## UNIVERSITY OF ILLINOIS ATURBANA - CHAMPAIGN

JAN 13 202 UIUC OPRS

Department of Kinesiology & Community Health



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December 3th, 2017.

## MEDICAL RELEASE

To the neurologist of	B
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Your patient has expressed interest in participating in a study investigating the effects of a training protocol for voluntary eye movements in Parkinson's Disease. We are including participants with unilateral involvement only, unilateral and axial involvement, or bilateral involvement without impairment of balance. Outcome measures include saccade latency, saccade duration, and saccade amplitude in reflexive and voluntary saccades. The protocol involves a large TV screen that will display visual targets arranged in circles of varying radii to train saccades up to 50 degrees of the participant's field of view. Participants will be asked to make approximately 120 saccades over the course of 30 min. There will be a total of eight training sessions. For participants with Parkinson's disease, pre and post assessments will be taken in an "ON" and "OFF medication state". For the "OFF-state", participants will be asked to start assessments at the time of scheduled medication intake but withhold from taking the medication for two hours. During the 2-hour testing session, in the "OFF-state", participants will perform an eye tracking assessment and the Movement Disorders Society-United Parkinson's Disease Rating Scale assessments, participants will be instructed to take their next

prescribed dose of medication and rest for 60 minutes before completing an "ON-state" eye tracking assessment. Please indicate whether in your medical judgment there are significant risks with a two-hour period delay in medication intake while performing assessments. If there is significant risk to your patient with disrupting medications they will be excluded from the study. Please complete the approval form in pages 4-7 if, in your medical judgment, this individual has no contraindications or serious health risks in participating in this study. Thank you for your time to read this summary and evaluate your patient's status for participation in this study. If you have any questions please do not hesitate to contact the principal

you for your time to read this	summary and evaluate	your patient's status	for participation in
this study. If you have any que	stions please do not h	esitate to contact the	principal
investigator in the study,			
Thank you.			
To be completed by the pa	rticipant		
Date			
Name			
Address			
City/State/Zip			
Phone()	_E-mail		_
Date of Birth(MM/DD/YYYY)			
Primary Diagnosis			
Date of Diagnosis(MM/DD/YYYY)_			
Emergency Contact Name			
Relationship	Phone #(	)	_

Do you have any allergies? If yes, (please specify)	
Have you ever been diagnosed with one of the following?	
Diabetes(Y/N)(Insulin?) Parkinson's diseaseHypertensionPulmo	onary Disease
20/20 or Corrected Vision (specify)Retinal Disease(specify)	
Glaucoma Ischemic Optic NeuropathyPseudoexfoliation Syndrome	Ocular Surgery
Ocular Trauma Cataracts Orbital myositis	
Other neurological disorders/injuries apart from Parkinson's Disease?diagnosis?	If yes, what was the
Have you undergone brain surgery (ex: for Deep Brain Stimulation)	
Are you receiving Outpatient Therapy YESNO  Please list any Surgeries including dates	
Please list the medications that you are currently taking	

Has there been a change in use of medication in the last 6 months? YES	NO
Do you have a family member or caretaker that can be present and aid	in transport for the "OFF
medication state" assessment days? YESNO	
By signing this document, I give permission to have my neurologist mail	a medical release form to Prof.
Signature	Date
To be completed by the participant's neurologist:	
Participants Name	
Diagnosis (List All)	
Does your patient meet the UK Brain Bank Diagnostic Criteria for PD? (y	
Please check all the PD related impairments:	
unilateral involvement only	
unilateral and axial involvement	
bilateral involvement without impairment of balance	
Other/higher level of involvement	
Height Weight Pulse BP Sex_	

Physical Exam	Normal	Abnormal	Explanation of Abnormalities
Head/Neck			
Eyes/Vision			
Ears/Hearing			
Heart/Lung			
G.I.			
C.N.S.			
Skin	-		
Significant "ABNOR	MAL TEST" EKG/	X-RAY/LAB):	
Medications (please	e list)		
Change in medication	on in the past 6	months YES	_NO If yes, specify

Initiation of medications that influence walking/mobility/coordination in the past 30 days?

YESNO
Do you expect any changes in medication in the next two months?
YESNO
If your patient has PD, is your patient under Deep Brain Stimulation (DBS) treatment?
YESNO
If the participant has PD, is there significant risk associated with withholding from movement related
medications for two hours from the regularly scheduled intake time and resuming intake immediately
after the two hour delay period?
YESNO
Do you have any modifications to the medication intake delay time and continuation of intake after the two hour delay?
YESNO
IF YES, please clarify:
Is it safe for the participant to resume medications after a two hour delay period?
YESNO
IF NO, but there is s safe way to stop and resume medications, please provide Instructions on how stop

and resume medication:

Approval for participation: YESNO		
Comments/Restrictions:		
Address:		
City:State	e:	_ZIP;
Physician's Signature:		
Physician's Emergency Contact Telephone Number:		
Date:		
Permission to release patient information to the		
can be found at the bottom of page 3	3.	
215.65		
PLEASE MAIL TO:		

THANK YOU FOR YOUR PARTICIPATION.