Does bilateral stimulation of the subthalamic nucleus aggravate apathy in Parkinson’s disease?

V Czernecki, B Pillon, J L Houeto, M L Welter, V Mesnage, Y Agid, B Dubois

Objective: High frequency stimulation of the subthalamic nucleus (STN) dramatically decreases motor disability in patients with Parkinson’s disease (PD), but has been reported to aggravate apathy. The aim of this study was to analyse the effect of STN stimulation on motivation and reward sensitivity in a consecutive series of PD patients.

Methods: Apathy and reward sensitivity (Apathy Scale, Stimulus-Reward Learning, Reversal, Extinction, and Gambling tasks) were assessed in 18 PD patients treated by bilateral STN stimulation (“on” and “off” conditions) compared with 23 matched patients undergoing long term treatment with levodopa (“on” and “off” conditions).

Results: Apathy decreased under both STN stimulation and levodopa treatment, whereas explicit and implicit stimulus reward learning was unchanged.

Conclusions: Bilateral STN stimulation in PD patients does not necessarily have a negative effect on motivation and reward sensitivity and can even improve apathy provided patients have been appropriately selected for neurosurgery.

Bilateral high frequency stimulation of the subthalamic nucleus (STN) is an increasingly popular neurosurgical technique that decreases the severity of parkinsonian motor disability and levodopa induced motor complications by 60–80% and the required daily doses of levodopa by 40–80%. It has been proposed that the improvement of motor symptoms results from inhibition of the hyperactivity of the STN, thereby restoring deficient thalamocortical activation of the supplementary motor area during movement. As the motor, associative, and limbic components of the cortical-basal ganglia-cortical loops pass through the STN it is unlikely that the restricted stimulation of such a small target selectively influences the motor component without also affecting the associative and limbic neuronal systems.2 The effects of stimulation of the STN on the associative component include an improvement of psychomotor speed and working memory together with an impairment of the ability to inhibit inappropriate responses.3–5 Stimulation of the STN also influences the cortico-subcortical limbic circuitry,6 as suggested by the appearance7–10 or improvement11−12 of various psychic disturbances.

Apathy, defined as a motivation in affect, cognition and behaviour,13 is a characteristic feature of Parkinson’s disease (PD)14 that decreases under levodopa induced treatment15 but remains unchanged or aggravated under bilateral STN stimulation.16−18 Whether this change in motivation results from a direct effect of STN stimulation or from the decrease in the daily dose of levodopa remains unclear.19 Nevertheless, the occurrence of apathy after neurosurgery may partially offset the benefit of this treatment. The aim of this study was to compare the effects of stimulation of the STN to those of levodopa treatment on apathy and reward sensitivity in PD patients.

PATIENTS

Eighteen consecutive patients treated by bilateral stimulation of the STN were recruited for the study. The neurosurgical procedure was performed as previously described.19 The electrodes were accurately implanted in the STN as shown by post surgery magnetic resonance imaging (MRI). Stimulation was effective because it resulted in a significant improvement in the motor score in the “off-levodopa” state and enabled the dose of levodopa to be significantly decreased (Table 1).

Before surgery, the motor disability score (UPDRS Part III)20 was evaluated in the “off-levodopa” state, as defined by the Core Assessment Program for Surgical Interventional Therapy (CAPSIT).21 Ten months after surgery (mean ± SEM: 10.0 ± 0.9) parkinsonian motor disability and performance on the experimental procedure were evaluated in the following conditions: 1) “off-stimulation” (after stimulation had been switched off for at least 1 hour) and “off-levodopa” (after a night without drug treatment)—that is, deep brain stimulation (DBS) off, drugs off; 2) “on-stimulation” (after stimulation had been switched on for at least 1 hour) and “off-levodopa”—that is, DBS on, drugs off.

Twenty three consecutive levodopa treated patients, most of them being candidates for neurosurgery, were matched to the stimulated patients for age, duration of disease, motor disability, and cognitive functions (Table 1) and tested “on” and “off” levodopa.

None of the patients was demented (score ≤18 on the Mattis Dementia Rating Scale)22 or depressed (score <16 on the Beck or the Montgomery and Asberg depression rating scales).23 24 A “frontal score”,25 modified to rate patients on a 50 point scale,26 included the Modified Wisconsin Card Sorting Test,27 category and phonemic fluencies (animal names and words beginning with M in 60 seconds),28 and graphic and motor series.29 Verbal learning was assessed with a procedure sensitive to the frontal strategic components of episodic memory.30 No significant difference was observed between stimulated and levodopa treated patients.

