Towards Ambulatory Motor Monitoring: Measuring Dyskinesia During Activities of Daily Living

Webinar Will Begin at 12:00 PM EDT



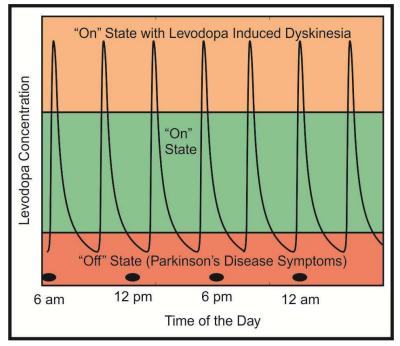
Outline

- Motor Fluctuations and Levodopa-induced Dyskinesia
- Challenges with Clinical Dyskinesia Assessment
- Intelligent Algorithms for Continuous Monitoring of Dyskinesia



Fluctuations and Dyskinesia

- Motor Fluctuations
 - Alternate between "OFF" and "ON" states over dose cycle
- Levodopa-induced
 Dyskinesia
 - Involuntary, episodic, and irregular movements
 - Most commonly occur at peak dose



Chronic Stages of Levodopa Therapy

Keijsers et al., Movement Disorders 18(1), 2003



Patient Impact



https://www.youtube.com/watch?v=CaJymwziF-M



New Therapy Development

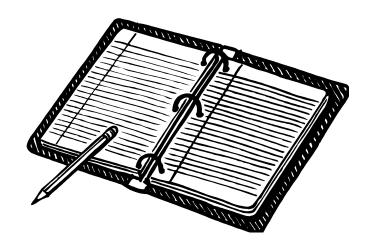
	THE MICHAEL J. FOX FO		
	OUR ROLE & IMPACT BLOG	UNDERSTANDING PARKINSON'S GET INVOLVED	
	HOME FOR RESEARCHERS FAPPLY FOR A GRANT S	2014	
	nicalTrials.gov ce of the U.S. National Institutes of Health	Example: "Heart attack" AND "Los Angeles" Search for studies: Advanced Search Help Studies by Topic Glossie	
Find	Studies - About Clinical Studies -	Submit Studies - Resources - About This Site -	
Home	> Find Studies > Search Results	Text Size	a 🔻
	407 studie	es found for: dyskinesia and parkinson's	
	PROUNAINI UUAL		
	이 같은 것 같은	Research (MJFF) wishes to engage researchers and drug tically treating levodopa-induced dyskinesia (LID). Applicants	

may submit a proposal focused on a relevant area of LID research (see full details below) for consideration of a one-year, \$125,000 award. See the Special Requirements section below for additional information.

GREATLAKES NEUROTECHNOLOGIES

Clinical Assessment of Dyskinesia





In-Clinic Assessment

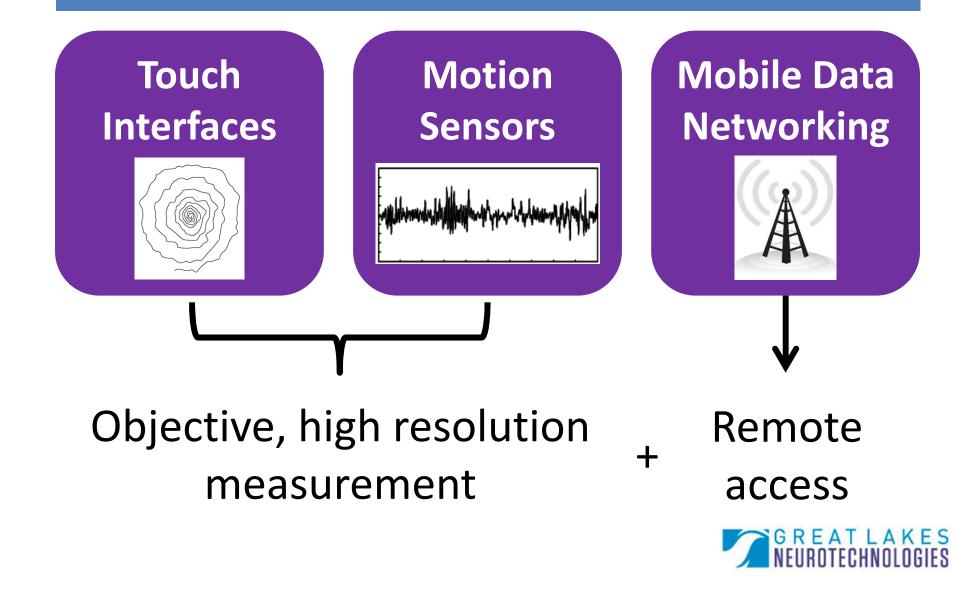
 Rating scales only provide a temporal snapshot of dyskinesia response, limited resolution

