

How to examine a patient with parkinsonism — tips and tricks

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ESSAY

ON THE

SHAKING PALSY.

CHAPTER I.

DEFINITION-HISTORY-ILLUSTRATIVE CASES.

SHAKING PALSY. (Paralysis Agitans.)

Involuntary tremulous motion, with less ened muscular power, in parts not in action and even when supported; with a propensity to bend the trunk forwards, and to pass from a walking to a running pace: the senses and intellects being uninjured.

How to define a patient with Parkinson's disease?

Patient 1

 65 years, mild tremor LUE, slightly changed walking with left leg, worsening of handwriting, feels more stiff and clumsy

Patient 2

• 54 years, several years of hyposmia, constipation, abnormal movements during sleep, hyperechogenic substantia nigra on USG, no parkinsonism

Patient 3

- 47 years, action dystonia left leg, no parkinsonism
- Later abnormal DaT scan, development of PD

What defines Parkinsons disease?

Motor features?

Other clinical symptoms?

DaT scan?

• Genetics?

• Presence of pathological synuclein in body fluids / peripheral tissue biopsies?

REVIEW



MDS Clinical Diagnostic Criteria for Parkinson's Disease

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The first essential criterion is parkinsonism, which is defined as
 <u>bradykinesia</u>, in combination with at least 1 of rest tremor or rigidity.
 Examination of all cardinal manifestations should be carried out as
 described in the MDS–Unified Parkinson Disease Rating Scale

Accuracy of clinical diagnosis of Parkinson disease

A systematic review and meta-analysis

Giovanni Rizzo, MD

ABSTRACT

- 11 studies with pathological confirmation of PD diagnosis
- Non-experts accuracy 73.8%
- Experts accuracy at first visit 79.6%
- Experts accuracy at follow-up 83.9%
- UKPDBB criteria accuracy 82.7%
- Accuracy in community-based study with subsequent pathological confirmation as low as 61.5%
- Most common reasons for misdiagnosis Essential tremor (community studies) and atypical parkinsonism (MSA, PSP) (clinic-based studies)

Classification of parkinsonian syndromes

Idiopathic Parkinson's disease (~80%)

Symptomatic secondary parkinsonism (~10%)

- Drug-induced
- •Toxic M.Wilson, Mn, CO, MPTP
- Traumatic
- Vascular
- Normal pressure hydrocephalus

Other neurodegenerative parkinsonisms (~10%)

- Multiple system atrophy
- Progressive supranuclear palsy
- Corticobasal degeneration
- Lewy body dementia
- •Huntington's disease, etc.

Pathological classification of synucleinopathies

Synucleinopathies	Tauopathies
• Parkinson's disease (+dementia)	Progressive supranuclear palsy
• Lewy body dementia	 Corticobasal degeneration
Multiple system atrophy	• Frontotemporal lobar degeneration

Slowness of movements

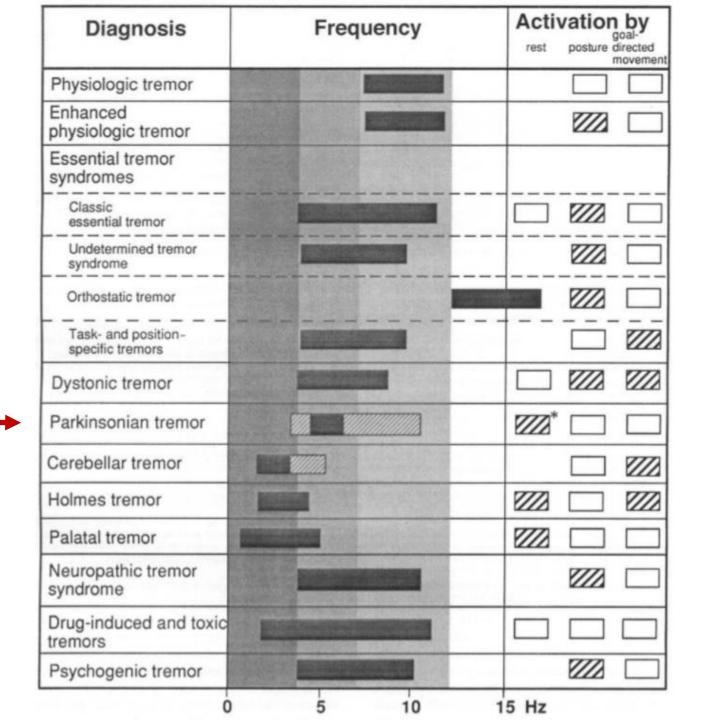
- Bradykinesia
- Paresis
- Dyssynergy related to cerebellar syndrome
- Apraxia/executive dysfunction
- "True bradykinesia" vs. other cause of slowness

