with rapid breakdown of rhythm, timing, and amplitude and fatigue over 5–10 repetitions. On writing there was “fast micrographia”, so that although the size of his writing was extremely small, its speed was normal. He could, however, use his hands normally for doing up buttons, tying laces, and pressing computer keys. Tapping tasks with his feet were difficult, as they were interrupted by freezing. Muscle power in the limbs and deep tendon reflexes were normal, and the plantar responses were flexor. He had no abulia. Treatment with trihexiphenidyl and levodopa had no effect. His family history was negative.

Routine blood counts, blood smears for acanthocytes, serum biochemistry including liver function tests, copper and caeruloplasmin, autoantibody screen, syphilis serology, alfa-fetoprotein, lactate, and pyruvate levels, white cell enzymes, and plasma amino acids, were all normal or negative. DNA analysis was negative for mitochondrial mutations (8344 and 8993). EEG, EMG, and nerve conduction studies and visual evoked potentials were normal. Brain MRI revealed bilateral and symmetrical signal changes (hypointensity in T1 sequences and hyperintensity in T2 sequences) in the GP, probably sparing the innermost part of the GPI (fig 1), but the rest of the brain appeared completely normal.

**Figure 1** (A) T2 weighted axial MRI image showing hyperintense bilateral and symmetrical signal changes in the GP, probably sparing the innermost part of the GP. (B) The same lesions are hypointense on T1 weighted coronal MRI image. The rest of the brain is normal on both sequences.

**METHODS**

To investigate different aspects of motor performance in our patient, and to compare him to healthy control subjects and patients with PD, we deliberately chose tasks for which we already had data from these two groups. All the investigations reported below were performed three and a half years after the incident which gave rise to the bilateral GP lesions. The patient was 29 years old at the time of the investigations.

**Reaction time tasks**

Simple reaction time (SRT), uncued choice reaction time (CRT), and precued choice reaction time (pCRT) tasks were performed as previously described. The tasks were carried out first with the right and then the left hand. For each task, a block of 100 trials was completed with each hand. For each of the SRT, uncued CRT, and precued CRT tasks, 20 trials were given for each of the five S1–S2 intervals (0, 200, 800, 1600, and 3200 ms) in pseudorandom order, with an equal number of responses to each of the four response keys. Median IT was calculated for both hands and for each of the five S1–S2 intervals.

**Pegboard and finger tapping**

The pegboard and finger tapping tasks were performed as previously described. The pegboard task involved peg placement using the Purdue pegboard. The patient performed the task with the right hand and the left hand separately, and with both hands simultaneously. On each occasion, the patient had to place as many pegs as possible in a 30 second period. In the finger tapping task, the patient was required to tap a response button repetitively using the index finger for 30 second. The tapping task was performed with the right hand, the left hand, and bimanually. In addition to performing the tapping and pegboard tasks individually, the patient also performed these tasks concurrently, tapping with one hand while placing pegs with the other.

**Flex and squeeze tasks**

Flex and squeeze tasks were performed as previously described. The patient sat with his forearm flexed at 90°, resting on a manipulandum, with his hand holding a vertical grip containing a strain gauge. The patient performed two types of movement, a 15° flexion of the elbow and an isometric 30 Newton (N) hand squeeze. Three conditions were assessed: simple flexion performed alone, the two movements performed simultaneously, and finally the two movements performed sequentially in the order squeeze then flex. For the sequential task, the inter-onset latency (IOL) between the onsets of the squeeze and flex was also determined.

**Movement related cortical potentials**

Movement related cortical potentials (MRCPs) were recorded as previously described. The patient was required to perform self-initiated joystick movements at the rate of once every 5–10 seconds, in a randomly chosen direction—left, right, forwards, or backwards. Twelve EEG channels were recorded. Electrodes were placed in three sagittal rows, according to the 10–20 international system and with three additional electrodes in FC sites (F3, FC3, C3, P3 on the left side, Fz, FCz, Cz, Pz on the midline, and F4, FC4, C4, P4 on the right side). EMG activity of the first dorsal interosseous and the abductor pollicis brevis was recorded with surface electrodes. EEG activity was back-averaged off line after realignment of the traces to movement onset, a grand average being calculated for each hand. The early (NS1) and the late (NS2) components of the averaged MRCP were identified by visual inspection. The onset of the early component was set at the onset of the rise of the slope from baseline and the end at...
the point of change of the slope, which corresponded to the onset of the late component. The end of the late component corresponded to the time of onset of EMG activity; the slopes of each component were measured.

