The effect of unilateral posteroventral pallidotomy on the kinematics of the reach to grasp movement

Kerry M B Bennett, John D O’Sullivan, Richard F Peppard, Peter M McNeill, Umberto Castiello

Abstract

Objective—to assess postoperative effects of unilateral posteroventral pallidotomy on the organisation of upper limb movement.

Methods—A three dimensional kinematic system (ELITE, BTS Italy) was used to record reach to grasp movements to objects of either small (0.7 cm) or large (8 cm) diameter placed at a reaching distance of either 20 or 30 cm. Four patients with Parkinson’s disease were assessed in “off” (12 hours without medication) and “on” (1 hour after administration of medication) preoperatively and postoperatively.

Results—Duration of the movement and the time spent in arm deceleration were significantly reduced after surgery. However, movement patterning according to object size was adversely affected. Postoperatively, all four patients showed an abnormal pattern of a longer movement duration, and three showed a longer time of reaching arm deceleration, for reach to grasp movements to the large object than for those to the small object.

Conclusion—Posteroventral pallidotomy seems to be beneficial in reducing bradykinesia of upper limb movements but may have “costs” to movement patterning, particularly for reach to grasp movements to objects of differing sizes. This study raises interesting questions about the role of the globus pallidus interna in coordinating stimulus bound visual information with appropriate motor patterning.

Keywords: Parkinson’s disease; pallidotomy; kinematics; reach; grasp; human; movement

Stereotaxic posteroventral pallidotomy was initially attempted by Leksell in the early 1950s, and is now regaining popularity as a relatively safe and effective neurosurgical technique for selected patients with Parkinson’s disease.1-6 Preoperative and postoperative assessments to determine the effectiveness of pallidotomy have usually been with clinical tests such as the unified Parkinson’s disease rating scale (UPDRS)7 and the core assessment program for intracerebral transplantation (CAPIT8). Most studies have reported significant clinical improvements. Laitenen et al.9 reported some lowering of the classic Parkinsonian signs of bradykinesia, rigidity, and tremor. Dogali et al.9 reported an average improvement of 38% on the CAPIT score and of 68% on the UPDRS score up to 12 months after pallidotomy. Similarly, Baron et al.10 showed an average improvement of 30.1% on the UPDRS after 1 year. Although many studies report a lessening of bradykinesia and improvements to gait,12-14,19,20 a recent study by Johansson et al.21 found no changes to gait patterns or bradykinesia, but significant effects on tremor and dyskinesia induced by medication.

Despite calls from several researchers,12-15 very few studies have assessed the effectiveness of pallidotomy using objective measurement instruments. Bennett et al.6 reported the use of a three dimensional kinematic system to assess the reach to grasp movement of patients with Parkinson’s disease preoperatively and postoperatively in both “off” (12 hours without medication) and “on” (1 hour after administration of medication) states. Preliminary results indicated that the movement was faster (by more than 150 ms) after unilateral posteroventral pallidotomy. Using a computerised target location paradigm, Simuni et al.17 also reported bilateral improvements postoperatively to movement speed, and in movement accuracy (see also Fookson et al.22). Johansson et al.23 conducted an optoelectronic analysis of a movement whereby medicated patients lifted an object from the floor and placed it on a shelf. Posteroventral pallidotomy resulted in no reduction in the time taken to perform this gross multicomponent action; however, more actions were performed per 30 second test sequence postoperatively, suggesting some shortening of the time between each action.

In the current study, a three dimensional kinematic system (ELITE, BTS Italy) was used to assess the effectiveness of surgery on an everyday action. This system allows for accurate measurement in both the temporal (accuracy<10 ms) and spatial (accuracy<0.5 mm) domains. It gives data about movement organisation, including its displacement, velocity and acceleration/deceleration profiles, in three dimensional space. It is thus a valuable tool for providing quantitative accurate measures of the effects of posteroventral pallidotomy on movement organisation.

