The metabolic effects of limbic leucotomy in Gilles de la Tourette syndrome

G V Sawle, A J Lees, N F Hymas, D J Brooks, R S J Frackowiak

Abstract
Regional cerebral oxygen metabolism was measured before and after limbic leucotomy in a patient with Gilles de la Tourette syndrome, obsessive compulsive disorder, and obsessional slowness. The preoperative scan showed hypermetabolism in the caudate nuclei, which normalised after operation. It is proposed that the beneficial effects of this operation on both tics and obsessive compulsive behaviour are mediated by disruption of abnormal neural activity in basal ganglia-thalamocortical loops.

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Patients with the Gilles de la Tourette syndrome may also have obsessive compulsive disorder, and indeed genetic studies have suggested that these conditions, together with chronic tic disorder, might share a common hereditary basis.1 A further association with obsessive compulsive disorder is obsessional slowness, a condition where slowness dominates the clinical picture.2 Treatments for Gilles de la Tourette syndrome include behavioural therapy and dopamine blocking agents; clinical response to the latter implying an underlying dopamine receptor supersensitivity.3 Treatments for obsessive compulsive disorder include behavioural therapy, a variety of pharmacological agents, and various forms of frontal leucotomy; clinical response to the latter suggesting an overactivity in frontal cortex-basal ganglia connections. Positron emission tomography (PET) studies using tracers of energy metabolism in these conditions have shown hypermetabolism in the caudate nuclei and frontal cortex.4 5 Studies of the dopaminergic system have been normal.6

Case report
A 45 year old man complained of compulsions to destroy things of value, to harm himself, to twitch and shout obscenities, and to hoard unnecessary objects. At the age of seven he had developed a tic of the head and neck associated with an indistinct vocalisation. At the age of 10 he had an urge to break windows, expose himself, burn money, or destroy his clothes, against his better judgment. When aged 35, he developed an urge to echo things he heard on television, in a crowd, or at the cinema. Daily activities were by now constantly interrupted by compulsions to harm himself, twitch or shout obscenities, leading to marked slowness in dressing, bathing, and eating. He took two hours to get out of bed in the morning after waking because of concern about which clothes he should put on and whether they were clean. His tendency to hoard objects led to his being charged with theft and losing his job as a laboratory technician. On psychiatric examination his mental state was dominated by obsessions and compulsions, but he was not clinically depressed and there were no psychotic features. On neurological examination he had facial tics and coprolalia. He had difficulty initiating movement but once started was able to execute movements fluently. There were no other neurological signs. An MRI scan was normal. Neuropsychological assessments revealed a Weschler Adult Intelligence Score (WAIS) verbal IQ of 108 and a performance IQ of 104.

Although an intense programme of behavioural therapy initially led to a clear reduction in his obsessions and compulsions, follow up 2 months later showed a reversion to his former state. He was treated with a number of drugs including carbamazepine, chlorpromazine, clonazepam, haloperidol, pimozide, procyclidine, propranolol, and sulpiride, in each case without sustained benefit. Despite continued medical treatment he deteriorated further, becoming totally disabled by his tics and compulsions. Following approval by the Mental Health Commission he was referred to the Atkinson Morley's Neurosurgical Unit where he underwent bilateral limbic leucotomy in September 1989. Two pairs of lesions were produced by thermocoagulation, the first in the anterior hypothalamus and the second in the inferior part of the cingulate gyrus. Each lesion was approximately 1 cm in diameter. Postoperative neuropsychological assessments (1 month later) revealed a WAIS verbal IQ of 114 and a performance IQ of 97.

In the first 3 postoperative months he had a considerable reduction in his obsessions.
The metabolic effects of limbic leucotomy in Gilles de la Tourette syndrome

Figure 1-01

Pre-leucotomy
Post-leucotomy

Table Mean (of left and right) cerebral metabolic rate for oxygen consumption (CMRO₂) normalised to global metabolic rate