**Contingent negative variation**

Contingent negative variation (CNV) was recorded as above, only the task performed by the subject was different. This was a ready-go task in which a pair of tones was presented via headphones. The warning tone (S1) of 500 Hz was followed by an imperative stimulus (S2, a tone of 1 kHz); the S1–S2 interval was 2.5 seconds. The subject was instructed to perform the left wrist extension task on hearing S2. The mean amplitude of the CNV from 500–1000 ms after S1 was measured.

**Cognitive function**

Our patient had neuropsychological assessment three years after his first admission to hospital. A battery of tests of cognitive function was used to assess IQ, memory, language, and visuospatial abilities and executive function. Current IQ was measured on the Wechsler Adult Intelligence Scale-Revised (WAIS-R). Estimates of premorbid IQ were obtained from the National Adult Reading Test (NART). Memory for words and faces was assessed with the Recognition Memory for Words (RMW) and Faces (RMF). To assess language and visuospatial abilities, the Graded Naming Test and the Silhouettes subtests of the Visual Object Space Perception battery were completed.

**RESULTS**

To allow comparison with the previously published data (in healthy controls and PD), all the results for the different reaction time tasks, and for the pegboard and finger tapping tasks, are given as mean values for the right and the left hand.

**Reaction time tasks**

**Unwarned and warned SRT tasks**

**Initiation times**

The mean (standard deviation) initiation time (IT) for the unwarned SRT was 456 (73) ms. The patient benefited from warning signals preceding the go-signal (at different intervals of 200–3200 ms), and his ITs were relatively constantly around 350 ms (mean 344 (11) ms) after different warning signal intervals.

**Movement times**

The mean (SD) movement time (MT) for the unwarned SRT was 193 (SD 56) ms. When the warning signal preceded the go signal (at different intervals of 200–3200 ms), the mean (SD) MT was 188 (26) ms—that is, at about the same level as in the unwarned trials.

**Use of advance information**

**Unwarned SRT versus unwarned and uncued CRT**

The ability to prepare movements in advance was assessed by comparing ITs in the unwarned SRT and unwarned and uncued CRT trials. Because the nature of the response is known in advance of stimulus presentation in SRT it can be preprogrammed, which gives SRT a speed advantage over CRT. In the present case, the difference between the mean ITs in the unwarned and uncued CRT (mean 587 (SD 128) ms) and the unwarned SRT (mean 456 (SD 73) ms) was 131 ms (fig 2A).

**Precued CRT versus uncued CRT**

In a precued CRT task, a precue presented 200, 800, 1600, or 3200 ms before a go signal provides full advance information about the particular response required on that trial. If this advance information provided by the precue is used by the subject to pre-programme responses before the go signal, then precued CRTs should be faster than uncued and warned CRTs. On average, at different precue intervals, there was no difference between the mean ITs in the precued CRT (mean 489 (SD 93) ms) and the uncued and warned CRT (mean 494 (SD 271)). Only for the 1600 ms interval was the precued CRT (mean 373 (SD 68)) clearly faster than the uncued CRT (mean 481 (SD 110)). Figure 3A summarises the reaction times in warned SRT, warned and uncued CRT, and precued CRT tasks. In terms of speed of movement execution during the various RT conditions, our case obtained mean MTs of 187.5 ms across the warned and unwarned SRT trials, a mean MT of 341.1 ms for the uncued CRT, and 186.1 ms for the precued CRT conditions.