The investigated movement was that of reaching to grasp an object, a natural everyday movement for which well used neuronal channels are recruited. As such, the learning or subject anxiety effects which may confound more experimental tasks are minimised. The reach to grasp movement has been well characterised by a series of studies beginning with the research of Jeannerod in the early 1980s.19 It is said to
Patients with Parkinson’s disease in early disease stages and “on” medication generally show normal kinematic patterning of movement. Hence, when reaching to grasp an object, apart from the expected generalised slowing, patients with Parkinson’s disease show the characteristic pattern of a longer deceleration phase of arm reaching and an earlier timing of maximum grip aperture for small than for large diameter objects.20–25 When reaching to grasp objects placed at different distances patients with Parkinson’s disease show the characteristic pattern of an increase in the amplitude of peak reaching velocity and a relatively later hand opening peak for longer reaches.20–25

The aim of the current study was to determine if the preoperative organisation pattern of the reach to grasp movement in Parkinson’s disease is altered as a result of posterolateral pallidotomy. For this purpose, intrinsic (size) and extrinsic (distance) properties of the object were manipulated to establish whether both the global parameters, such as movement duration and movement patterning, are affected by surgery. Given the results from previous studies of medicated patients with early stage Parkinson’s disease who show normal movement patterning it was predicted that a lesion to the globus pallidus should not disrupt movement patterning. Further, given brain imaging findings of enhanced activity in cortical areas downstream from the globus pallidus interna, and clinical reports of a decrease in bradykinesia after pallidotomy, it was predicted that the reach to grasp movement should be quicker. The results support only this hypothesis but raise some questions about the role of the basal ganglia, in particular, the globus pallidus interna, in movement organisation.

### Materials and methods

**Patients**

Four patients with Parkinson’s disease underwent unilateral pallidotomy: patient 1, a 71 year old woman with Parkinson’s disease of 21 years’ duration, patient 2, a 54 year old man with Parkinson’s disease of 4 years’ duration, patient 3, a 75 year old woman with Parkinson’s disease of 11 years’ duration, and (4) patient 4, a 63 year old man with Parkinson’s disease of 12 years’ duration. The daily levodopa equivalent dosage (100 mg of regular levodopa=130 mg of controlled release levodopa=10 mg bromocriptine=1 mg pergolide) was 610 mg, 100 mg, 500 mg, and 1245 mg for patients 1, 2, 3, and 4 respectively. For three patients (1, 3, and 4), intervention was suggested because of severe dyskinesia induced by medication, and motor fluctuations. For patient 2, surgery was suggested because of intolerance to levodopa and poor relief of symptomatology with alternative medication. Morning clinical assessments, including CAPIT scores and UPDRS, were performed in the week before, and 1 (patient 1) or 3 (patients 2–4) months postoperatively. These assessments were undertaken in the off state (after 12 hours of withdrawal from parkinsonian medication) and in the best on state, after the morning medication (patients 1 and 2: 100 mg levodopa/25 mg cabergida; patients 3 and 4: 200 mg levodopa/50 mg cabergida). Table 1 shows the score of each patient for each section of the UPDRS in the off state preoperatively. Table 2 gives the values of the dyskinesia score and some CAPIT scores in off and on states for each patient before and after surgery.

No patients showed visual field defects and all had normal or corrected to normal visual acuity. All patients gave informed consent. The project was approved by both hospital and university ethics committees.
SURGICAL PROCEDURE
Surgery took place after withdrawal from Parkinson’s disease related medication for at least 12 hours. The lesioning protocol was based on that used by Dogali et al. With a Cosman-Roberts-Wells (Radionics, Burlington, MA, USA) stereotaxic frame affixed to the patient’s skull, a three dimensional volume MRI data set was acquired (Siemens 1.5 Tesla scanner) and images reconstructed parallel to the line joining the anterior and posterior commissures (AC-PC plane) and at right angles to this. The anatomical target (the most posterior and ventral aspect of the globus pallidum interna) was 2–3 mm anterior to the mid-commissural point, 5–6 mm below the AC-PC line, and 18–21 mm lateral to the midline. The final preoperative target was selected after inspection of the reformatted axial and coronal images with particular reference to the internal capsule and optic tract.
A 10 mm frontal burr hole was made 2–3 cm lateral to the midline positioned so as to make the electrode trajectory 0–20° to the sagittal plane and 60–80° to the axial AC-PC plane. Through a 1.5 mm diameter guide tube extending up to 5 mm into the cortex, an Ohye semi-microelectrode was introduced into the brain and advanced towards the target. Neuronal recording was started 10 mm from the target with signals amplified, passed through an AD converter, and inspected visually and aurally (Viking II Electrodiagnostic System, Nicolet Diagnostic Instruments, Madison WI, USA).