Patient Diaries

 Self-assessment at home at regular intervals, confounded by patient awareness, compliance

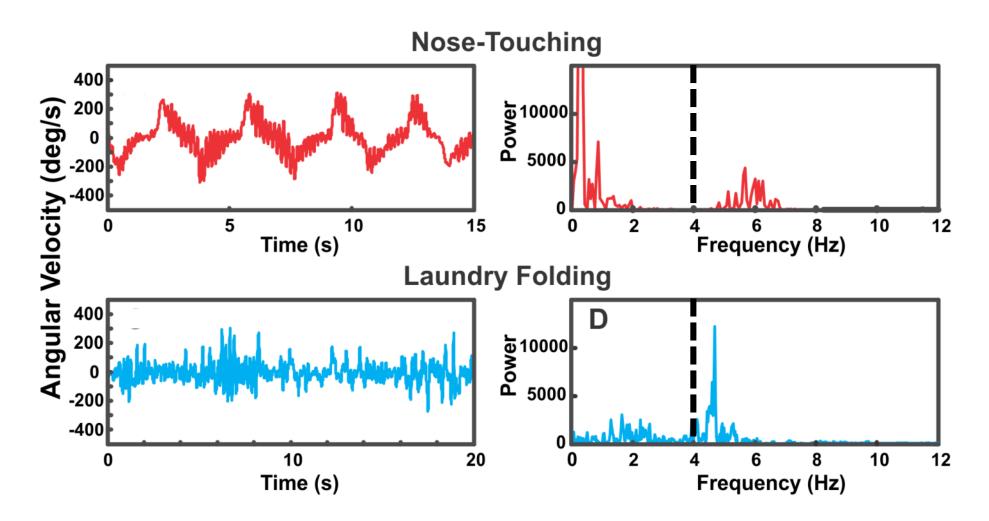


Technology-based Assessment



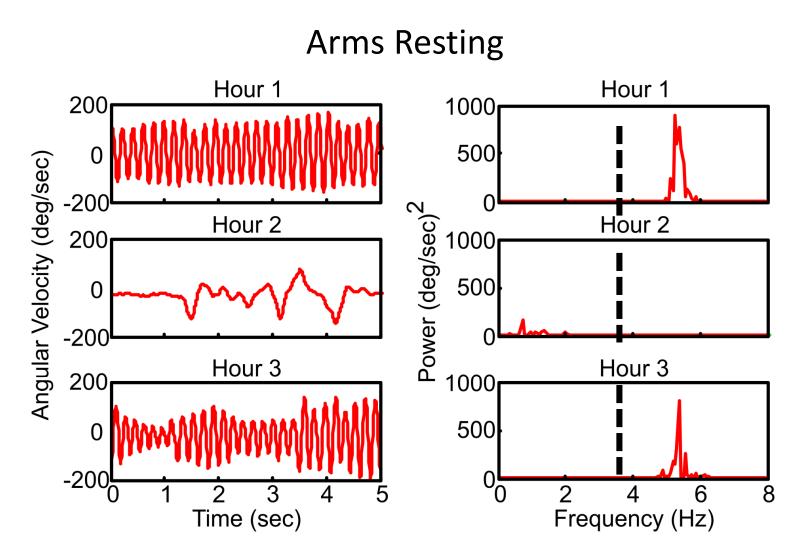


What exactly do motion sensors capture?



