

Motivation

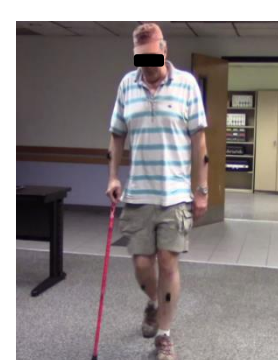
- ◆ Sensor-based technology is needed for implementing personalized therapeutic approaches in Parkinson's disease (PD)¹.
- ◆ Current approaches measure bradykinesia using intermittent standardized UPDRS motor tasks, such as finger tapping (Item 23).
- ◆ Continuous sensor-based monitoring of whole-body bradykinesia during daily activity is important, yet remains to be achieved.

Objective

- ◆ To develop a software platform of wearable sensors and real-time algorithms for automated detection of body bradykinesia during unscripted activities of daily living.

Approach

- ◆ Defined presence and absence of body bradykinesia based on impairments listed in Item 31 of UPDRS.
- ◆ Selected sensors that can provide **both** muscle activity (EMG) and inertial movement (Gyro and Accelerometer) to characterize motor impairments of body bradykinesia.
- ◆ Because body bradykinesia manifests differently during gait, walking and non-walking segments were automatically classified prior to body-bradykinesia detection.
- ◆ Assessed body bradykinesia motor impairments by tracking changes in the magnitude of sensor-based metrics (e.g. reduced limb velocity, reduced limb amplitude and poverty of movement).



- Walking impairments**
- Reduced Arm Swing
 - Reduced Step Length
 - Hesitancy



- Non-walking impairments**
- Reduced Amplitude
 - Reduced Velocity
 - Hesitancy
 - Poverty of Movement

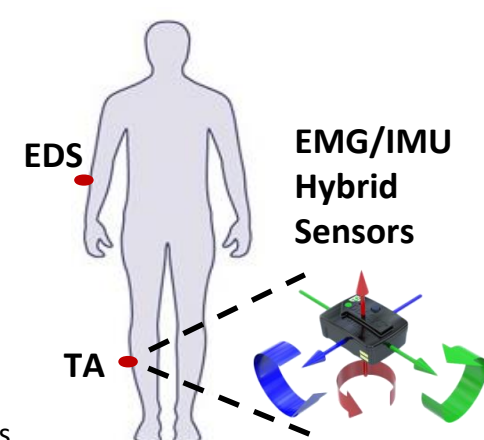
Data Collection

(1) Subject Population

PD Subjects	Training Data*	Testing Data*
Number	n = 8	n = 8
Age (y)	57.5 ± 12.5	63.2 ± 12.1
Male/Female	6/2	6/2
PD Duration (y)	8.6 ± 5.4	5.4 ± 1.7
Total Data	1000 min	1000 min
Prevalence (%)**	58.7	76.4
Hoehn-Yahr (On)	II-III	II-III

* Algorithms trained and tested using different data sets
** % of Total Data w/ bradykinesia

