SJD Institutional Review Board Title: Recruitment Plan Form Code: SJDIRB Form 3.6 Version: 02

Section 1. To be filled to	up by the Principal Investigator.			
1. Protocol Title		2. Principal Investigator/s		
3. Sponsor		4. Co-Investigator/s		
5. Study Sites		6. Target number of respondents under SJDIRB		
TARGET POPULATION				
7. Describe the characteristics of the individuals you plan to recruit (e.g., age, gender, ethnicity, health status, etc.).		8. Provide a rationale for the chosen population.		
RECRUITMENT PLANS: Check all that apply and provide details for each method selected:				
OFlyers/Posters (attach copies)	9. Where will they be posted?		10. For how long?	
OSocial Media	Specify platforms and provide sample posts How will confidentiality be			
OPresentations	maintained? 13. Where and to whom?			
OReferrals from healthcare providers/ community organizations	14. Provide details			
OAdvertisements	15. Specify where ads will be placed and provide copies			
○Other	16. Specify			
	COMMUNITY ENGAGI	EMENT ACTIVITIES		
17. Describe planned activities	e.g., health fairs, community meetings, workshops, presentations at community centers, partnerships with local organizations			
18. Purpose of activities	e.g., raise awareness of the study, build trust, address community concerns, ensure culturally appropriate recruitment strategies			
19. How will these activities support ethical recruitment?				





SJD Institutional Review Board



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RECRUITMENT MATERIALS CONTENT (e.g. Posters, Powerpoint presentations, etc.)			
20. Confirm that all recruitment materials include the following:	OStudy title OName of the principal investigator/contact person OPurpose of the research study (in lay language) OParticipant eligibility criteria OTime commitment required OLocation of the research OStatement about private responses for social media recruitment OIndication of compensation (if applicable) ODeadline for responding (if applicable) OContact details of research ethics reviewer (San Juan de Dios Institutional Review Board)		
21. Language and Tone - Confirm that recruitment materials adhere to the following:	 OUse of "participant" instead of "patient," "subject," or "volunteer." OUse of simple/lay language, avoiding jargon. OAvoidance of acronyms, abbreviations, or mnemonics unless well-known and non-sensationalistic. OClear explanation of investigational treatments. OAvoidance of coercive or sensationalistic language. OAvoidance of stereotypes in text and images. ONo promises of favorable outcomes beyond those in the informed consent. ONo claims about the safety or efficacy of investigational products. (If applicable) OAccurate description of medical treatment provided (if any). OCompensation not featured as the lead element. ONo mention of commercial sponsors or product manufacturers. 		
ETHICAL CONSIDERATIONS			
22. Describe any potential ethical concerns related to recruitment and how they will be addressed.			
23. If any required information is omitted from recruitment materials, explain the rationale.			
24. Recruitment Timeline: Provide estimated start and end dates for recruitment.			





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25. Monitoring and	
Evaluation:	
Describe how the	
effectiveness of	
the recruitment	
plan will be	
monitored and	
evaluated.	

Name of Principal Investigator	Signature and Date

This form, along with the submitted recruitment materials, will be reviewed by the IRB to ensure ethical and responsible recruitment practices.