Table 3  Mean (SD) values of selected parameters before surgery in off and on states

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Patient 1</th>
<th>Patient 2</th>
<th>Patient 3</th>
<th>Patient 4</th>
</tr>
</thead>
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<tr>
<td>Movement initiation time (ms):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>518 (98)</td>
<td>524 (63)</td>
<td>411 (95)</td>
<td>363 (82)</td>
</tr>
<tr>
<td>On</td>
<td>518 (148)</td>
<td>528 (74)</td>
<td>401 (61)</td>
<td>321 (62)</td>
</tr>
<tr>
<td>Difference</td>
<td>—</td>
<td>—</td>
<td>—</td>
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<tr>
<td>Movement duration (ms):</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
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<td>1595 (159)</td>
<td>1027 (76)</td>
<td>1230 (104)</td>
<td>1145 (105)</td>
</tr>
<tr>
<td>On</td>
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<td>957 (60)</td>
<td>998 (113)</td>
</tr>
<tr>
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<td>-92</td>
<td>-273</td>
<td>-147</td>
</tr>
<tr>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>41 (8)</td>
<td>38 (5)</td>
<td>39 (5)</td>
<td>41 (5)</td>
</tr>
<tr>
<td>On</td>
<td>41 (6)</td>
<td>41 (4)</td>
<td>45 (4)</td>
<td>47 (5)</td>
</tr>
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<td>—</td>
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<td>+6</td>
</tr>
<tr>
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<td></td>
<td></td>
</tr>
<tr>
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<td>928 (121)</td>
<td>657 (70)</td>
<td>757 (96)</td>
<td>685 (91)</td>
</tr>
<tr>
<td>On</td>
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<td>560 (65)</td>
<td>527 (60)</td>
<td>531 (96)</td>
</tr>
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<td>-77</td>
<td>-231</td>
<td>-155</td>
</tr>
<tr>
<td>Amplitude of maximum grip aperture (mm):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>82 (7)</td>
<td>69 (5)</td>
<td>87 (6)</td>
<td>72 (7)</td>
</tr>
<tr>
<td>On</td>
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<td>72 (4)</td>
<td>76 (5)</td>
<td>60 (3)</td>
</tr>
<tr>
<td>Difference</td>
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<td>—</td>
<td>-11</td>
<td>-12</td>
</tr>
<tr>
<td>Time to peak grip aperture (%):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>70 (8)</td>
<td>77 (9)</td>
<td>70 (8)</td>
<td>71 (7)</td>
</tr>
<tr>
<td>On</td>
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<td>78 (6)</td>
<td>69 (5)</td>
<td>69 (9)</td>
</tr>
<tr>
<td>Difference</td>
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<td>—</td>
<td>—</td>
<td>—</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>238</td>
<td>117</td>
<td>95</td>
<td>94</td>
</tr>
<tr>
<td>On</td>
<td>137</td>
<td>67</td>
<td>74</td>
<td>174</td>
</tr>
<tr>
<td>Difference</td>
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<td>-50</td>
<td>-22</td>
<td>+80</td>
</tr>
<tr>
<td>Variability of time to peak grip aperture (ms):</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Off</td>
<td>125</td>
<td>96</td>
<td>96</td>
<td>79</td>
</tr>
<tr>
<td>On</td>
<td>80</td>
<td>56</td>
<td>47</td>
<td>92</td>
</tr>
<tr>
<td>Difference</td>
<td>-45</td>
<td>-40</td>
<td>-49</td>
<td>+13</td>
</tr>
</tbody>
</table>

Differences are given only when significant t test results were obtained (p<0.01). No tests were performed on the variability values.
USA). Based on characteristic neuronal discharge patterns, the ventral and dorsal margins of the globus pallidus were determined.

