Parallel Appearance of Compulsive Behaviors and Artistic Creativity in Parkinson’s Disease

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Key Words
Parkinson’s disease · Impulse control disorder · Gambling · Punding · Mesolimbic · Dopamine · Ventral striatum · Creativity

Abstract
A 55-year-old male with idiopathic Parkinson’s disease developed three behavioral changes under combination therapy with selegiline, cabergoline and levodopa. Co-existent behaviors included severe pathological gambling, punding and novel skills in writing poetry (published poetry books). Brain [18F]fluorodopa PET imaging showed decreased tracer uptake in the striatum contralateral to the predominant motor symptoms, consistent with the clinical diagnosis of Parkinson’s disease. Uptake in the ventral striatum was markedly high. Brain MRI before and after behavioral changes showed no pathological findings. The patient was diagnosed as having Parkinson’s disease together with DSM-IV criteria-fulfilling pathological gambling and punding-like stereotyped behavior. There are no established criteria for the classification of emerged artistic creativity, although there are descriptions of the phenomenon in the literature. Inspired by the case, we conducted a preliminary survey— including 290 patients with Parkinson’s disease—exploring the possible relationship between creativity and impulsive-compulsive behaviors. The case, supported by the results of the survey, adds to the cumulative evidence of the association between dopaminergic medication and enhanced creativity, and suggests a possible linkage between increased artistic creativity and impulsive-compulsive behaviors in Parkinson’s disease. Furthermore, it could be speculated that the high mesolimbic dopamine function might relate to the behavioral changes observed in this patient, and is suggestive of the overlapping neurobiological mechanisms of compulsive behaviors and artistic creativity.
Introduction

Over the past decade, it has become evident that Parkinson’s disease and dopamine replacement therapy are associated with the increased frequency of impulse control disorders, such as pathological gambling, compulsive shopping, binge eating and hypersexuality [1]. Other repetitive behaviors, such as punding, walkabout and hobbyism, are also common in Parkinson’s disease patients, and multiple behavioral problems occur frequently [2]. Anecdotal reports of changes in artistic creativity due to antiparkinsonian medication have been previously reported. For instance, a Parkinson’s disease patient was reported to have developed novel poetic writing skills after starting drug treatment [3], and another patient started producing large amounts of artistic drawings [4]. Deep brain stimulation for Parkinson’s disease has induced similar effects in one patient [5], and some patients, who had already shown artistic talent before the diagnosis, have experienced changes in artistic style either due to dopaminergic medication [6] or subthalamic stimulation [7]. It has also been suggested that a former artistic profession in Parkinson’s disease patients could, in fact, be a risk factor for dopamine dysregulation syndrome, which is characterized by addiction to dopamine replacement therapy and stereotyped behaviors, such as punding [8].

Case Report

A 55-year-old right-handed man was diagnosed with Parkinson’s disease in 2001 (onset at the age of 46 years). The first symptoms of the disease were rest tremor in the left hand and slight rigidity in the left upper extremity. Brain MRI at the time of the diagnosis showed no pathological findings. Antiparkinsonian medication was started at the time of the diagnosis. Medication changes over the course of the disease are presented in table 1. The patient had graduated from high school, corresponding to 12 years of formal education, and had various occupations after school including working as a janitor and a survey interviewer, before retiring due to motor symptoms of the disease 2 years after the diagnosis.

Approximately 5 years after the diagnosis, the patient gradually developed a gambling problem mostly for coin-operated slot machines. Before the diagnosis, the patient had no gambling problems and no experience in slot machine gambling. Earlier, while working as an interviewer in casinos, he remembered being ‘surprised’ that some people could get addicted to gambling. At the time of the brain PET imaging in 2010, he played more than 10 h/week on slot machines and spent more than 1,000 €/month on electronic machine gambling. His gambling had caused him significant debts. In 2010, pathological gambling was confirmed according to DSM-IV criteria. Maladaptive gambling behavior was indicated by 7 out of 10 DSM-IV items (cutoff ≥5 items) and the South Oaks Gambling Screen (SOGS) score was 8 out of 20 (cutoff ≥5 points). The questionnaire for impulsive-compulsive disorders in Parkinson’s disease (QUIP) indicated pathological gambling (5 out of 5 items, cutoff ≥2 items), but did not indicate any other impulse control disorders.