<table>
<thead>
<tr>
<th>Anatomical region</th>
<th>Mean (SD) CMRO₂ normal (n=6)</th>
<th>Mean (SD) CMRO₂ pre-op</th>
<th>Mean (SD) CMRO₂ post-op</th>
<th>% Change CMRO₂ post-op</th>
</tr>
</thead>
<tbody>
<tr>
<td>Caudate</td>
<td>0.94 (0.08)</td>
<td>1.30</td>
<td>1.05</td>
<td>-19.40</td>
</tr>
<tr>
<td>Putamen</td>
<td>1.06 (0.12)</td>
<td>1.19</td>
<td>1.24</td>
<td>4.70</td>
</tr>
<tr>
<td>Anterior cingulate</td>
<td>0.91 (0.10)</td>
<td>0.94</td>
<td>0.76</td>
<td>-19.10</td>
</tr>
<tr>
<td>Medial frontal cortex</td>
<td>0.97 (0.03)</td>
<td>0.98</td>
<td>0.84</td>
<td>-14.30</td>
</tr>
<tr>
<td>Thalamus</td>
<td>1.01 (0.10)</td>
<td>1.20</td>
<td>1.05</td>
<td>-12.50</td>
</tr>
<tr>
<td>Frontal lobe</td>
<td>0.93 (0.06)</td>
<td>0.93</td>
<td>0.92</td>
<td>-1.50</td>
</tr>
<tr>
<td>Temporal lobe</td>
<td>1.02 (0.05)</td>
<td>1.03</td>
<td>1.05</td>
<td>2.40</td>
</tr>
<tr>
<td>Parietal</td>
<td>1.10 (0.03)</td>
<td>1.05</td>
<td>1.10</td>
<td>5.50</td>
</tr>
<tr>
<td>Occipital lobe</td>
<td>1.04 (0.04)</td>
<td>1.10</td>
<td>1.12</td>
<td>1.60</td>
</tr>
</tbody>
</table>

and by 6 months he subjectively reported an "overall 70% improvement". He was less inclined to tear his clothes but his tics, which were relatively mild preoperatively, were unchanged. By 1 year postoperation he reported a further improvement in his compulsions and by 14 months he had almost no signs of Tourette syndrome, though he was still troubled by urges to tear the buttons from his clothes. By 19 months he was living independently with no clinical signs of Tourette syndrome and he reported a substantial improvement in his obsessions.

Fifteen months before and 21 months after operation the patient underwent measurement of regional cerebral oxygen metabolism by PET using the steady state method with ¹⁸O₂, C¹⁵O₂, and C¹⁴O as tracers. Quantitative data were obtained by applying stereotactically positioned regions of interest to the image sets. The data were normalised to global metabolism and compared with data from six male age matched volunteers. All PET studies were performed under ethical approval and isotopes were administered with permission from the Administration of Radioactive Substances Advisory Committee of the United Kingdom (ARSAC). At the time of his initial PET study (June 1988) he was taking chlorpromazine 150 mg daily and propranolol 40 mg daily. At the time of surgery he was receiving chlorpromazine 100 mg daily only. At the time of his second PET study (June 1991) he was taking chlorpromazine 150 mg daily.

Results

The figure shows selected slices from pre- and postoperative images of oxygen metabolism. Representative slices from the level of the basal ganglia, and the anterior cingulate are shown for each study. The principal change demonstrated is a reduction in the rate of oxygen metabolism in these two structures.

Quantitative data for selected regions of interest are presented in the table. No other region showed such a marked reduction in metabolic rate following leucotomy; indeed many other regions showed a slight increase.

Preoperatively the most striking hypermetabolism was measured in the caudate, although the thalamus was also hypermetabolic. The anterior cingulate, medial frontal cortex, and all whole lobe values were normal. Orbital frontal cortex values were not elevated.

Postoperatively the greatest reductions in measured metabolic rate were in the caudate and the anterior cingulate, with lesser changes in the medial frontal cortex and thalamus. The only whole lobe value to show a fall was the frontal lobe.