Increased muscle tone

- Rigidity
- Spasticity
- Interposition of tremor

Tremor

- Resting vs. action tremor
- Asymetrical vs. Symetrical
- Where?
 - Head more likely ET or dystonic!
 - Chin, lips, tongue PD more likely
- Postural vs. Reemergent -transient tremor supression
- Myoclonus (more likely multiple system atrophy, corticobasal degeneration)



Motor features

Responsive

- Bradykinesia
- Rigidity
- Tremor
- Off freezing
- Off dystonia

Poorly responsive

- Postural instability
- Speech disorders
- Dysphagia
- On freezing
- On dystonia

TABLE 1. MDS Clinical Diagnostic Criteria for PD—Executive Summary/Completion Form

The first essential criterion is parkinsonism, which is defined as bradykinesia, in combination with at least 1 of rest tremor or rigidity. Examination of all cardinal manifestations should be carried out as described in the MDS-Unified Parkinson Disease Rating Scale.³⁰ Once parkinsonism has been diagnosed:

- Diagnosis of Clinically Established PD requires:
 - 2. At least two supportive criteria, and

1. Absence of absolute exclusion criteria

3. No red flags

Diagnosis of Clinically Probable PD requires:

- 1. Absence of absolute exclusion criteria
- Presence of red flags counterbalanced by supportive criteria
 If 1 red flag is present, there must also be at least 1 supportive criterion
 If 2 red flags, at least 2 supportive criteria are needed
 No more than 2 red flags are allowed for this category

Supportive criteria

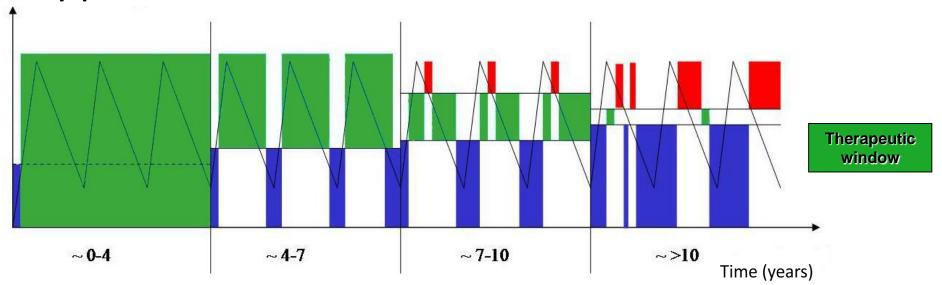
Clear and dramatic beneficial response to dopaminergic
 therapyjednoznačné a dramatické zlepšenie po dopaminergnej liečbe.

- Presence of L-dopa induced dyskinesia
- Rest tremor of a limb documented on clinical examination

Presence of either olfactory loss or cardiac sympatethic denervation on

MIBG SPECT

Typical course of PD



Early stage

Wearing-off

Good response to short off periods dopaminergic treatment

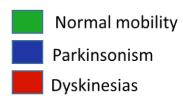
Wearing –off with dyskinesias

Predictable peakof-dose dyskinesias

On-off fluctuations

Unpredictable fluctuations

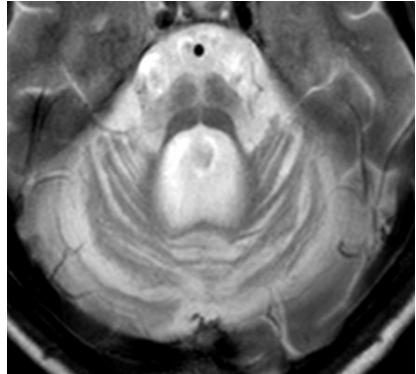
Very narrow therapeutic window

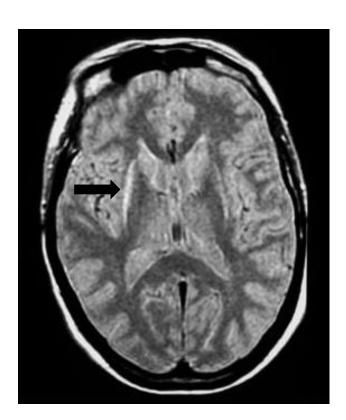


Imaging - MRI

- Standard MRI finding in PD normal "age-related"
- Done to exclude other disorders