Tremor can be differentiated from voluntary motion by taking advantage of separation in the frequency spectrum

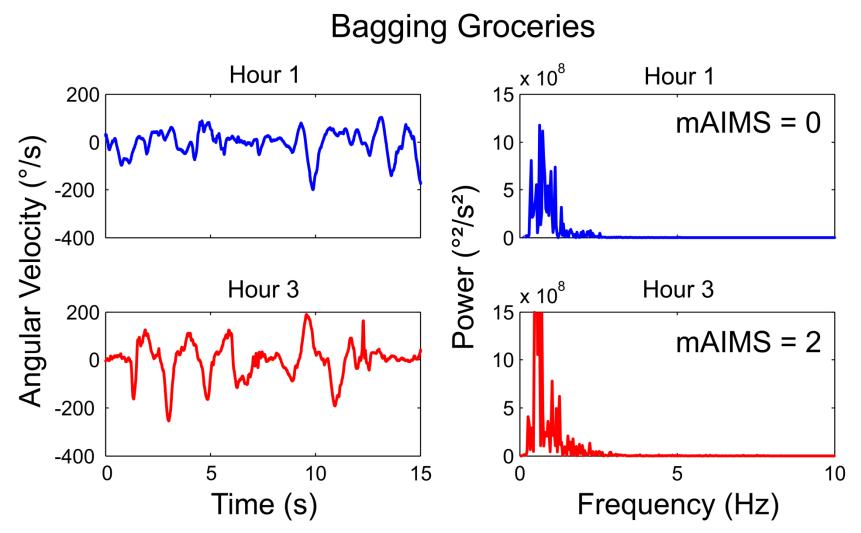




Same principles can be used to quantify dyskinesia when there is no voluntary motion

Mera et al., Journal of Parkinson's Disease 3(3), 2013

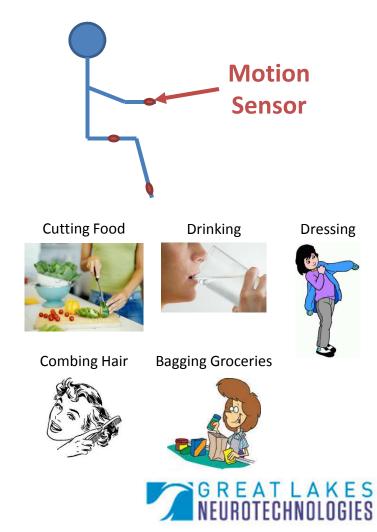




Quantifying dyskinesia during routine activities is significantly more challenging because of kinematic and spectral overlap

Motion Sensor Dyskinesia Quantification During ADLs

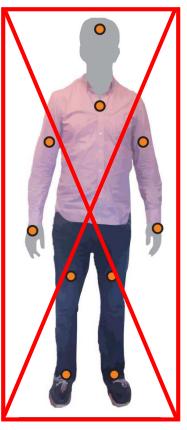
- Motion sensor units positioned on hand, thigh, and heel
- Representative scripted ADLs performed over a 3-hr period after levodopa dose
- Motion sensor data saved, videos scored by blinded raters using m-AIMS





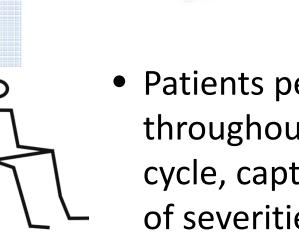
Goals

- Develop an intelligent algorithm that can rate dyskinesia severity across a range of routine activities
- Determine a minimal set of motion sensors to minimize patient burden



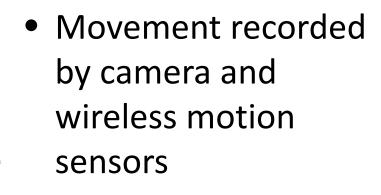






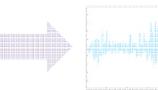
• Patients perform tasks throughout dose cycle, capturing range of severities





Computational Model



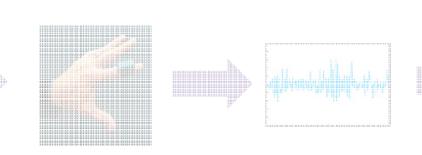






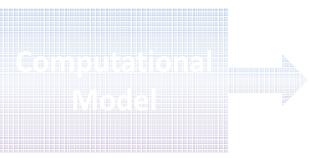
 Blinded clinicians score dyskinesia using modified AIMS scale