The semi-microelectrode was then replaced with a lesioning probe (1 mm diameter, 2 mm exposed tip, Radionics, Burlington, MA, USA) which was lowered under continuous impedance monitoring to 8 mm above the target. Electrical stimulation through the probe (0.2 ms pulse duration, 2, 5, 50, and 100 Hz, 0–10 V) was conducted at 8, 4, and 2 mm dorsal to the target, at the target, and at 2 mm ventral to the target. During stimulation, the face and limbs were observed for muscle contraction and patients were asked to report phosphenes, to determine relation to the internal capsule and optic tract respectively. Up to four trajectories (3, 4, 3, and 1 respectively for patients 1–4) were analysed to determine the optimal target. After a test radiofrequency thermocoagulation lesion at 70°C for 20 seconds, permanent lesions at 80°C for 60 seconds were performed at 2 mm levels from the target to 8 mm dorsal along the chosen trajectory, to produce a cylindrical lesion. Speech, vision, and motor function were monitored throughout the lesioning process and formal visual fields tested postoperatively. Figure 1 shows an example of the lesion site as confirmed postoperatively (at 1–3 days) by MRI.

**KINEMATIC ASSESSMENT PROCEDURE**

The kinematic assessment was performed during morning sessions in both off and on states in the week before surgery and 1–2 months after surgery. The day of these assessments differed from that of the clinical assessment. The off state was 1 hour after the following first dose medication for each subject: patient 1 levodopa/benserazide 25/6.25 mg, levodopa/benserazide (controlled release) 100/25 mg, pergolide 0.375 mg, amantadine 100 mg, selegiline 5 mg; patient 2 pergolide 0.5 mg, amantadine 100 mg; patient 3 levodopa/carbidopa 100/25 mg, selegiline 5 mg; patient 4 levodopa/carbidopa 125/12.5 mg, levodopa/carbidopa (modified release) 200/50 mg, benserazide 10 mg.

Movements were recorded with the ELITE three dimensional kinematic analysis system (B/TIS Italy). Greater detail of this system is given in Castiello et al. The cameras detect infrared reflections of small markers (0.25 cm diameter) attached to the following points of the reaching limb: (a) wrist-radial aspect of the distal styloid process of the radius; (b) index finger-radial side of the nail; and (c) thumb-ulnar side of the nail. Under normal lighting conditions, the participant was seated in front of the table working surface (1×1 m). Before each trial, the right or the left hand was placed on the table in the mid-sagittal plane 15 cm from the thorax. In this position the shoulder was flexed (5–10°), the elbow flexed, the forearm semipronated, and the wrist was in 10–15° of extension. The index finger and thumb were held gently opposed, and the ulnar border of the hand rested on a pressure sensitive starting switch. The target was a perspex cylinder of either small (0.7 cm) or large (8 cm) diameter and 8 cm height placed 20 cm or 30 cm directly in front of the starting switch. Each trial began with an acoustic signal representing the “go” command for the patient to reach and grasp the cylinder. No instructions were given as to the speed of movement or its spatial boundaries. For each size/distance combination the participant performed 15 trials with each hand.

The ELIGRASP (B/TIS, 1994) software package was used to give a three dimensional reconstruction of the marker positions and to filter the data. The transport component was assessed by analysing the trajectory, velocity, and acceleration profiles of the wrist marker. The manipulation component was assessed by analysing the trajectory of each of the hand markers, and the distance between these two markers. Movement initiation time, so called because no emphasis was placed on a rapid response, was taken from release of the starting switch. The end of the movement was taken as the time when the fingers closed on the object and there was no further change in the distance between the index finger and thumb. The dependent variables were temporal measures of the grasp and reach components (see
Unilateral posteroventral pallidotomy on the reach to grasp movement

Table 4  Mean (SD) values of selected parameters in the off state before and after surgery