Abbreviations: DBS, deep brain stimulation; CAPSIT, Core Assessment Program for Surgical Interventional Therapy; MRI, magnetic resonance imaging; PD, Parkinson’s disease; STN, subthalamic nucleus
To compare the effects of STN stimulation and levodopa on apathy and reward sensitivity, the patients were randomly assessed either in the “on” state and a day after in the “off” state or in the “off” state and a day after in the “on” state. They were assessed early in the morning before taking their usual drugs.

The Ethical Committee of the Salpêtrière Hospital approved the study and all subjects gave informed written consent.

METHODS

There were two assessments at a 24 hour interval for each patient. Each assessment lasted about 40 minutes allowing the patients to stay in a stable motor state. For each assessment, the tests were presented in the same order: Apathy Scale, Stimulus Reward Learning 1 and Reversal, Gambling task, Stimulus Reward Learning 2, and Extinction.

Apathy scale

Fourteen questions—for example, “Do you have plans and goals for the future?”—were read by the examiner. For each question, the subject was given four possible answers: “not at all”, “slightly”, “some”, or “a lot”. Scores ranged from 0 to 42, with higher scores indicating more apathy. A score of 14 was used as a pathological cut off level. Subjects were asked to answer according to how they felt at the time of the examination. This scale has been shown to be reliable for the evaluation of apathy in PD.31

Stimulus Reward Learning, Reversal, and Extinction

The tasks were adapted from Rolls et al.32 In Stimulus Reward Learning 1, the subject first learned to touch one of two highly discriminable coloured fractal images that appeared randomly on a video monitor equipped with a touch screen. Different patterns were used for the first and second assessment. The subject gained one point for touching the correct pattern or not touching the incorrect one, and lost one point for not touching the correct pattern or touching the incorrect one. If the pattern was touched, it was immediately replaced by a message telling the subject whether a point had been gained or lost. If the pattern was not touched it disappeared after 7 seconds and was replaced by a message telling the subject whether a point had been gained or lost. A pleasant rising tone also emphasised correct responses, whereas incorrect responses were signalled by a short unpleasant tone. A running total of obtained points was displayed on the screen. The subjects were asked to try to gain as many points as possible. They advanced to each new trial at their own pace, by pressing the space bar on a keyboard, until a criterion of nine correct responses out of ten trials had been reached. The score consisted of the number of trials needed to attain the criterion.

Once the Stimulus Reward Learning criterion had been reached, the Reversal task automatically occurred without warning—the relationship between the patterns and the rewarding or punishing consequences being reversed. Testing continued for 30 trials and further reversals occurred whenever the criterion of nine successive correct responses was reached again. The scores consisted of the number of reversals in 30 trials, the number of trials and the number of errors for the first reversal, the last error trial for the first reversal, and the total number of commission errors (previously correct stimuli touched) and omission errors (previously incorrect stimuli not touched). Reversal has been shown to reflect the ability to shift a mental set on the basis of affective cues (affective shifting) in contrast to tests such as the Wisconsin Card Sorting Test that instead evaluate the ability to shift on the basis of a cognitive process (attentional shifting).33

Stimulus Reward Learning 2, before Extinction, consisted of the same procedure as before Reversal, but with different patterns. After the Stimulus Reward Learning criterion had been reached, the Extinction task automatically took place. In this condition points were won each time the subject refrained from touching one pattern and were lost by touching it. The scores were the number of trials needed, the last error trial, the total number of perseveration errors (previously correct stimuli touched) and omission errors (previously incorrect stimuli not touched). Extinction has been shown to reflect the ability to shift on the basis of a cognitive process (attentional shifting).33

The Gambling task

Bechara et al provided the computerised version of the task.34 The subject sees on the screen four decks of cards labelled A, B, C, and D, each of them being programmed to have 60 cards. Using a mouse, he or she can click on a card from any of the four decks. Every time the subject picks a card, a
Influence of STN stimulation on apathy and reward sensitivity

The apathy score significantly improved under STN stimulation ($F(1,16) = 8.5; p = 0.01$), but the reward sensitivity scores did not change (Table 2). As the state of stimulation and the order of assessment could interact with the performance on Stimulus Reward Learning, Reversal and Extinction, and the Gambling task, we reanalysed these variables, comparing the two subgroups on the first assessment only. No difference in these variables was noted between the “on-stimulation” state and the “off-stimulation” state. Overall, these results show an improvement of apathy under stimulation, but no change of reward sensitivity.