()snotsturno Model /

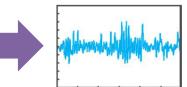




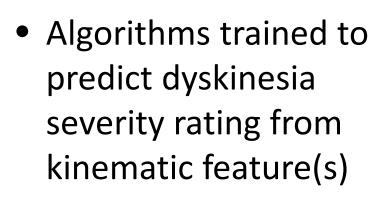
 Kinematic features extracted from motion data using signal processing techniques











Computational Model

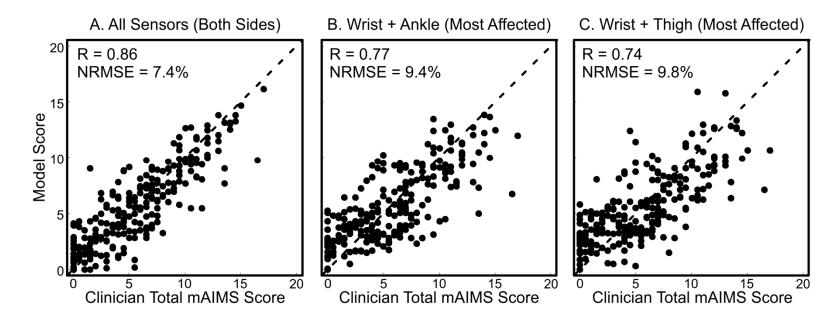


 Process repeated using different combinations of sensors and locations

Computational Model



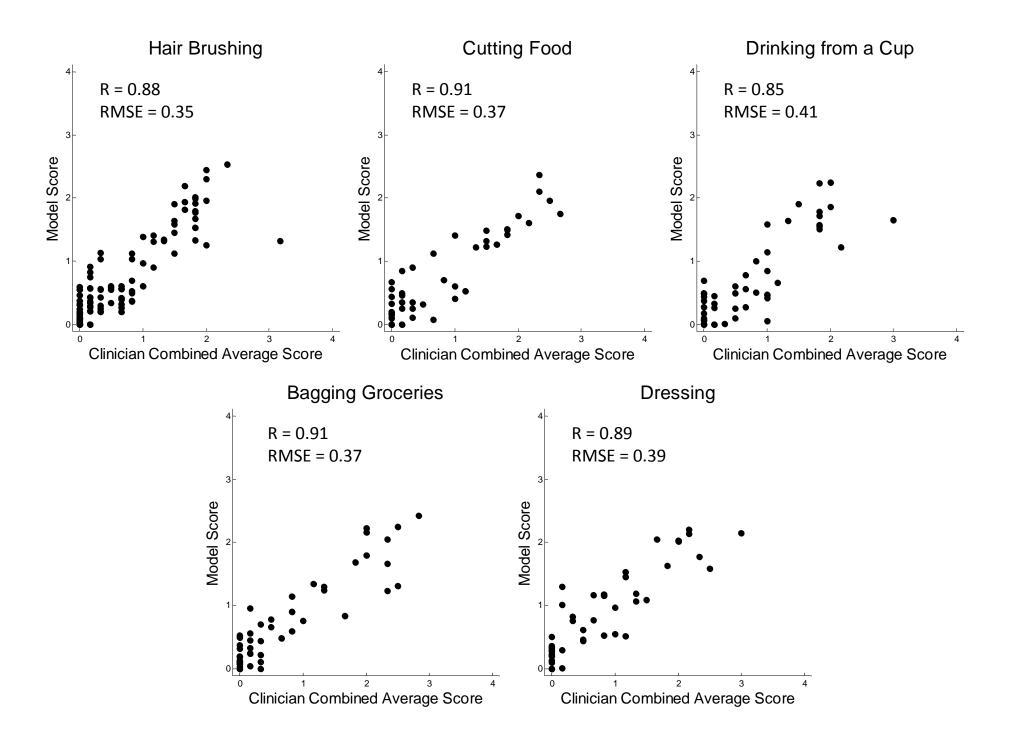
Severity Scoring Model



 Linear regression models were developed to predict total mAIMS score averaged across both raters using kinematic features as inputs