Discussion

In considering the pathogenicity of both Gilles de la Tourette syndrome and obsessive compulsive disorder it is pertinent to consider the possibility of defects in the "lateral orbitofrontal" and "limbic" basal ganglia thalamocortical circuits respectively (for review see Alexander, et al.). The lateral orbitofrontal circuit comprises a loop from lateral orbitofrontal cortex via ventromedial caudate to rostromedial substantia nigra reticulata and dorsomedial globus pallidus interna, the latter structures projecting via medial thalamic nuclei back to the lateral orbitofrontal cortex. The limbic circuit is most easily visualised from the perspective of the ventral striatum which receives projections from the anterior cingulate and medial orbitofrontal cortex, as well as from many areas in the temporal lobe. It projects to the ventral pallidum which in turn projects via medial thalamus back to the anterior cingulate and medial orbitofrontal cortex. An aberrant loop to and from the orbitofrontal cortex involving the caudate nucleus, nucleus accumbens, globus pallidus, and dorsomedial nucleus of the thalamus has recently been proposed as a pathogenetic mechanism in obsessive compulsive disorders. The operation of limbic leucotomy selectively targets the anterior cingulate cortex and might therefore be expected to specifically influence the limbic basal ganglia thalamocortical circuit.
This appears to have been successfully achieved in our patient since the caudate (including ventral striatum), anterior cingulate, medial frontal cortex, and thalamus all show a marked reduction in metabolic rate.

Other possible explanations for both the clinical change and metabolic change comprise either behavioural or pharmacological intervention. Neither seems likely, however, as the patient had never shown a sustained response to behavioural therapy, and he was receiving the same dose of chlorpromazine at the time of each PET scan.

Two other recent reports have described patients with Tourette syndrome submitted to the same operative procedure. One patient obtained a sustained resolution of destructive (particularly self-injurious) behaviour together with an improvement in his tics, another derived persistent but modest improvement in ritualistic activities, obsessions, and functional abilities. A third patient, who had delayed speech and motor development since early childhood, in addition to the clinical diagnosis of Tourette syndrome, underwent bilateral cingulotomy twice. After the first operation he showed marked improvement for 10 days only. After repeat operation he showed moderate improvement in ritualistic behaviour. Larger patient numbers have been reported for the treatment of obsessive compulsive disorder using a similar operation, where more than 50% of patients have been reported to achieve a sustained marked improvement.

Several patients with Tourette syndrome have been treated by coagulation of rostral intralaminar and medial thalamic nuclei. In three cases a marked improvement in the frequency of tics was reported, while in others the same operation failed to produce sustained benefit. In our patient the change in thalamic activity following cingulotomy was less marked than the changes in either the cingulate or caudate, nevertheless, it is possible that some of the improvement was directly related to changing neural activity in this structure.

In Gilles de la Tourette syndrome previous PET studies using 18$F$-fluorodeoxyglucose have suggested glucose hypermetabolism in the basal ganglia, while in obsessive compulsive disorder the principal findings (persisting after normalisation to hemisphere values) have been in the frontal lobe. We previously measured regional cerebral oxygen metabolism in a group of patients with obsessional slowness in the context of obsessive compulsive disorder (but none of whom had Tourette syndrome) and found hypermetabolism in the orbital frontal cortex, premotor cortex, and midfrontal cortex. Caudate values were normal. In this Tourette patient the most striking finding was of hypermetabolism in the caudate nuclei, while metabolic rates throughout the frontal cortex were normal. The inter-relationship between Tourette syndrome and obsessive compulsive disorder is complex, patients at either end of an apparently mixed spectrum showing either pure Tourette syndrome or pure obsessive compulsive disorder. Since the patients we previously studied had pure obsessive compulsive disorder and this patient had additional Tourette syndrome, it may be that the different metabolic pattern in this patient is a manifestation of a different neurobiological mechanism for obsessive compulsive disorder in the context of Tourette syndrome.

Other evidence implicating a disorder of the caudate nuclei in obsessive disorders includes the occasional observation of focal striatal MRI changes in obsessive compulsive disorders and evidence of obsessive thoughts and compulsive behaviour in children suffering from Sydenham's chorea; nevertheless the functional bases of Gilles de la Tourette syndrome and obsessive compulsive disorder are unclear. Abnormalities in a number of neurotransmitter systems have been implicated as have second messenger systems such as adenosine 3'5'-monophosphate. Functional imaging studies have demonstrated overactivity of the cingulate gyrus and basal ganglia, while abnormalities in the periaqueductal grey and midbrain tegmentum have been suggested by analogy with encephalitis lethargica and studies on the anatomy of vocalisation. In view of the striking male predominance in Tourette syndrome, a role for sex hormone influences mediated via excitatory neurotransmitter mechanisms has been suggested.

This report underlines the significance of metabolic changes in the basal ganglia in Gilles de la Tourette syndrome, demonstrating a congruence between postoperative changes in metabolic activity and recorded clinical improvement.

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