Individual results showed a significant improvement of apathy under stimulation in nine patients, no significant change in eight patients, and a significant aggravation in only one patient. The nine patients who significantly improved differed from the other patients in only two ways: a shorter duration of the disease ($10.4 \pm 1.4$ years versus $15.0 \pm 1.0$ years; $F(1,16) = 6.5; p = 0.02$) and a lower UPDRS score without stimulation and without treatment ($26.3 \pm 4.3$ versus $39.7 \pm 3.4; F(1,16) = 5.9; p = 0.03$). They did not differ from the other patients for the doses of levodopa or of dopaminergic agonists converted in Levodopa Equivalent Doses (Table 3; $6 \text{ mg of ropinirol} = 100 \text{ mg of levodopa}$; $1 \text{ mg of pergolide} = 100 \text{ mg of levodopa}$; $10 \text{ mg of bromocriptine} = 100 \text{ mg of levodopa}$). In the group with improvement of apathy under stimulation, eight patients received dopaminergic agonists (ropinirol for two patients with a mean daily dose of $21.0 \pm 9.0$ mg; pergolide for five patients with a mean daily dose of $1.9 \pm 0.6$ mg; bromocriptine for one patient with a mean daily dose of $40 \pm 0.0$ mg). In the group without improvement of apathy under stimulation, six patients received dopaminergic agonists (ropinirol for three patients with a mean daily dose of $8.3 \pm 2.3$ mg; pergolide for two patients with a mean daily dose of $2.2 \pm 0.7$ mg; bromocriptine for one patient with a mean daily dose of $40 \pm 0.0$ mg). None of the stimulated patients received other drugs.

Relation between severity of apathy and clinical and experimental variables in stimulated patients

At baseline, six stimulated patients (33%) had an apathy score higher than 14—a score considered as pathological. They were significantly more impaired ($p<0.05$) than the other stimulated patients on all variables of Reversal. This was not the case for Stimulus Reward Learning or the Gambling task.

A matrix of correlation showed significant correlations ($p<0.05$) between the severity of apathy and the frontal message is displayed on the screen indicating the amount of money he or she has won or lost. A green bar on the top of the screen also changes according to the amount of money won or lost. The subject is asked to win as much money as possible, and, if he or she cannot, to avoid losing money as much as possible. The experiment shuts off automatically when 100 cards have been selected. The subject must progressively discover that decks A and B are disadvantageous (big gains but bigger losses), whereas decks C and D are advantageous (small gains but even smaller losses). The scores consisted of the number of advantageous choices (C+D) minus disadvantageous choices (A+B) for each of the five blocks of 20 cards and for the total of the 100 cards. Deficits on this task have been shown to be related to lesions or dysfunction of the orbitofrontal cortex and to be independent of working memory deficits, which are related to lesions of the prefrontal dorsolateral cortex.

Data analysis

The same analyses were performed for all variables: 1) analysis of the effects of stimulation using ANOVA with repeated measures (“on-stimulation” state versus “off-stimulation” state); 2) comparison of stimulated patients who improved or not improved under stimulation and of apathetic and non-apathetic patients between groups using ANOVA; 3) comparison of the effects of stimulation with those of levodopa, using ANOVA with repeated measures, with the two groups (stimulated patients versus levodopa treated patients) as a between factor and condition of treatment (“on” treatment [with stimulation or levodopa] versus “off” treatment [without stimulation or levodopa]) as a within factor. To evaluate apathy score changes under stimulation among individual patients each test score was transformed to a standard $z$ score, using total sample baseline means and standard deviation (SD). The clinical criterion of more than 1.0 SD above or below the mean was used to tally improvement or deterioration under stimulation. Correlations were searched for between the severity of apathy and all the cognitive or clinical characteristics of the stimulated patients.

RESULTS

Table 2  Effects of bilateral stimulation of the subthalamic nucleus and levodopa treatment on apathy and reward sensitivity