**Performance on the pegboard and tapping tasks**

All the values given are means for the right and left hand. During the period of 30 seconds, the patient performed 185 unimanual taps. Bimanually, he was able to produce about the same number of taps (177 taps, 96% of unimanual condition) compared with the unimanual conditions. The number of pegs inserted unimanually was 13 during the period of 30 seconds. Compared with unimanual performance, the number of pegs inserted decreased slightly during the bimanual condition (11 pegs, 85% of unimanual condition).
In a more complex task, in which the patient placed pegs with one hand, and did tapping with the other, the number of taps clearly decreased (104 taps, 56% of unimanual condition). However, the number of pegs inserted remained at the same level compared with the unimanual condition (12 pegs, 92% of unimanual condition). The results of the pegboard and finger tapping tasks are summarised in table 1.

**Performance of the flex and squeeze tasks**

Movement times for the two tasks performed separately are for the right hand. In the simultaneous and sequential tasks, hand squeeze was performed with the left hand, and elbow flexion with the right hand. In the sequential task, hand squeeze was performed first, followed by elbow flexion. The movement time for squeeze alone was 158 ms. Hand squeeze performance slowed down to some extent when performed simultaneously with elbow flexion (190 ms; 20% increase in movement time). The movement time for elbow flexion alone was 209 ms. Elbow flexion performance remained at about the same level when performed simultaneously with hand squeeze (226 ms; 8% increase in movement time). In the sequential task, performance of elbow flexion clearly slowed down (297 ms; 42% increase in movement time). The results of the flex and squeeze tasks are summarised in table 2.

**Movement related cortical potentials**

Compared with the previous data for healthy controls, the movement related cortical potentials (MRCPs) preceding voluntary joystick movements were abnormal. The first component (NS1) (usually starting at about 1.5 seconds before movement onset) was absent prior to movement of either hand, as shown by the flat early slope. The later (NS2) component was present (amplitude of 5µV on the left; amplitude of 7µV on the right) (fig 4).

**Contingent negative variation**

There was no recordable contingent negative variation (CNV) in an S1/S2 task using an interval of 2.5 seconds and an imperative movement of wrist extension on the left (fig 5).

**Cognitive function**

Three years after the insult, the Verbal and Performance IQs of our case were 102 and 106, respectively. Both of these were in the average range. On the NART, his estimated premorbid IQ of 117 was in the high average range. Comparison of the verbal IQ on the WAIS-R and the premorbid estimate of IQ from the NART suggested moderate cognitive under

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**Table 1** Performance of pegboard and finger tapping tasks under unimanual, bimanual, and dual task conditions

<table>
<thead>
<tr>
<th></th>
<th>Present case</th>
<th>Controls (n = 13)</th>
<th>PD (n = 7)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age (years)</strong></td>
<td>29</td>
<td>38 (range 21–62)</td>
<td>61 (10)</td>
</tr>
<tr>
<td><strong>Pegboard</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>(unimanual)</strong></td>
<td>13</td>
<td>16 (2)</td>
<td>9 (2)</td>
</tr>
<tr>
<td><strong>(bimanual)</strong></td>
<td>11</td>
<td>14 (2)</td>
<td>7 (3)</td>
</tr>
<tr>
<td><strong>Unimanual pegboard (%)</strong></td>
<td>85</td>
<td>87 (8)</td>
<td>71 (19)</td>
</tr>
<tr>
<td><strong>Tapping</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>(unimanual)</strong></td>
<td>185</td>
<td>173 (19)</td>
<td>113 (22)</td>
</tr>
<tr>
<td><strong>(bimanual)</strong></td>
<td>177</td>
<td>165 (26)</td>
<td>108 (30)</td>
</tr>
<tr>
<td><strong>Unimanual tapping (%)</strong></td>
<td>96</td>
<td>94 (10)</td>
<td>95 (13)</td>
</tr>
<tr>
<td><strong>Pegboard</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>(with tapping)</strong></td>
<td>12</td>
<td>15 (2)</td>
<td>10 (2)</td>
</tr>
<tr>
<td><strong>Unimanual pegboard (%)</strong></td>
<td>92</td>
<td>93 (9)</td>
<td>106 (9)</td>
</tr>
<tr>
<td><strong>Tapping</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>(with pegboard)</strong></td>
<td>104</td>
<td>150 (22)</td>
<td>61 (31)</td>
</tr>
<tr>
<td><strong>Unimanual tapping (%)</strong></td>
<td>56</td>
<td>87 (10)</td>
<td>55 (27)</td>
</tr>
</tbody>
</table>