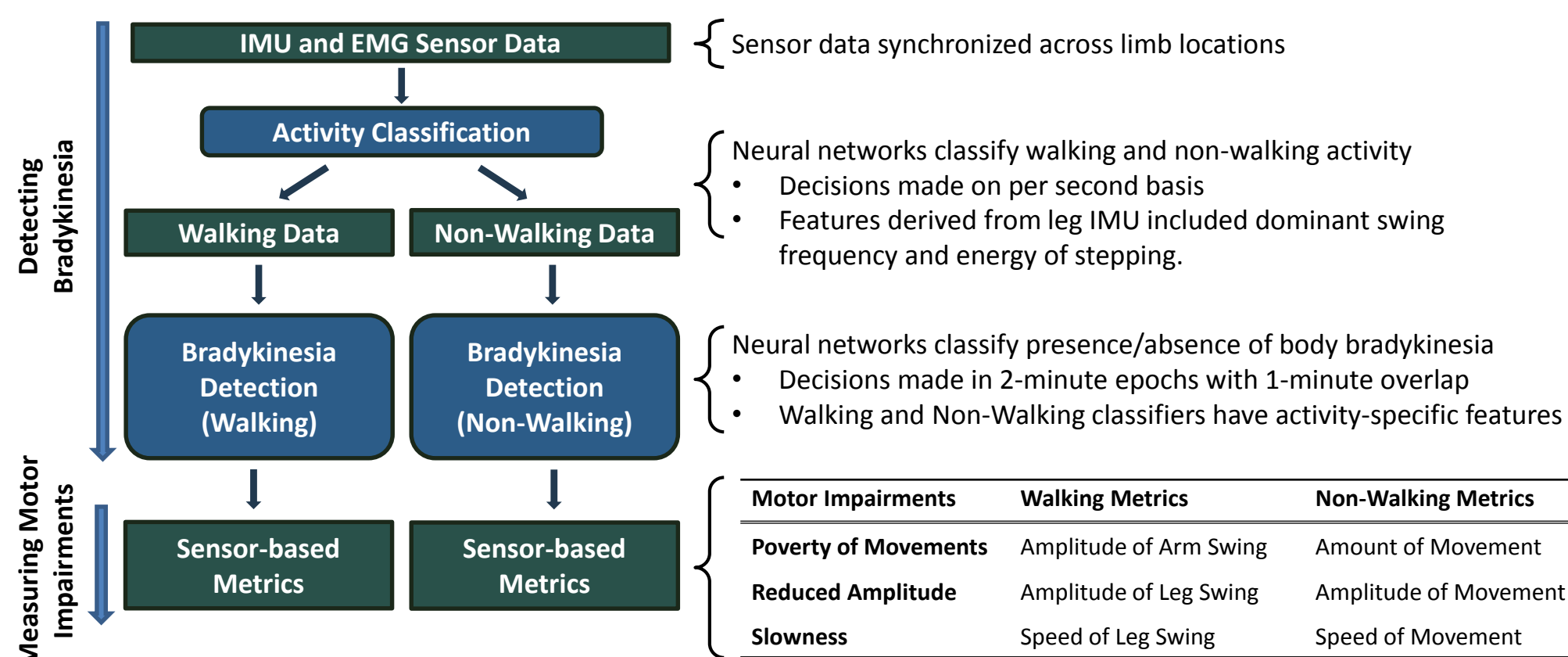
(2) Sensor Placement



(3) Data Acquisition Protocol

- ◆ Trigno™ wireless sensors (Delsys Inc) recorded sEMG and IMU measurements from upper- and lower-limb (see Fig.)
- ◆ Data were recorded continuously during 3 hours of unscripted activity in a simulated home setting.
- ◆ Video recordings were annotated by movement disorder experts to determine the presence/absence of body bradykinesia (based on Item 31 of UPDRS)

Algorithm Design

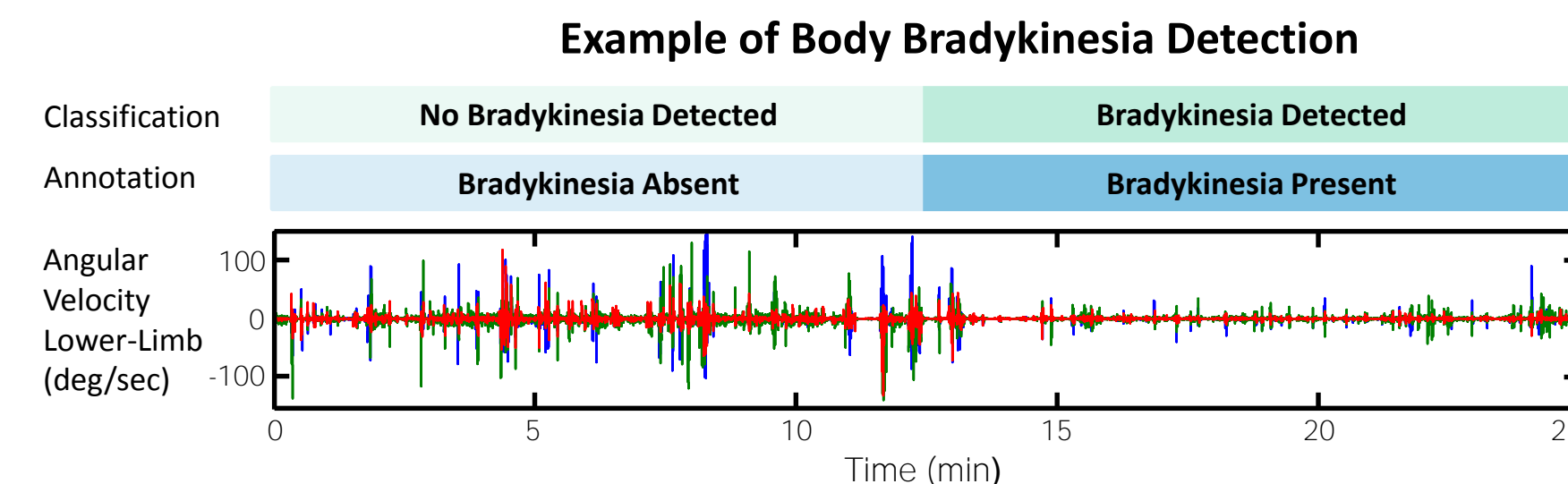


Bradykinesia Classification Results

Accuracy of Body Bradykinesia Detection

- The activity classifier separated walking and non-walking test data with 99.5% accuracy.
- Combined body bradykinesia classification accuracy under all activities was 96.5%.