<table>
<thead>
<tr>
<th></th>
<th>Stimulation</th>
<th>Levodopa</th>
<th>Group (p value)</th>
<th>Treatment (p value)</th>
<th>Inter (p value)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Apathy Scale (total score)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>“on” treatment</td>
<td>11.2 ± 0.9</td>
<td>11.0 ± 1.5</td>
<td>0.91</td>
<td>&lt;0.0001</td>
<td>0.55</td>
</tr>
<tr>
<td>“off” treatment</td>
<td>13.4 ± 1.2</td>
<td>14.0 ± 1.5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2. Stimulus Reward Association</td>
<td></td>
<td></td>
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<tr>
<td>Learning (number of trials)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>“on” treatment</td>
<td>19.2 ± 3.9</td>
<td>22.6 ± 5.6</td>
<td>0.37</td>
<td>0.30</td>
<td>0.81</td>
</tr>
<tr>
<td>“off” treatment</td>
<td>23.1 ± 4.6</td>
<td>28.8 ± 4.8</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Reversal (number in 30 trials)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>“on” treatment</td>
<td>1.6 ± 0.2</td>
<td>1.3 ± 0.2</td>
<td>0.48</td>
<td>0.28</td>
<td>0.43</td>
</tr>
<tr>
<td>“off” treatment</td>
<td>1.3 ± 0.2</td>
<td>1.3 ± 0.2</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Extinction (last error)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>“on” treatment</td>
<td>8.1 ± 1.1</td>
<td>14.2 ± 2.5</td>
<td>0.13</td>
<td>0.99</td>
<td>0.13</td>
</tr>
<tr>
<td>“off” treatment</td>
<td>10.5 ± 1.6</td>
<td>11.8 ± 1.9</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3. Gambling Task (advantageous minus disadvantageous choices)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>“on” treatment</td>
<td>25.4 ± 10.2</td>
<td>13.4 ± 6.9</td>
<td>0.39</td>
<td>0.65</td>
<td>0.56</td>
</tr>
<tr>
<td>“off” treatment</td>
<td>19.4 ± 9.6</td>
<td>14.2 ± 6.4</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Values expressed as mean ± SEM. The ANOVA results are shown as probability levels for the group effect (patients with stimulation versus patients with levodopa), the treatment effect (on treatment/off treatment), and the interaction between these two factors (Inter).
score, the number of perseverations on the Wisconsin Card Sorting test, and the total recall score of the Grober and Buschske test. In contrast, no correlation was found between the severity of apathy and severity of depression, age at disease onset, duration of the disease, UPDRS scores with or without stimulation, or levodopa dose.

**Comparison of the effect of stimulation and that of levodopa**

There was no group effect and no interaction between group and condition (Table 2). The effect of treatment was limited to the apathy scale ($F(1,37) = 18.7; p < 0.0001$). Overall, therefore, the results were similar for the two groups of patients, with an improvement of apathy both under stimulation and levodopa treatment, but no change in reward sensitivity; however, it must be highlighted that the group with levodopa was tested after withdrawal of 982 mg of levodopa (plus dopaminergic agonists) whereas the group with stimulation was tested after withdrawal of 133 mg of levodopa (plus dopaminergic agonists). Unfortunately the doses of dopaminergic agonists or other drugs were not collected for the group with levodopa.

**DISCUSSION**

At baseline (without treatment), 33% of the patients treated by STN stimulation and 39% of the patients treated by levodopa could be considered as apathetic (apathy score $> 14$). These results suggest, therefore, that apathy is not an inevitable consequence of neurosurgery. In addition, apathy improved under stimulation in patients with a moderate disease duration (about 10 years) and moderate disease severity (UPDRS III score of about 26). The improvement of the apathy score was of the same magnitude whether patients were treated by STN stimulation or with levodopa (Table 2). This result is in contradiction with recent studies that showed an aggravation of apathy under STN stimulation. In the latter studies, however, the patients had severe personality disorders before surgery or were older. We can conclude that apathy is improved by both STN stimulation and levodopa treatment provided strict inclusion criteria for neurosurgery are applied.

**Table 3** Treatment of patients whose apathy improved or not improved under stimulation

<table>
<thead>
<tr>
<th></th>
<th>Improved</th>
<th>Not improved</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients</td>
<td>9</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Levodopa (mg/day)</td>
<td>161.1 ± 86.5</td>
<td>105.6 ± 83.5</td>
<td>&gt;0.10</td>
</tr>
<tr>
<td>Agonists (LED mg/day)</td>
<td>230.5 ± 57.2</td>
<td>140.7 ± 47.3</td>
<td>&gt;0.10</td>
</tr>
<tr>
<td>Total treatment (LED mg/day)</td>
<td>391.7 ± 133.1</td>
<td>246.2 ± 81.3</td>
<td>&lt;0.10</td>
</tr>
</tbody>
</table>

Values expressed as mean ± SEM; LED, mean daily dose of levodopa equivalent$^{18, 29}$

**ACKNOWLEDGEMENTS**

INSERM (Institut National de la Santé et de la Recherche Médicale) and Assistance Publique supported the study. We thank A Bechara for providing us with the computerised version of his Gambling task, A M Bonnet and the nurses of the Fédération de Neurologie and Centre d’Investigation Clinique for their contribution. We are also grateful to Leon Tremblay and Mathias Pessiglione for helpful comments, Magali Volteau for statistical advice, and Nikki Horne for revising the English.
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