PD data is taken from Brown et al., and data for control subjects from Fuller and Jahanshahi. Data shown is the mean of the right and left hand. PD and control data is expressed as mean (SD).
functioning. His memory for words (RMW 41/50 correct, 10th percentile) and faces (RMF 37/50 correct 5th percentile) were impaired. On the Graded Naming Test he obtained a score (22/30) in the superior range. His visuospatial abilities (VOSP Silhouettes score 25/30) were in the average range. On a relatively easy test of set shifting, the Weigl test, he performed adequately and correctly sorted by both rules.

**DISCUSSION**

Clinical characteristics of parkinsonism caused by lesions of globus pallidus

Bilateral isolated lesions of globus pallidus (GP) are rare. In a large series of basal ganglia lesions reported by Bhatia and Marsden, there were only 15 out of 240 such cases. Only four of these 15 cases (27%) had parkinsonism. One presented with parkinsonism alone, whereas others had additional abulia or abulia associated with either speech disorder or obsessive-compulsive behaviour. In addition to these, there are only a few other cases reported with parkinsonism caused by bilateral pallidal lesions. Hawker and Lang reported a case with severe akinetic rigid syndrome after sudden respiratory arrest and subsequent bilateral pallidal lesions. Feve et al and Bucher et al reported four cases each, and Haaxma et al described one case, who had an additional lesion of the right substantia nigra. After the initial insult, the symptoms of parkinsonism of our case developed after a delay and subsequently progressed. We believe, however, that this was due to his pallidal lesions as the development of delayed onset cases of parkinsonism or other extrapyramidal symptoms after acute insults of varying causes have been well documented before. It is also unlikely that the parkinsonism was due to alcoholism as it has been reported that alcohol withdrawal induced parkinsonism is likely to disappear during relatively short follow up, which was not the case with our patient.

As can be seen from table 3, the most common causes for these lesions were either brain hypoxia or carbon monoxide (CO) poisoning. Regarding the nature of parkinsonism, it appears that, in these published cases, axial motor problems (gait and speech disorder with possible freezing and postural instability) seemed to dominate over brady/akinesia (defined mainly by rapid alternating (distal) limb movements), rigidity, and tremor of the limbs (table 3). This was also the case with our patient, who had a speech problem, “fast micrographia”, and impaired postural reflexes, but no distal limb akinesia or tremor. However, the clinical picture of parkinsonism of our case differed from that of typical PD, as the main clinical findings in PD are rest tremor, distal bradykinesia, and limb rigidity. The case reported by Haaxma et al, who also underwent detailed investigations of motor function, presented with a striking gait disorder, postural instability, and general slowness. It is also notable that most of these cases had lesions affecting both the internal (GPI) and external (GPe) segments of the globus pallidus, or alternatively, the lesions were restricted to GPe only. It has been suggested that, within GP, it is mainly the lesions of the

![](http://jnnp.bmj.com/)

### Table 2

<table>
<thead>
<tr>
<th>Present case</th>
<th>Controls (n = 9)</th>
<th>PD (n = 10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>29 47 (16) 58 (10)</td>
<td>58 (10)</td>
</tr>
<tr>
<td>Squeeze (alone)</td>
<td>158 156 (23) 221 (29)</td>
<td>221 (29)</td>
</tr>
<tr>
<td>Squeeze (simultaneous)</td>
<td>190 156 (27) 309 (77)</td>
<td>309 (77)</td>
</tr>
<tr>
<td>Increase in time (%)</td>
<td>20 10 60</td>
<td>60</td>
</tr>
<tr>
<td>Squeeze (sequential)</td>
<td>182 150 (25) 254 (38)</td>
<td>254 (38)</td>
</tr>
<tr>
<td>Increase in time (%)</td>
<td>4 5 15</td>
<td>15</td>
</tr>
<tr>
<td>Flexion (alone)</td>
<td>209 229 (41) 349 (77)</td>
<td>349 (77)</td>
</tr>
<tr>
<td>Flexion (simultaneous)</td>
<td>226 216 (30) 357 (154)</td>
<td>357 (154)</td>
</tr>
<tr>
<td>Increase in time (%)</td>
<td>8 6 60</td>
<td>60</td>
</tr>
<tr>
<td>Flexion (sequential)</td>
<td>297 244 (35) 445 (110)</td>
<td>445 (110)</td>
</tr>
<tr>
<td>Increase in time (%)</td>
<td>42 7 28</td>
<td>28</td>
</tr>
<tr>
<td>IOL</td>
<td>212 244 (33) 425 (46)</td>
<td>425 (46)</td>
</tr>
</tbody>
</table>