Together with slot machine gambling, the patient developed punding-like compulsory behavior, which involved picking wild berries. From the year 2006 onwards, he gradually spent more and more time picking berries. In 2010, the patient saw himself as being obsessed with picking wild berries together with the problematic gambling habit. In the autumn of 2010 alone, he picked hundreds of liters of lingonberries. He described going to the woods early in the morning, losing his sense of time and place (similar to gambling), and stayed in the woods for up to 12 h. While picking, he felt no urge to gamble. He started picking berries before dawn, parking his car next to the woods and using the headlights of his car to find berries. In 2011, due to motor fluctuations and sudden off-periods, he had to regularly interrupt berry picking and return to his car for levodopa dosing.

The patient had shown no interest in arts or poetry before the diagnosis and dopamine replacement therapy, which was confirmed by his family and friends. Seven years after the diagnosis (2 years after gambling and punding started), he started writing poems. He predominantly wrote
early in the morning (3–7 a.m.) before taking morning medications. The themes of his poems were a mixture of love, disease and nature. His poetry was considered interesting by a commercial publisher, and his first poetry book was published 1 year later, and the second book was published in 2010. He showed no interest in other forms of art, such as visual arts or music.

At the time of the PET imaging in 2010, the patient was on Hoehn and Yahr stage 2.0, and his total Unified Parkinson’s Disease Rating Scale (UPDRS) score was 53 [UPDRS I = 2 (out of 16), UPDRS II = 12 (out of 52), UPDRS III = 36 (out of 108), UPDRS IV = 3 (out of 23)]. Blood count, plasma glucose, plasma alanine aminotransferase, plasma gamma-glutamyl transferase, fasting plasma creatinine and plasma carbohydrate-deficient transferrin (CDT) were normal. Urine drug testing for cannabis, cocaine, opiates, phencyclidine, methadone, dextropropoxyphene and benzodiazepines was negative. Brain 1.5T MRI continued to show no pathological findings. The patient underwent a structured interview (SCID-I) by a consultant psychiatrist, which did not reveal any lifetime Axis-I psychiatric disorders or ADHD. The patient scored 10 points (out of 63) in Beck Depression Inventory, indicating that he did not suffer from clinically significant depression. Temperament and character inventory (TCI) scores were 27/40 (27 out of 40) for novelty seeking, 16/35 for harm avoidance, 20/24 for reward dependence, 5/8 for persistence, 36/44 for self-directedness, 34/42 for cooperativeness and 7/33 for self-transcendence. Barratt Impulsiveness Scale (BIS-11) total score was 76, and subscores were 18 (range 8–32) for attentional, 23 (11–44) for motor and 35 (11–44) for non-planning impulsiveness.

Brain [18F]fluorodopa-PET was performed with an ECAT EXACT HR+ scanner in 3D mode, injected dose 168 MBq, 90-min scanning. Patlak analysis with occipital cortex as reference, antiparkinsonian medications had been discontinued for 12 h prior to tracer injection, and carbidopa 150 mg had been given orally 1 h prior to injection, as described previously [9]. Striatal deficit in tracer uptake was seen especially in the putamen contralateral to the predominant symptoms of the disease, which was in line with the diagnosis of idiopathic Parkinson’s disease [right caudate K(t) = 10^{-3} min^{-1}, 7.41, left caudate 9.35, right putamen 2.99, left putamen 4.21, right ventral striatum 10.10, left ventral striatum 10.40, right medial orbitofrontal cortex (mOFC) 2.26, and left mOFC 2.36]. As compared to the other subjects of our previously reported study of Parkinson’s disease patients without known increase in creativity, with or without impulse control disorders, the left [whole group K(t) = 8.40 (6.46–10.40) 10^{-2} min^{-1} and right ventral striatal [8.77 (7.31–10.10)], and the left caudate [7.39 (4.87–9.35)], [18F]fluorodopa uptake of the present patient was the highest of the group (n = 19, one patient with self-reported increase in creativity was excluded from the analysis) [9].

After PET imaging, cabergoline was rapidly stopped due to pathological gambling. Soon after the stopping of cabergoline, pathological gambling behavior decreased without significant changes in punding or poetic creativity. In parallel with cabergoline discontinuation, his motor symptoms increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared. The levodopa dose was therefore markedly increased and restless legs symptoms appeared.

