

Supplementary Table 1 Diagnostic criteria for neuroleptic malignant syndrome

Diagnostic criteria	Scoring	Scores in our patient
Exposure to a dopamine antagonist, or dopamine agonist withdrawal, within the past 72 hours	20	Onset of symptoms 3 days after the addition of quetiapine
Hyperthermia (> 100.4°F or > 38.0°C on at least 2 occasions, measured orally)	18	A temperature of 98°F at presentation
Rigidity	17	Patient had 'lead pipe' rigidity in all 4 limbs
Mental status alteration (reduced/ fluctuating level of consciousness)	13	Clear sensorium
Creatine kinase elevation (at least 4 times the upper limit of normal)	10	Normal levels
Sympathetic nervous system lability, defined as at least 2 of the following: 1. Blood pressure elevation (systolic or diastolic $\geq$ 25% above baseline) 2. Blood pressure fluctuation ($\geq$ 20 mm Hg diastolic change or $\geq$ 25 mm Hg systolic change within 24 hours) 3. Diaphoresis 4. Urinary incontinence	10	Only one criterion, i.e., urinary incontinence, was present

Hypermetabolism, defined as heart-rate increase ($\geq 25\%$ above baseline) AND respiratory-rate increase ($\geq 50\%$ above baseline)	5	Normal
Negative work-up for infectious, toxic, metabolic, or neurologic causes	7	All were negative
A total score of ≥ 74 is required for the diagnosis of neuroleptic malignant syndrome	74	The score of 37 in our patient was well below the threshold, ruling out neuroleptic malignant syndrome

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Supplementary Table 2 Non-motor symptoms of Parkinson's disease

Symptom	Features	Rates	Onset	Course
Olfactory dysfunction	Anosmia/hyposmia	90%	Precedes motor symptoms Present at any stage of PD.	It may be associated with greater severity and rapid progression of PD. Risk factor for developing PD. May predict the onset of dementia in PD.
Visual disturbances	Impaired contrast sensitivity, colour discrimination, convergence, diplopia, dry eye syndrome	22% - 78%	Early-stage feature of PD.	Increase with disease progression.
Pain	Musculoskeletal/nociceptive, radicular/neuropathic, dystonia-related, primary/central PD,	30%-85%	Early-stage feature of PD.	-

	akathisia-related					
Sleep disturbances	Insomnia, fragmentation, awakening, periodic limb movements, restless leg syndrome, daytime hypersomnolence, sleep apnoea	sleep	20%-90%	Precedes symptoms. Mid-stage feature of PD.	motor	Prevalence increases with increasing illness duration.
Rapid eye movement sleep behaviour disorder (REMBD)	Parasomnia characterised by loss of muscle atonia and dream-enactment behaviours	is	25%-50%	Precedes symptoms by 12-14 years. Early-stage feature of PD.	motor	Patients with REMBD have an 33%- 90% risk of eventually developing PD. Associated with more severe PD.
Constipation	Constipation is often associated with a sensation of incomplete evacuation	is often	28%-80%	Precedes symptoms by 20 years in about half of the patients.	motor	Increases the risk of developing PD by 3 to 4.5-fold.

Gastrointestinal dysfunctions	Dysphagia, gastroparesis, small intestinal bacterial overgrowth, bowel dysfunction, hypersalivation	25%-67%	It can precede motor symptoms. Mid-stage feature of PD.	-
Hypotension	Orthostatic hypotension Post-prandial hypotension Nocturnal/labile hypertension	30%-60%	Late-stage feature of PD. 80% patients are asymptomatic.	Related to the duration and severity of PD and the use of dopaminergic medications. More common in patients with dementia, such as those with Lewy Body dementia
Urinary	Increased frequency,	25%-	Late-stage feature of	-

symptoms	urgency & urge incontinence, nocturia, incomplete bladder emptying, retention	50%	PD	
Sexual dysfunction	Men - erectile dysfunction, difficulty reaching orgasm, premature ejaculation Women - low sexual desire, arousal & orgasmic difficulties	53%- 68%	Can precede motor - symptoms Late-stage feature of PD	
Diaphoresis	Paroxysmal drenching sweats	-	-	-
Fatigue	Feelings of tiredness or exhaustion, independent of depression or sleep disturbances	33%- 80% %	It can precede motor symptoms. Early-stage feature of PD.	Not associated with duration or severity of PD & anti-parkinsonian medications.
Psychiatric symptoms	Anxiety, depression, psychosis, apathy, impulse control disorders	50%- 90%	It can precede motor symptoms. It can be present at	More than one disorder in 45% of the patients. Prevalence increases with

				any stage of PD.	increasing duration.	illness
					Depression increases the risk of developing PD.	
Cognitive impairment	Subjective decline, mild impairment, dementia	cognitive cognitive	30%- 80%	It can precede motor symptoms. It can be present at any stage of PD.	Prevalence increases with increasing duration.	illness