Data for separate (alone) tasks are for the right hand. In the simultaneous and sequential tasks, hand squeeze was performed with the left hand, and elbow flexion with the right hand. In the sequential task, hand squeeze was performed first, followed by elbow flexion. Percentage increase in time values refer to the respective alone values. PD and control data are taken from Benecke et al and expressed as mean (SD). IOL, interonset latency in the sequential task.

Figure 4  Data for movement related cortical potential (MRCP) before self paced voluntary movement.

Figure 5  Contingent negative variation.
Motor performance examined with different experimental tasks

The motor performance of the present case is compared, when available, to the previously published data of normal control subjects and patients with PD.

Preparation for movement

There is general consensus that SRTs are prolonged in PD.7 The present subject had slightly slower reaction times (ITs) in unwarned (mean 456 (SD 73) ms) and warned (mean 344 (11) ms) SRT trials compared to the control group of the same age range (mean age 39 (SD 10) years), who had mean unwarned SRTs of about 430 ms.26 Although his SRTs were slightly faster compared with patients with PD (mean age 64 (SD 8) years), who had mean warned SRTs of about 400 ms,3 the relative slowness of SRTs in our case compared with age matched control subjects was in the same range of magnitude (about 50 ms) to that reported for patients with PD compared with their age matched control subjects.1 This suggests that the unwarned SRTs of our young patient with bilateral lesions of GP were prolonged in a similar fashion as previously reported in PD.1,2 The same was true for unwarned and uncued CRTs, which showed differentially greater slowness than SRTs (fig 2.) in the present case relative to the age matched controls (difference of 94 ms for CRTs compared with difference of 15 ms for SRTs between our case and young controls) and in PD patients compared with their elderly age matched controls (difference of 145 ms for CRTs compared with difference of 81 ms for SRTs between PD and elderly controls).3,33 Although the methods to study simple and choice reaction times used by Haaxma et al33 were somewhat different from ours, their overall results were in line with our case, as both the simple and choice reaction times of their case were slower compared with control subjects.

In healthy controls, SRTs are faster than CRTs34 in part because, due to its invariant nature, in SRT subjects have the possibility to prepare in advance the movement they will have to perform at the time of the go-signal. Our case also benefited from this advance information, as his SRTs were clearly faster than his CRTs (fig 2). This has also been reported in PD.3,34 The magnitude of the CRT-SRT difference in our case (130 ms) was similar to that previously reported in PD (a mean difference of 148 (SD 71) ms), and clearly greater than that reported in controls of the same age range (mean 60 (SD 51) ms), or in controls age matched with PD (mean 86 (SD 70) ms).33 As evident from figure 2 and as noted above, the greater CRT-SRT difference in our case relative to young controls and in PD patients relative to elderly controls is due to the fact that their CRTs show a differentially greater slowness than their SRTs compared with age matched controls. Thus, the current case with bilateral pallidal lesions also shows the differential slowing of CRTs previously found1 in patients with Parkinson’s disease.

Fully cued CRTs provide another way to study the subject’s ability to prepare movements in advance—in particular the time needed to use advance information about the forthcoming movement to speed up response initiation. Healthy control subjects can take advantage of the precue to speed up their CRTs to the level of SRTs, if this information is presented at least 800 ms before the go signal1 (fig 3B). Although PD patients can also benefit from a precue, they seem to be slower in preparing expected precued responses, since they require a longer S1–S2 interval of 3200 ms to do this1 (fig 3C). At only one interval, 1600 ms, was the precued
CRT of our case faster than his warned and uncued CRT, and equivalent to his SRT, but he failed to maintain this benefit with a longer interval of 3200 ms (fig 3A). This suggests that, as with parkinsonian patients (but in contrast to control subjects), our case did not benefit from the extra information about the hand and direction of movement provided by the precue.

