

Protocol Number: Protocol Number: PI: Protocol Synopsis: Sponsor/CRO: As part of the pre-study activities for the upcoming protocol, please provide the following information regarding your access to the required population and your site's initial plan for recruiting participants in this trial.			
	Based upon review/search of available databases, document the number of participants fit protocol criteria and would be contacted for participation in trial:		
On v	what sources are you basing this number?		
	Medical Record Chart Review (i2b2, ICD-10 code search)		
	Community Database		
	Research Database		
	Other:		
	Please list the potential challenges you see to enrolling participants and what you ld implement to overcome these issues:		
	Inclusion /Exclusion criteria too strict		
	Distance/ Participant lives too far away/Participant Travel or Transportation Issues/Participant does not want to drive to UAB due to traffic/Parking		
	Protocol requires too much from participant: procedures/frequency of visits/duration of protocol (lasts for years)		
	Study/Protocol will not pay participant for time to participate		
	Age of participant population		
	Investigational Product (IP) administration route/ requirements (multiple doses, injections, refrigeration)		
	Randomization deterrent		
	Seasonal illness/ Time of year for enrollment		
	Other:		



	Social Media (Facebook, Twitter, Snapchat, Instagram, Google Ads)
	Newspaper/ Magazines/written advertising outlets
	Community Outreach (Navigators, Health Advisors)
	Television (local affiliates such as ABC, CBS, NBC, FOX)
	Electronic Signage (local advertisers provide billboard space)
	Radio
	Video Recordings in waiting areas
	Other Print Materials & Mailings (Flyers, Brochures, pamphlets)
	MD/Outside Community referrals (engage professional networks)
	Clinical Faculty Engagement through Faculty meetings or listservs
	<u>ClinicalTrials.gov</u>
	Research Match
	NCATS (Recruitment Innovation Center)
	Other
	source(s) other than those noted above will be used, please provide details and per of participants who could be contacted for participation in this trial:
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	e the costs for the above resources included in your budget? Yes or No

FIER ONE

Based upon the above data, please provide a written description of your site's Recruitment Plan for the study.

- Include details of the number of participants that will be scheduled for screening within 4 weeks of the study opening at your site:
- What is the goal for the number of participants to be enrolled per month until target enrollment has been reached?
- What is your initial plan for recruiting participants (what tactics will you use):

TER TWO

5. What are your contingency plans for recruitment if your recruitment plan is not yielding enrollment goals, document the triggers or timelines for implementation should they be required?

First Contingency Plan



Retention Plans: Have you thought about how you are going to retain your participants for the duration of the trial/protocol? Will you provide Participant Retention Materials? Yes or No						
						Reminder Materials: appointment cards, text reminders, calendars
						Birthday cards
	Holiday cards					
	Bags to carry Investigational Product (IP)					
	Newsletter about protocol					
	Regular personal contact					
	Provide transportation					
	Provide parking					
	Regular expressions of gratitude for participating					
	Provide meal /snacks for participant					
	Inform participant of results of protocol					
	Other					
Did you include retention materials in budget? Yes or No						
Who will provide the retention materials?						