Survey

Is the co-existence of the three behavioral changes coincidental in our patient? The possible linkage between impulsive-compulsive behaviors and artistic creativity in this patient led us to preliminarily investigate the phenomenon in a population of Parkinson’s disease patients. We performed a postal survey inquiring the issue from January to August 2011. The survey was sent to 376 patients from our previous study [10] (patients who had indicated that they were willing to participate in further studies), which investigated the prevalence of impulse control disorders in Finnish patients with Parkinson’s disease. The survey included the QUIP together with questions regarding possible artistic creativity before and after the diagnosis of Parkinson’s disease. A total of 296 (78.7%) of 376 patients returned the survey, but 6 patients were excluded because of inadequate data. Overall, 38.7% (n = 108 of 279) of the patients screened positive for at least one impulse control disorder in QUIP and 19.3% (n = 54 of 280) of the patients reported increased artistic creativity after the diagnosis of Parkinson’s disease. Impulse control disorders as screened with QUIP were significantly more frequent in patients reporting increased creativity after the diagnosis than in patients without creativity increase (54.7 vs. 34.6%, respectively; n = 270, χ² = 7.31, d.f. = 1, p = 0.007). Moreover, 33.3% (18 of the 54) of the patients with an increase in creativity subjectively
linked the enhanced creativity directly to dopamine replacement therapy. The prevalence of creativity
or impulse control disorders did not vary according to the side of predominant motor symptoms or to
the type of medication used by the patient.

Discussion

The clinical symptoms and findings of the patient indicated the diagnosis of
Parkinson’s disease which was later supported by [18F] fluorodopa PET and normal
findings in two brain MRIs, before and after the appearance of behavioral changes. The
diagnosis of pathological gambling was confirmed according to DSM-IV criteria. To
date, there are no generally accepted diagnostic criteria for punding, nor is it defined in
the DSM-IV. However, the continuum of punding to hobbies has been described as
repetitive actions; from simple and meaningless behavioral rituals to those such as
collecting, drawing, cleaning or categorizing, etc., overrunning physiological needs or
social responsibilities, while patients may or may not enjoy the act or retain the insight
of the irrationality of their behavior [11]. In this patient, the excessive berry picking
clearly fulfilled the description above and was classified as punding, although the
behavior was not completely stereotyped simple behavior without meaning.
Furthermore, although the patient enjoyed berry picking, he subjectively felt that
picking was compulsive.

In the context of Parkinson’s disease, artistic productivity/creativity could be more
related to the disease or the treatment than intrinsic artistic motivation. Although the
increase in artistic creativity is multifactorial after the diagnosis of a chronic disease
(including art produced in patient associations; the heightened interest in arts due to
psychological factors after diagnosis and increased free time after premature
retirement), the present data suggests that, irrespective of the cause, artistic creativity
is associated with co-existent problems of impulse control in patients with Parkinson’s
disease. However, it should be noted that a previous study [12] provided contradictory
results compared to ours, which could relate to differences in power to detect
differences (number of patients 36 and 290, respectively), or in methodology as the
previous study included only patients with completely novel artistic productivity,
whereas the present study also included patients with prior artistic activity. It should
also be noted that the survey was a retrospective study and that the positive screen for
impulse control disorders with QUIP is not comparable to clinical diagnosis. We also
lack clinical data to confirm self-reported Parkinson’s disease diagnoses. Therefore, the
results should be interpreted with caution and only as supportive material to the case
described here.

Whether the increase in artistic production of our patient is a reflection of increased
creativity, i.e. something that is novel and meaningful, or merely just an increase in
basal arousal, can be questioned. Creativity has been defined as the ability to produce
work that is at the same time novel and meaningful, as opposed to trivial or bizarre
[13]. The poetry of the patient was accepted twice by a commercial publisher, which
suggests that the artistic level of his work was above average. From a neurobiological
and neuropharmacological perspective, one would expect dopamine-enhancing
therapies to merely increase dopaminergic drive and activation. However, increased
creative drive can secondarily improve creative skill [14]. First, there can be a practice
effect: the more the patient paints, writes or acts, the better the quality of his work.
Another option is that high motivation increases the number of ideas produced, and therefore, a proportional increase in the number of novel and exceptional ideas.