Preparation to move can also be investigated by recording the activation of specific brain areas before self-paced voluntary movements. Movement-related cortical potentials (MRCP) have two major components. The early component (earlier BP or NS1) appears 1–2 seconds before movement, and the second component (late BP or NS2) begins about 650 ms before the onset of EMG activity, and rises more steeply than the earlier one. It has been shown that the first component is reduced in patients with PD when assessed off medication.44 As in PD, the MRCPs of our case were abnormal as the first component was absent prior to the movement of either hand; the later component was present. The CNV is a slow negative potential occurring between a warning (S1) and an imperative (S2) stimulus, and it reflects processes related to preparation of a forthcoming movement and anticipation of the imperative stimulus. In PD, the amplitude of the CNV is reduced and correlates with the severity of the disease.2 Our case had no recordable CNV when an S1–S2 interval of 2.5 seconds was used.

Execution of simple and complex movements

In the RT tasks, the movement times (MTs) provide information about the speed of execution of simple aimed movements. In general, the MTs in SRT tasks are faster than in CRT tasks both in healthy controls and PD, but MTs do not differ significantly between precued and uncued CRT conditions.34 Compared with age matched controls, patients with PD have slower MTs.1 On average, in the SRT task, the MTs of the present case were similar to those of young age matched controls.24 In contrast, his MTs in the uncued CRT task were even slower than those of the elderly control subjects.1 In fact our case showed an abnormal pattern of MTs, in that his MTs for the precued CRT task (mean 186.1 ms) were considerably faster (difference of 155 ms) than his MTs for the uncued CRT condition (mean 341.1) and similar to his MTs for the SRT task (mean 187.5 ms). This pattern was not previously seen in either of the young or old control groups or the patients with PD.1,34

Whether the CRT tasks provided fixed start and end points for the movement execution, the movements in the elbow flexion and hand squeeze tasks were self-terminated. Our case was slower in executing aimed movements with a fixed end point as his MTs in the SRT and uncued CRT tasks were slow, although he performed individual elbow flexion and hand squeeze movements at a normal speed. This contrasts with previous findings in parkinsonian patients, in whom it has been suggested that they may have greater difficulties in executing self-terminating movements.45

For patients with PD, simple repetitive movements such as finger tapping and peg placement tasks (involving distal movements) and simultaneous or sequential movements (for example, flex and squeeze or concurrent peg placement and tapping) are more difficult to execute than simple and single movements.2 Furthermore, the more complex the task, the greater the deterioration in the performance of PD patients compared with control subjects, as indicated by a marked drop of the number of taps during simultaneous placing of pegs with the other hand.46 Indeed, this dual task demonstrated one of the most relevant similarity between PD patients and our case (table 1, bottom line). Similarly, as in PD,7 the speed of elbow flexion in our case slowed down in the sequential task (table 2). Compared with PD, there were, however, some differences in simultaneous and sequential flex and squeeze tasks. In contrast to the previously published data in PD,16,47,48 the speed of elbow flexion in our case was faster (and about the same as in middle aged (mean age 47 years, range 31–67 years) control subjects), and it remained the same whether executed on its own or simultaneously with a squeeze of the other hand. In addition, the IOL between the sequential tasks was much shorter than in PD, and slightly shorter than in middle aged control subjects,33—that is, it appeared to be normal. In this regard, the performance of a similar case reported earlier18 resembled PD more closely, as the speed of elbow flexion in that case was slower during both the simultaneous and sequential tasks, and the IOL was about three times longer compared with controls. The reasons for the different results between these two cases with bilateral lesions of GP are not known. One possible explanation could be that whereas our case had isolated GP lesions, the case reported by Haaxma et al17 had an additional lesion in the right substantia nigra. It is also possible that the size and the exact sites of the lesions within GP were different in these two cases.