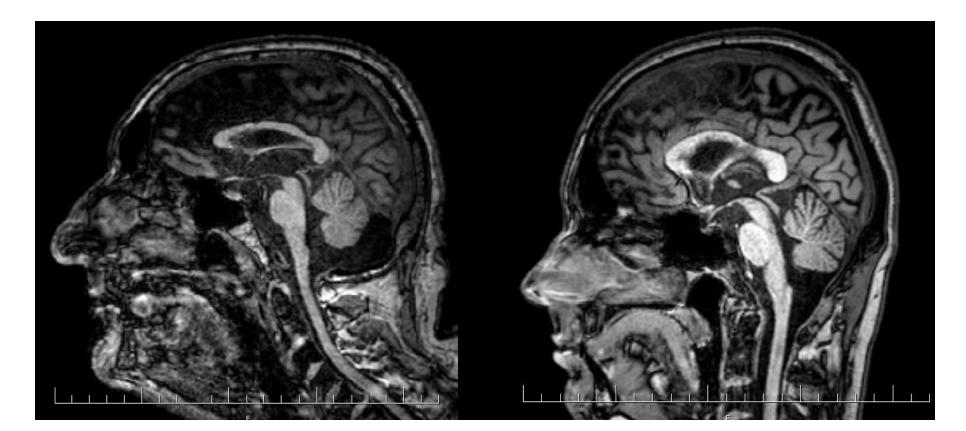






Imaging - MRI

- Standard MRI finding in PD normal "age-related"
- Done rather to exclude other disorders



Imaging – DaT scan

- Does not differentiate PD from other forms of neurodegenerative parkinsonism!
- Differentiates
 - Drug-induced parkinsonism
 - Essential tremor
 - Dystonic tremor
 - Functional parkinsonism
 - Lewy body dementia vs. Alzheimer's disease
- SWEDD scans without evidence of dopaminergic deficit

Imaging – MIBG + IBZM SPECT

• To differentiate PD vs. atypical parkinsonism

- MIBG SPECT cardial sympatethic denervation in PD, not in MSA
 - PD pathological (usually dramatic) reduction of radiotracer uptake
 - MSA, PSP normal radiotracer uptake

- IBZM SPECT examination of postsynaptic D2 receptors
 - Normal uptake in PD (only presynaptic deficits)
 - Reduced uptake in atypical parkinsonism

Therapeutic test

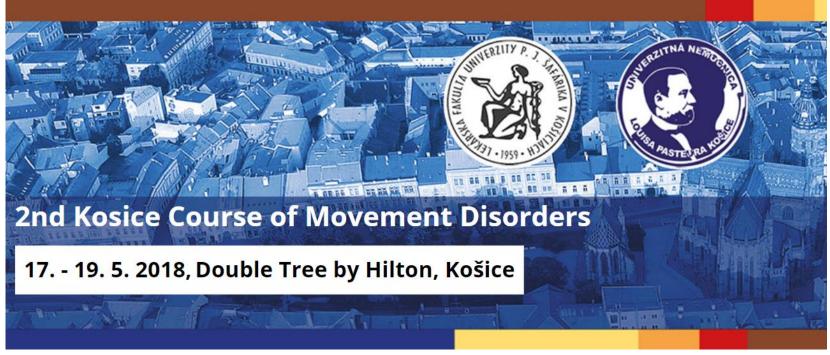
- L-dopa challenge (single dose administration) not reliable
 - Positive might support PD
 - Negative (single dose administration) doesn't mean anything
- Apomorphine unreliable similar to L-dopa challenge
- Necessary to administrate high doses for longer time (~1000mg for at least 1 month, if therapeutic benefit not observed on lower doses)

 Usually no/minimal response to chronic L-dopa administration in atypical parkinsonism (MSA may be an exception in first few years)

Conclusions

• Parkinson's disease has many faces

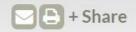
 However, some faces are very atypical and these need to be distinguished...



www.expy-ke.sk

www.movementdisorders.org

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