The patients showed markedly high novelty seeking and impulsiveness, as measured with TCI and BIS-11, compared to an earlier report including Parkinson’s disease patients with and without impulse control disorders [15]. In addition, the PET imaging revealed exceptionally high ventral striatal and left caudate fluorodopa uptake suggesting high mesolimbic dopamine function, which has been speculated to be of importance in creativity [14] and impulse control disorders [11].

The patient had been on a maximal dose of cabergoline for 7 years and had suffered from pathological gambling and punding for 5 years before the treatment was stopped because of behavioral problems. As the result of the change in treatment, the urges for gambling decreased without changes in punding or creativity. There is literature to suggest that punding-like repetitive behaviors are particularly associated with levodopa treatment [2], whereas impulse control disorders, such as pathological gambling, are particularly associated with dopamine agonists [1]. Although one is tempted to interpret that, in this case, punding and creativity were related more to treatment with levodopa, and pathological gambling more to cabergoline, impulse control disorders and punding are known to frequently co-exist and there are multiple other factors, apart from medication, that are associated with repetitive behaviors in Parkinson’s disease. If deep brain stimulation is performed for this patient in the future, a reduction in levodopa dose will be possible, and further behavioral changes could emerge.

The increased risk of impulse control disorders in Parkinson’s disease seems to be associated not only with dopaminomimetic medication, but also with several other factors, such as younger age, smoking, family history of gambling problems, and depression [1, 10, 15]. Here we have described a patient with early-onset Parkinson’s disease, who developed an impulse control disorder and punding together with novel poetic creativity. The case and our survey add to the cumulative evidence of the association between dopaminergic medication and enhanced creativity, and suggest a linkage between increased artistic creativity and impulsive-compulsive behaviors in Parkinson’s disease. [18F]Fluorodopa PET imaging revealed markedly high mesolimbic dopamine function, which could relate to the behavioral changes described here, and is suggestive of the overlapping neurobiological mechanism of impulsive-compulsive behaviors and creativity. Our preliminary survey of 290 patients further suggests that the parallel appearance of these behaviors is not an exceptional rarity in Parkinson’s disease. Larger epidemiological and neuroimaging studies are needed to investigate the relevance of these findings, with respect to the clinical characteristics and the neurobiological background of the phenomenon.

Acknowledgements

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Disclosure Statement

Dr. Joutsa has received speaker honoraria from Boehringer Ingelheim. Dr. Martikainen has received speaker honoraria from Boehringer Ingelheim, speaker honoraria and consultant fees from UCB Pharma, and consultant fees from Orion Pharma and GlaxoSmithKline. Dr. Kaasinen has received speaker honoraria and/or travel grants from Orion-Pharma, Abbott, UCB Pharma and Lundbeck; and serves as a member of the advisory board of UCB Pharma and as a consultant for Orion-Pharma and Lundbeck.

Table 1. Changes in the antiparkinsonian medication and the behavior of the patient

<table>
<thead>
<tr>
<th>Time after diagnosis years</th>
<th>Medication, daily dose</th>
<th>Non-motor behavioral changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>Diagnosis of Parkinson's disease, selegiline 5 mg</td>
<td>–</td>
</tr>
<tr>
<td>1</td>
<td>Selegiline 5 mg, cabergoline 1 mg</td>
<td>–</td>
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<tr>
<td>2</td>
<td>Selegiline 5 mg, cabergoline 2 mg</td>
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<tr>
<td>3</td>
<td>Selegiline 5 mg, cabergoline 2 mg, levodopa/carbidopa 100/25 mg</td>
<td>Pathological gambling, punding</td>
</tr>
<tr>
<td>4</td>
<td>Selegiline 10 mg, cabergoline 2 mg, levodopa/carbidopa 200/50 mg</td>
<td>Creativity, pathological gambling, punding</td>
</tr>
<tr>
<td>5</td>
<td>Selegiline 10 mg, cabergoline 2 mg, levodopa/carbidopa 200/50 mg</td>
<td>Creativity, pathological gambling, punding</td>
</tr>
<tr>
<td>6</td>
<td>PET imaging, selegiline 10 mg, cabergoline 2 mg, levodopa/carbidopa 300/75 mg</td>
<td>Creativity, punding</td>
</tr>
<tr>
<td>7</td>
<td>Selegiline 10 mg, pramipexole 0.18 mg, levodopa/carbidopa/entacapone 750/187.5/1,000 mg under consideration for deep brain stimulation</td>
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References