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Supplementary Table 3 Psychiatric symptoms associated with Parkinson's disease

	Prevalence	Characteristics	Prodromal phase	Risk factor for PD	Course
Depression	Clinically significant symptoms: 30%-40% Minor depression: ¹ 22% - 37% Dysthymia: 13% -22.5% Major depression: 17%-25%	The most common disorder. Two to threefold more than the general population Comorbid anxiety and depression are found in 50%-80% of patients with PD.	Precedes the onset of motor symptoms in up to 30% of patients with PD by 4-6 years	Increases the risk of developing PD by 2-3 times in the 2-5-year period before diagnosis	Early PD – 10% -27% patients have depression Late PD – 60% have depression
Cognitive impairment	Subjective cognitive decline: ² 25% -40% Mild cognitive impairment: ³ 20%-30% Dementia: point prevalence 24% - 41%	Continuum from SCD to MCI and dementia The point prevalence of dementia is 4-6 times higher than in the general population Types include Alzheimer's, vascular, Lewy body, and Parkinson's disease dementia	Cognitive impairment can be present for decades before the manifestation of motor symptoms	Decline in executive dysfunction and working memory deficits has been reported as a risk factor for PD	Early PD – 10% -20% patients have cognitive deficits The mean time to dementia onset is 10 years. Prevalence at 10 years: 50%-75%; 20 years: 83%
Anxiety	Point prevalence: average 31% (range 25% - 43%)	Anxiety disorder, non-specific: 13% - 30% Generalised anxiety disorder: 11% -14% Panic disorder: 7%-30%	Precedes the onset of motor symptoms in	Anxiety, by itself or associated with depression,	Anxiety is often related to the withdrawal of

Psychosis	Lifetime prevalence: 49%	Social phobia: 14%	up to 30% of patients with PD developing PD by 20 years	dopaminergic medication or to the wearing off of medication effects
	Clinically significant symptoms: 26%	Specific phobia: 13%		
	Psychosis: 25%-40%	Minor/ ill-formed hallucinations include passage (feeling like something is passing at the outer visual field), presence (feeling that someone is close by) hallucinations, and illusions (distorted perception of an actual stimulus). They are typically not disruptive. They occur early, insight is preserved, and there is no cognitive impairment	Psychosis has been reported to occur even in untreated PD in the prodromal phase	Dopaminergic medication can cause psychosis in 20% - 35% patients with PD.
	Complex visual hallucinations: 15% - 40%			
	Minor visual hallucinations: 25% - 50%			
	Non-visual hallucinations: 10% - 35%			
Bipolar disorder	Delusions: 4%	Major/complex/ formed hallucinations are more commonly visual and impairing. They occur about 10 years after onset, insight is lost and cognitive impairment is common		Prevalence of psychosis increases to 60% - 70% after 12 years
	Prevalence: 11%	Patients with BD and PD have increased prevalence of catatonia, depression,	Some patients BD increases the risk of PD by 2-6	Patients with BD and PD have a poorer

(BD)			delusions, cognitive deficits, impulse disorder long times in the 3- 6- course and outcome, control disorders, and somatic symptom before PD onset year period before and treatment disorders.	diagnosis	response.
Apathy	Prevalence: average 40% (range 16% -70%)	Lack of motivation, initiative, and concern, diminished goal-directed activity, reduced interests and emotional display. Overlaps with depression and dementia.	-	-	Early-stage feature of PD. May predict dementia and cognitive decline over time.
Impulse control disorders	Prevalence: 14% -29% of patients with PD have one or more impulse control disorders	Pathological gambling, compulsive buying, pathological sexual behaviour, and compulsive eating.	-	-	Related to the use of dopaminergic medications. Prevalence increases with time.

¹ Dysthymic symptoms not present for 2 full years or more

² Subjective cognitive decline (SCD)- self-perceived decline in cognition with normal performance on tests

³ Mild cognitive impairment (MCI) - decline in cognition associated with subtle deficits on tests and impairment of complex activities of daily living

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