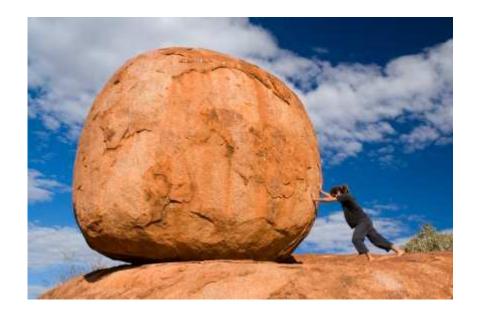




Conclusions

- A motion sensor system can accurately capture dyskinesia during routine activities
 - Provide an objective tool for quantifying motor symptom fluctuation in the context of daily life
- Ongoing study to validate algorithms in ambulatory setting





Current Commercial Technology

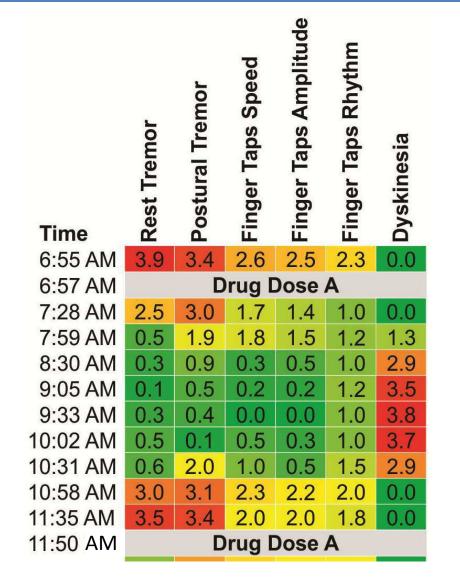




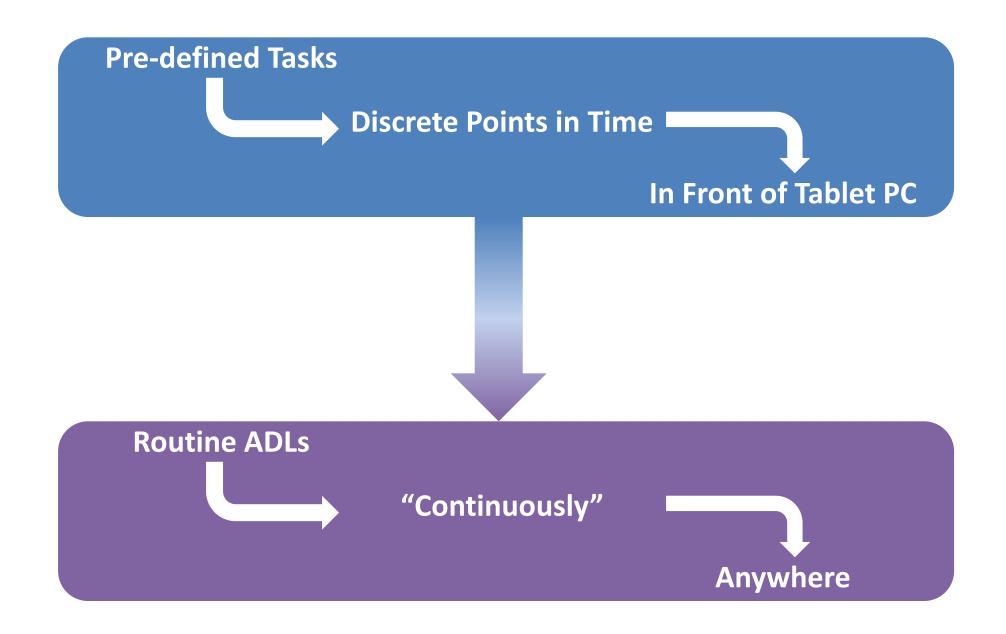




Capturing Fluctuations









Upcoming Features

 Moving towards system that can easily provide objective measures of medication state and physical mobility with minimal patient burden through continuous monitoring





Acknowledgements



Michelle Burack, MD, PhD E. Ray Dorsey, MD, MBA



Dustin Heldman, PhD Joseph Giuffrida, PhD Thomas Mera, MS



Zoltan Mari, MD

NIH/NINDS 7R43NS071882 NIH/NIA 9R44AG044293

Questions?

Please contact: Christopher Pulliam, PhD Senior Biomedical Engineering Researcher cpulliam@glneurotech.com