<table>
<thead>
<tr>
<th>Patient</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
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<tr>
<td>Movement initiation time (ms):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>518 (98)</td>
<td>524 (63)</td>
<td>411 (95)</td>
<td>363 (82)</td>
</tr>
<tr>
<td>After</td>
<td>517 (145)</td>
<td>379 (31)</td>
<td>352 (54)</td>
<td>369 (59)</td>
</tr>
<tr>
<td>Difference</td>
<td></td>
<td>-15</td>
<td>-29</td>
<td>-13</td>
</tr>
<tr>
<td>Movement duration (ms):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>1595 (159)</td>
<td>1027 (76)</td>
<td>1230 (104)</td>
<td>1145 (105)</td>
</tr>
<tr>
<td>After</td>
<td>1335 (147)</td>
<td>818 (56)</td>
<td>1011 (68)</td>
<td>1055 (89)</td>
</tr>
<tr>
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<td>-90</td>
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<tr>
<td>Time to peak reach velocity (%):</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>41 (8)</td>
<td>38 (5)</td>
<td>39 (5)</td>
<td>41 (5)</td>
</tr>
<tr>
<td>After</td>
<td>36 (6)</td>
<td>42 (4)</td>
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<td>48 (6)</td>
</tr>
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<tr>
<td>Reach deceleration time (ms):</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>928 (121)</td>
<td>637 (70)</td>
<td>757 (96)</td>
<td>686 (91)</td>
</tr>
<tr>
<td>After</td>
<td>861 (114)</td>
<td>479 (55)</td>
<td>548 (60)</td>
<td>555 (83)</td>
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<tr>
<td>Difference</td>
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<td>-209</td>
<td>-131</td>
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<td>Amplitude of maximum grip aperture (mm):</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>82 (7)</td>
<td>69 (5)</td>
<td>87 (8)</td>
<td>72 (7)</td>
</tr>
<tr>
<td>After</td>
<td>51 (5)</td>
<td>83 (4)</td>
<td>65 (4)</td>
<td>105 (8)</td>
</tr>
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<td></td>
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<td>70 (8)</td>
<td>71 (7)</td>
</tr>
<tr>
<td>After</td>
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<td>-9</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
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<td>117</td>
<td>95</td>
<td>94</td>
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<tr>
<td>After</td>
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<td>83</td>
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<td>Difference</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Before</td>
<td>125</td>
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</tr>
<tr>
<td>After</td>
<td>110</td>
<td>48</td>
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<td>70</td>
</tr>
<tr>
<td>Difference</td>
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<td>-48</td>
<td>-26</td>
<td>-9</td>
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</tbody>
</table>

Differences are given only when significant t test results were obtained (p<0.01). No tests were performed on the variability values.

Castiello et al. (11) for details). For each parameter, independent t tests were conducted between (a) the preoperative off and on states to determine the effects of medication preoperatively, and (b) the preoperative and postoperative off states to determine the effects of surgery. Throughout the reporting of the results, differences reflect a t test which was significant at an a level of at least 0.01.

Results

Clinical tests

Table 2 shows a summary of the results of clinical tests conducted before and after unilateral pallidotomy. For all patients the score obtained for the motor component of the UPDRS was highest in the off state preoperatively and lowest in the on state postoperatively, with the mean motor UPDRS motor score in the off state improving by 30% after surgery. The stand-walk-sit time value was included to give a general idea of the speed of a gross motor action. This time was greater in the off than in the on state and generally decreased postoperatively. The two point test was included to indicate the results for a gross upper limb movement, of relevance to the movement assessed in the current study. However, results for this test were inconsistent. Patients 2 and 4 showed improvement in both limbs after surgery, patient 1 showed slight worsening in the ipsilesional hand, and patient 3 showed a slight bilateral increase in this time. The medication induced dyskinesia of patients 1, 3, and 4 improved postoperatively (mean preoperative score 7.6, mean postoperative score 6, representing an average 21% improvement), particularly in the contralesional limb.

Kinematic assessment

Preoperative off state v preoperative on state

The patterning of movement before surgery varied greatly across the four patients who participated in this study, and it is because of this heterogeneity that the results for each patient are presented separately.

Reach to grasp kinematics: data collapsed according to object size and target distance

Table 3 shows the mean preoperative values of a selected number of parameters for each subject in the off and on states. Very few showed a common medication effect. One exception was movement duration, which showed a decrease ranging from 92–378 ms after medication. This decrease in movement duration was supported by significantly increased amplitudes of the peaks of arm reaching acceleration, velocity, and deceleration. Another exception was deceleration time, which showed a clear decrease in absolute terms (range 77–231 ms). By contrast, movement initiation time showed no change with medication, and there were no clear pattern of results for any of the transport or manipulation kinematic parameters. For example, patient 1 showed no change to transport parameters with medication whereas patients 2, 3, and 4 showed later temporal settings of peak velocity, acceleration, and deceleration after medication. Similarly, medication affected the amplitude of maximum grip aperture (the maximum distance between the thumb and index finger markers during the opening/closing cycle) in three of four patients, but the direction of this change varied (for example, it was greater for patient 1 but less for patients 3 and 4). Three of the four patients
A comparison of reach to grasp kinematics according to object size (small v large).