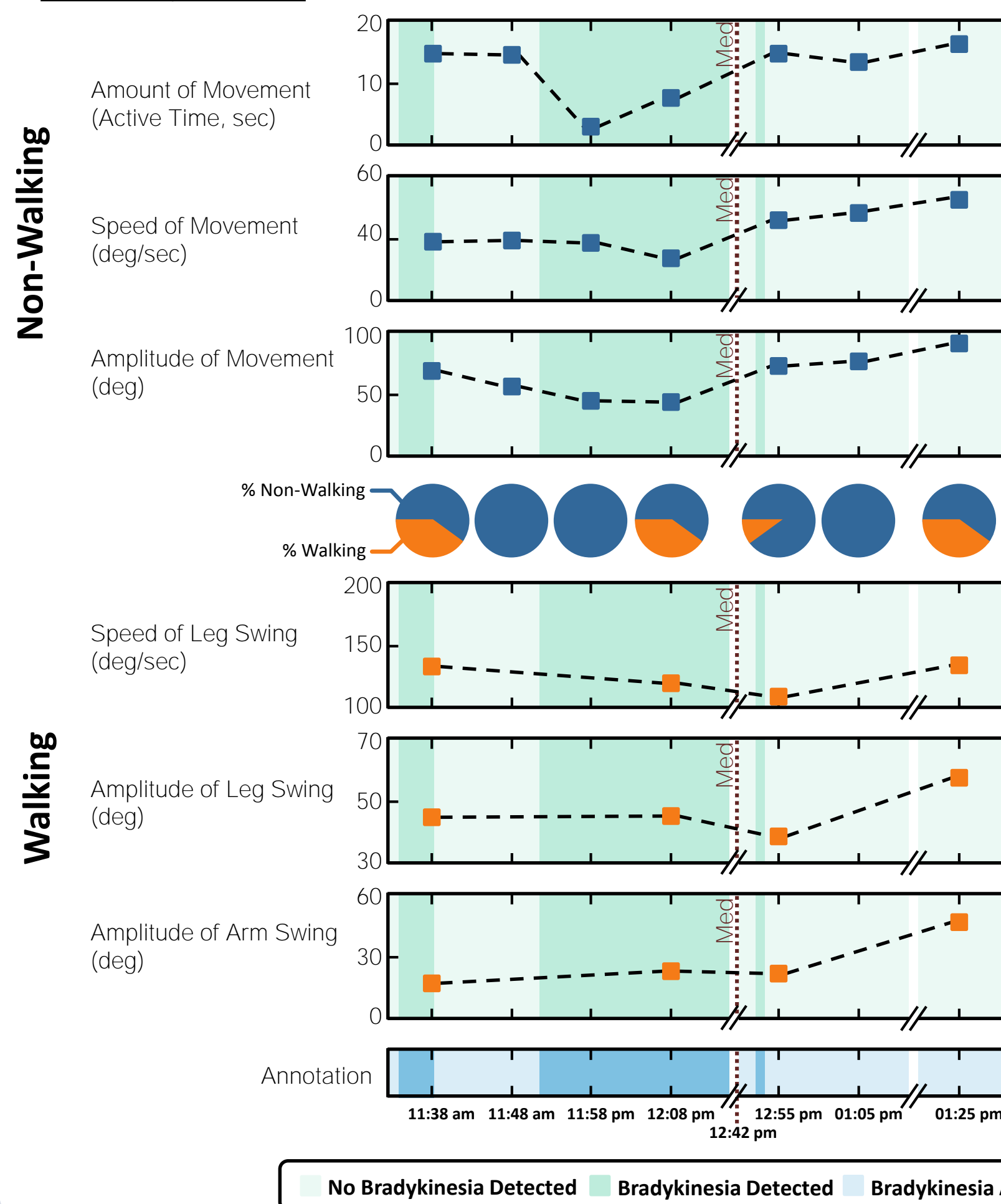
Bradykinesia Classification	Sensitivity (%)	Specificity (%)	Accuracy (%)
Non-Walking	99.4	90.9	97.4
Walking	92.6	100	93.8
Overall			96.5



