



Date Form Completed:

1. Project Title:		
2. Chart Wills (Farmed and and		
2. Short Title (5 words or less):		
3. Brief Project Description:		
4. Protocol # (If applicable)		1
IRB Protocol Number:	Approval Date:	Expiration Date:
IACUC Protocol Number:	Approval Date:	Expiration Date:
5. Anticipated Start Date:	Proje	cted End Date:
-	•	
6. Name of the Principal Investigator		
Name:		Dept:
Address:		
Phone:		Fax:
Email:		
7. Name of the Primary Contact Person e.	g. responsible for runnir	ng project on-site
Name:	,,	Dept:
Address:		
Phone:		Fax:
Email:		
Cell phone for emergency contact (e.g. scanner is down, s	cnow storm atc.):	





8. Please check one: ☐ Funded Research Project	☐ Unfunded Pilot Study			
For Pilot Projects Only (obtaining data with subjects for future funded projects):				
Number of hours of MR scan time requested for pilot study (20 hours max.):				
If data is to be used for a grant application, please provide potential funding source and application deadline:				
Date email request sent to dalsop@bidmc.harvard.edu :				
Note: Pilot projects must be approved in writing before completing this form. If you exceed approved hours, you must reapply for additional hours. Please contact MRI Research at MRIResearch@bidmc.harvard.edu with any questions.				
For ALL Funded Accounts: Source (e.g. NIH, etc.) of Funding:				
Duration of Funding: Start Date:	End Date:			
Grant Number if applicable (e.g. NIH, etc):				
Billing Information:				
Research Administrator:	Phone:			
Address:	Email:			
Please provide the names and email addresses where the invoice should be	sent to (i.e. PI, CRA, Research Administrator, etc.):			
Imaging Protocol:				
Was this study derived from BI Pilot MR data? Yes No For industry-sponsored studies, did your sponsor provide a specific protocol a If yes, please email this to MRIResearch@bidmc.harvard.edu	nd imaging manual?			
Is Pre-Protocol Development Requiredsetting up the protocol/sequences were Yes No Number of hours requested for protocol development (should not exceed 5) Name of person you are working with from MRI Research re: protocol development (should not exceed 5)	hours):			
If protocol is known, please list sequences:				
9. Which magnet will be used: 1.5T 3T Both 1.5T and 3T	West Campus MRI			





10. Estimated duration and total number of scanning sessions requested. *Reminder, a standard study slot consists of a 45 minute scan and 15 minutes for setup, breakdown, and cleaning. Please also include the time to set up study equipment. Studies will be booked and charged in 30 minute increments. The time may be adjusted accordingly after the pre-protocol phase is complete. Please contact Fotini Papadopoulou, MRI Research Technologist, with questions related to scan duration @ 617-667-2156 or email fpapadop@bidmc.harvard.edu					
Total number of subjects:					
Total number of scans/subject:					
Estimated scan time/subject:					
11. Will healthy volunteers be scan	ned?			Yes	☐ No
Number of healthy volunteers to be	e scanned:				
Frequency of scan(s) per healthy vo	olunteer and interval:				
12. Will patients be scanned?				Yes	No
Number of patients to be scanned:					
Frequency of scan(s) per patient an	id interval:				
Are these clinical scans that need to be booked in CCC and charged to the patient's insurance?					
Will your study use LAR's to obtain consent? If yes, will the LAR/a family member familiar with the patient's medical history be able to accompany them to the MRI to complete and review the MRI Safety Form? Yes No					
13. Will contrast be used?:			Yes	☐ No	
If yes:	Turne does 8 venter				
	Type, dose & route: Administered by:	prior to scan	during scan		
	Type, dose & route: Administered by:	prior to scan	during scan		
14. Will medications other than contrast be administered prior to the scan or during the scan?: Yes No					
	Type, dose & route: Administered by:	prior to scan	during scan		
15. Is a clinical interpretation of the images required? If yes, is a Radiology read on BIDMC letterhead, as opposed to a formal read in the medical record sufficient: Yes No If no, please explain what is needed:					

Revised: 8/2020





16. Will animals be used in this study:	Yes No			
If yes, please describe the details of the study in an attac	ched document or email to MRIResearch@bidmc.harvard.edu			
17. Do you or your sponsor have a preferred form of d If yes, please describe:	lata transmittal (Sponsor upload, SFTP, XNAT, etc.)?			
18. Do you or your sponsor require the having the orig	ginal copy of the safety checklist (as opposed to a copy)?			
19. Will hazardous chemicals, inhalational anesthetics of the study in an attached	-			
20. Will you be bringing any objects or devices that will need to enter the MRI Room during this study?				
21. Will you be using any non-FDA approved devices d If yes, please describe:	luring the MRI Scan? Yes No			
I attest the information provided in this application is current Research scanners, MRI Policy and Procedures and ensure fir	and accurate. I will adhere to the BIDMC policies as stated in the Ansin nance responsibility for the cost of the study.			
Printed Name of Principal Investigator: (Electronic Signature)	Date:			
Checklist for Submission Completed Requisition for MR Scan Time (submit e-version with electronic signature) Research Protocol (Part B of BI IRB application is sufficient), Part A, and Research Staffing Form Current IRB Approved consent form (electronic PDF preferred) IRB approval letter from home institution (electronic PDF Preferred) Part E and H of BIDMC IRB application if using contrast agents or other medications, along with IRB approval documentation Submit all documents to the MRI Research at MRIResearch@bidmc.harvard.edu				
ADMINISTRATIVE USE ONLY				
Date Forms Received:	MRI Schedule Study ID Title:			
	Data Transfer Method:			
Approved Scan Rate:				
Approved Scan Time Slot:	Approval from MR Research:			