Many previous studies have shown that patients with Parkinson’s disease in early disease stages and in the on state demonstrate normal movement patterning with regard to object size.10-25 However, in the current study it was only the first pallidotomy patient (patient 1) who showed the classic small-large patterning in both the on and off states before surgery. Reflecting the greater accuracy requirements, movement duration for this patient was greater for the small than for the large object, and this was supported by lower amplitudes in the peaks of velocity, acceleration, and deceleration for the small object. Similarly, deceleration time (the time taken for the reaching arm to come in on the object) was greater for the small than for the large object.

Figure 2 shows that for the parameters of movement duration and deceleration time, medication served to reduce the small/large differences for patient 1 but a normal patterning was preserved. By contrast, medication preoperatively for patient 2 led to an inappro-
Unilateral posteroventral pallidotomy on the reach to grasp movement

Velocity profiles obtained from reach to grasp movements (patient 2) to the small and large cylinders (A) before and (B) after surgery.

Figure 4

Comparison of reach to grasp kinematics according to object size (small v large)

Surgery clearly affected the kinematics of the transport component according to target object size. Figure 3 illustrates that no patient showed the classic patterning postoperatively. In all four patients postoperatively, movement duration for the small object was less than that for the large object. For patients 1 and 2 this signified a reversal of the normal off preoperative pattern. For patient 4 this signified a worsening of the abnormal patterning and in the case of patient 3 a change to the small/large difference of abnormal patterning. Figure 4 shows an example of velocity profiles obtained from reach-to-grasp movements to the small and large cylinders by patient 2. Before surgery, movement duration was longer for the small than for the large object. After surgery this pattern was reversed.

The postoperative results for deceleration time (time from acoustic signal to release of starting switch) was lower for two patients (2 and 4) postoperatively. Many transport and manipulation component parameters showed no consistent pattern (table 4). For example, peak reach velocity was relatively earlier for patient 1, later for patient 2 and patient 4, and showed no change for patient 3. The amplitude of maximum grip aperture was consistently affected by surgery but the pattern of this change varied across patients.

A comparison of reach to grasp kinematics according to reaching distance (20 cm v 30 cm)

Patterning according to target distance was not changed as a result of surgery. All four patients continued to show the normal patterning of lower amplitudes of the peaks of velocity, acceleration, and deceleration for the 20 cm than for the 30 cm distance. For example, in the case of patient 2, preoperatively the amplitude of the peak of acceleration was 3320 mm/s² for the 20 cm reaching distance and 3723 mm/s² for the 30 cm distance. Postoperatively these values were 5038 mm/s² and 6714 mm/s² respectively (recalling that movements were of lower duration postoperatively). For the manipulation component, all patients showed the normal patterning of an earlier peak grip aperture for the 20 cm than for the 30 cm reach.

Discussion

The main aim of this study was to determine the effects of unilateral posteroventral pallidotomy on motor actions by assessing kinematically the performance of the everyday action of reaching to grasp an object. The pat-
terning of this action has been well characterised by a host of previous research studies, with consistent findings that kinematics differ according to intrinsic (for example, size) and extrinsic (for example, distance) characteristics of the object to be grasped. Hence, the interest in this study was in determining the effect of unilateral pallidotomy on these characteristic movement patternings.

At a global level, the clear surgical benefit was to movement speed and duration. Postoperatively, all four patients with Parkinson’s disease showed lower movement durations and movements which were of greater velocity. For three patients (2, 3, and 4), the reduction was primarily due to decreases in the time taken to hone in on the target object (deceleration time). For example, patient 3 showed a reduction in movement duration of 219 ms postoperatively, with 209 ms of this being to the deceleration phase. Patient 1 was the only patient to show greater time reductions in the accelerative than in the decelerative phase of movement.

The probable explanation for this enhancement of movement speed is that pallidotomy results in easier excitation of the motor association areas. Using PET, it has been demonstrated that the motor association cortex (supplementary motor area and premotor cortex) shows greater movement related activity after pallidotomy. By removing (or decreasing) inhibitory pallidal influences upon thalamocortical circuitry, the association cortex is thought to be more “ready” or “set” for activation. The reduction in movement duration probably reflects the improved time efficiency of the supplementary motor area and premotor cortex in reaching the threshold for activation.

