



WMSU-REO-FR-005.02 Effective Date: 04-June-2025

## **INFORMED CONSENT ASSESSMENT FORM**

Study Protocol Title*	RxVision: OCR-Based Medical Prescription Reade	r using TroCR and E	BioBERT			
WMSU REO Code		Type of Review	□Expedited	□Full		
Researcher*	Regine B. Bagalangit, Roland Jay J. Pada Ushlie Mae U. Ungaya	ERP	□Chair	□Member		
Name of Adviser*	LUCY F. FELIX-SADIWA, MSCS	Institution*				
Name of Reviewer		Date Received				
Guide questions for reviewing the informed consent process and form						
If NO, please explain.	the informed consent of the participants?  ants provided with sufficient information regarding:	□Unable to Asses	3 103			
Purpose of the study?		□No	□Yes			
Expected duration of participation?		□No	□Yes			
Does the protocol include an adequate process for ensuring that consent is voluntary?		□No	□Yes			
Procedures to be carried out?		□No	□Yes			
Discomforts and inconveniences?		□No	□Yes			
Risks (including possible social, physical, emotional, and psychological)?		□No	□Yes			
Random assignment to experimental and control group?		□Not applicable	□Yes □No			
Benefits to the participants?		□No	□Yes			
Compensations/reimbu	rsements of expenses	□No	□Yes			

	re of Primary Reviewer		Review Date
			·
	Reasons for disapproval:		
	☐ Disapproved		
	☐ Major revisions required		
Recommendation:	<ul><li>□ Approved</li><li>□ Minor revisions required</li></ul>		
o to contact for pertinent	questions and/ for assistance in the research	n-related injury	/?
you nave any other cond ne	CITIS!		
ne informed consent writ guage that participants c you have any other cond		□No	□Yes
ent of confidentiality?		□No	□Yes
uties and responsibilities of the participants are duly stated?			□Yes
ticipants may withdraw f	rom the study anytime without any penalty?	□No	□Yes