Results for Tracking the Effects of Dopamine Replacement on Body Bradykinesia

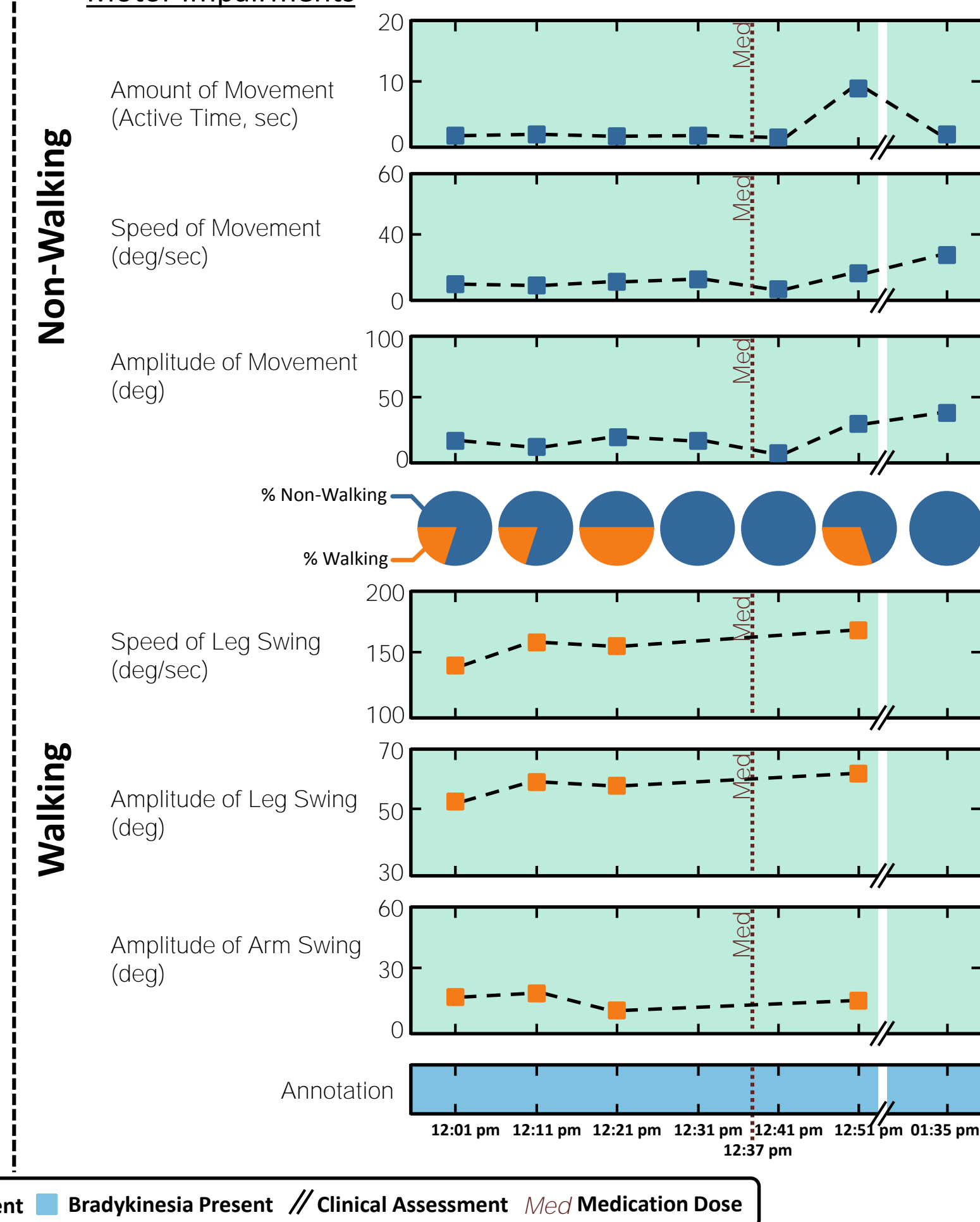
Case 1: This patient responds to medication intake with a change in body bradykinesia assessment that coincides with improvements in motor impairments.

Motor Impairments



Case 2: This patient responds to medication intake with no changes in body bradykinesia assessment but he has improvements in motor impairments during non-walking which can inform the clinician of changes in body bradykinesia severity.

Motor Impairments



Conclusion

First achievement of a wearable system that is used during normal daily activities to provide:

- Body Bradykinesia classification with <5% error
- Objective metrics to quantify motor impairments associated with body bradykinesia

This proof-of-concept tool provides assessments and metrics that are responsive to PD medication.

Acknowledgments

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References

¹Movement Disorders, Vol 31, No 9 2016.