Counter to the benefits to movement speed are the apparent costs to movement patterning according to size. Postoperatively, no patients showed the normal kinematic patterning of longer movement duration and deceleration time for the small than for the large object. In all four patients, the duration of movement was lower, and the period spent in honing in upon the object was lower, for the small than for the large object. A clear confounding effect in this study is that three patients showed abnormality of the size function before unilateral pallidotomy. In itself, this is contrary to the results of many previous studies of patients with early stage Parkinson’s disease in whom no abnormality of movement patterning according to size has been reported. It also increases the difficulty of dissociating pre-existing abnormality from surgical effects. Certainly surgery does not improve or restore the size function. Indeed, the postoperative results indicate that surgery magnifies abnormality in three patients, and detrimentally changes the patterning of the one patient who showed normality preoperatively.

One interpretation of these results is that feedback mechanisms are altered by lesioning the posteroverentral pallidum. The main evidence for this is the inappropriate allocation of deceleration time according to object size. This suggests that visual information about object size is not appropriately relayed to motor pathways during the on line performance, particularly during the deceleration phase. At the neurophysiological level, such an interpretation is difficult to defend because the basal ganglia form part of a medial system which operates largely in a feedforward mode (see Goldberg for review). Lesioning the globus pallidum interna should thus theoretically affect feedforward rather than feedback processes. Yet the accelerative phase of the movement does not show abnormality in the size function postoperatively. Amplitudes of the peaks of arm reaching acceleration and velocity are lower for the small than for the large object. Such a result suggests that the initial part of the movement is patterned correctly, and could be interpreted as reflecting normal feedforward processes.

Many previous studies have shown that patients with Parkinson’s disease become more dependent on visual feedback to guide movement. Goldberg proposed that this could reflect greater reliance upon the responsive, feedback dependent, lateral, premotor system involving the arcuate premotor area and the cerebellum; in other words, the system reverts to alternative and healthier neural channels. With regard to pallidotomy, such a theory would predict no change to this use of feedback mechanisms with lesioning of the globus pallidus interna because the system would still revert to use of the lateral system. By contrast, these results suggest that feedback mechanisms have been disturbed by surgery, and thus point to a role for the basal ganglia in the processing of visual information during the on line performance of a motor action.

There are reasons for exercising caution in adopting the proposal that posteroventral pallidotomy affects feedback mechanisms, or that it promotes greater reliance on feedforward pathways. The first is that only four patients have been investigated, and within this group there was tremendous between patient variability in movement patterning preoperatively. Of note, however, is that the patients of this study showed similar degrees of clinical improvement to those of previous studies. For example, the mean off motor UPDRS score for these four patients demonstrated a 30% improvement—a result which resembles the 30% and 24% improvements reported by Lozano et al and Baron et al, respectively. Secondly, the effects of surgery are very specific. It is only the patterning of the transport (reach) component that shows the dysfunction. The size function of the manipulation component does not seem to be affected postoperatively, and suggestive of normal patterning, the timing of peak grip aperture is earlier for the small than for the large object. On the one hand this lends support to the argument that transport and manipulation are subserved by separate visuomotor pathways, of which only the transport is targeted by unilateral pallidotomy. An alternative explanation is that the patterning of peak grip aperture timing is under feedforward rather than feedback processing. A third reservation against accepting the feedback theory is that distance patterning of both the transport
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and manipulation components is not affected by surgery. Hence, the peak amplitudes of arm reaching acceleration, velocity, and deceleration will lower for a 20 cm than for a 30 cm distance both preoperatively and postoperatively. Again, however, these parameters may be under feedforward control.

In summary, unilateral posteroventral pallidotomy results in quicker reach to grasp actions but disruption to the kinematic organisation of reach parameters with respect to the intrinsic object characteristics of size. These results raise interesting questions about the function of the globus pallidus interna. Supporting much previous research into the function of the basal ganglia, the current study adds evidence for the notion that these nuclei are not responsible for the selection of specific muscles—there is no obvious disruption to the overt performance of the reach to grasp action as a result of surgery. However, the finding of stimulus bound effects on movement kinematics suggests that the globus pallidus interna plays a part in high level sensorimotor functions which couple particular sensory inputs to appropriate movement patterns.38 39 This idea is in line with that proposed by Moore40 who suggested that the basal ganglia are well placed to play a comparator function comparing motor output with feedback.

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