Cognitive function

Some of the previous studies have suggested that bilateral pallidal lesions may give rise to deficits on tests of executive function and/or passivity, inertia and lack of initiative (abulia).12,25,27 Abulia and apathy can also be seen in longstanding PD.44 However, our patient had no clinical evidence of abulia. Our patient had normal language and visuospatial abilities, but the neuropsychological assessment indicated memory impairment. We believe that the cognitive findings of our case may be a result of the GP lesions, as memory disturbance and deficits of recognition memory in patients with palilidal lesions have been previously reported.25,27 However it is also possible that the memory deficits of our subject may have resulted from the alcohol abuse and possible changes in other brain areas not visible in the MRI.

Comparison with the effects of pallidal surgery in PD

In the light of the present results, it is also interesting to look at the effects of pallidial surgery (unilateral pallidotomy) on the same experimental measures of motor function, and the effects of surgery on the contralateral side are discussed below. Unilateral pallidotomy has been reported to improve (especially ITs) SRTs and CRTs,39,40 although there is one study showing improvement of MTs, but not ITs, in these tasks.41 Interestingly, however, the IT in precued CRT improved after pallidotomy with a pre cue interval of 800 ms, but not with shorter or longer intervals.41 This resembles the results of precued CRT of our case (see above). In addition, repetitive hand movements, such as pronation/supination and finger tapping, and hand dexterity evaluated by pegboard test have been reported to be improved after pallidotomy.39–41 In the study by Limousin et al,37 pallidotomy had no significant effects on repetitive, distal, and fine movements measured by pegboard and finger tapping tasks, whereas it improved movement times in simple and simultaneous elbow flexion tasks. The use of different methods, the exact lesion sites within GP, and the proposed functional somatotopy of GP are likely to explain some of the discrepancies found in the above studies.42–45 However, in most of these studies, pallidotomy seemed to improve many of the experimental motor tasks that are abnormal in patients with PD and our case (compared with age matched controls). This gives indirect evidence that altered firing patterns of GP contribute to the execution of different motor tasks in PD.
What is the functional status of the lesioned GP in our previously healthy subject?

A general assumption is that if a specific nucleus is lesioned, either deliberately or by some disease related event, it becomes functionally silent. However, this is perhaps not the case. Firstly, most of the different measures of motor performance in our case with bilateral GP lesions were similar to those reported previously in patients with PD. Secondly, although there is widespread pathology in the PD brain, and although the principal lesion in PD is the degeneration of the nigrostriatal pathway, it is the resultant abnormal pallidal output that is generally considered to be one of the most crucial factors contributing to the motor symptoms. In PD, in conjunction with being functionally overactive with increased firing rate, the current pathophysiological models suggest that it is perhaps the abnormal firing pattern of the GP that is even more important—that is, the pallidal output becomes “noisy”. Given that the pallidal output is perhaps the only pathologically abnormal factor shared by both our subject and patients with PD, and their similar performance on many tests of motor function, it is tempting to speculate that the functional status of the GP is similar in our case and in patients with PD. This suggests that, instead of being functionally silent, the pallidal output in our case with an acquired pallidal lesion may be “noisy”, as in PD. This idea is supported by a recent report, in which patients with pallidal lesions (who had developed either parkinsonism or dystonia as a result of their lesions) were studied. Using functional MRI, Bucher et al. found that the pallida of these patients were still functionally active despite being lesioned, but direct electrophysiological recordings of these cases would be needed to confirm this. Taking into account the present results, the above findings on pallidal function, and the positive effects of pallidal surgery in PD, it seems possible that whereas the previously normal GP in a healthy person could be damaged and its output made “noisy” by a lesion, pallidotomy could normalise to some extent the “noisy” GP output in PD. It is also possible that if the non-lesioned part of the GP may become functionally unstable (noisy) and contribute to the abnormal output of the GP (together with the lesioned area). Finally, we compared the performance of our case with patients with PD, as we had data from this group. However, these results do not have to be specific to patients with pallidial lesions or PD. Indeed, other extrapyramidal conditions with pallidal pathology could have similar results on the different tasks evaluated.

Note added in proof:
It was recently reported that in experimental conditions in rats, damage to globus pallidus can induce nigral dopaminergic cell death (Wright et al. Eur J Neurosci 2004;20:1737–44).

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