Issue date: 21 October 2024 Valid Until: 21 October 2029



**NO: SAMM 946** 

(Issue 1, 21 October 2024 replacement of SAMM 946 dated 28 August 2025)

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LABORATORY LOCATION: (PERMANENT LABORATORY)  □ 本	Clinical Research Center, Sarawak General Hospital Jalan Tun Ahmad Zaidi Adruc, 93586 Kuching, Sarawak., 93586, SARAWAK MALAYSIA
ACCREDITED SINCE :	21 OCTOBER 2024
FIELD(S) OF MEDICAL TESTING:	CHEMICAL PATHOLOGY HAEMATOLOGY
SITE:	
1 . SITE LABORATORY(HQ) :	MOLECULAR LABORATORY, LEVEL 1, PATHOLOGY DEPARTMENT SARAWAK HEART CENTER, MALAYSIA
FIELD(S) OF MEDICAL TESTING:	MOLECULAR GENETIC

The standard used for assessment of this laboratory is MS ISO 15189:2022 (ISO 15189:2022, IDT).

A medical laboratory's fulfilment of the requirements of ISO 15189 means the laboratory meets both the technical competence requirements and the management system requirements necessary for it to consistently deliver technically valid test results. The management system requirements in ISO 15189 are written in language relevant to a medical laboratory's operations. Medical laboratories that implement ISO 15189 operate generally in accordance with the principles of ISO 9001. (See Joint IAF-ILAC-ISO Communiquè, November 2021)

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CENTRAL LOCATION	Clinical Research Center, Sarawak General Hospital Jalan Tun Ahmad Zaidi Adruc, 93586 Kuching, Sarawak., 93586, Sarawak
FIELD(S) OF MEDICAL TESTING:	CHEMICAL PATHOLOGY, HAEMATOLOGY

#### SCOPE OF MEDICAL TESTING: CHEMICAL PATHOLOGY

Specimen Tested	Type of Test/ Properties Measured/	Test Methods, Specifications/ Equipment/Techniques Used
Routine Chemistry Plasma	? – glutamyltransferase (GGT)	Multilayered Film method by using Fuji DriChem NX 500 as documented in CRCSGH-QWI-
	Amylase (AMYL)	LAB-3-002
	Alkaline Phosphatase (ALP)	Colorimetric and Non-Selective
	Total Cholesterol (TCHO)	Electrode method by using Roche Cobas c311 as
	High Density Cholesterol (HDL-C)	documented in CRCSGH-QWI- LAB-3-027
	Total Protein (TP)	
	Aspartate Aminotransferase (GOT/AST)	
	Glutamic pyruvic transaminase (GPT/ALT)	
	Magnesium (Mg)	
	Calcium (Ca)	
	Lipase (LIP)	
	Lactate Dehydrogenase (LDH)	
	Triglycerides (TG)	
	Total Bilirubin (TBIL)	
	Creatinine (CRE)	
	Blood Urea Nitrogen (BUN)	

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	Albumin (ALB)	
	Uric Acid (UA)	
	Inorganic Phosphate (IP) Direct Bilirubin (DBIL)	
	Glucose (GLU)	
	Electrolyte	
	1. Sodium (Na+), 2. Potassium (K+)	
	3. Chloride (Cl-)	
Routine Chemistry	Thyroid-Stimulating Hormone	Fluorescent measurement
-immunoassay	(TSH)	through Two Wavelength
Plasma		Detection (Top-Top) method by
	Free Thyroxine (FT4)	using TOSOH AIA-900 as
		documented in CRCSGH-QWI-
		LAB-3-003

#### **SCOPE OF MEDICAL TESTING: HAEMATOLOGY**

Specimen Tested	Type of Test/	Test Methods, Specifications/
	<b>Properties Measured/</b>	Equipment/Techniques Used
Routine Haematology	Complete Blood Count (CBC)	Cell Count by using Abbott Cell
Whole Blood		Dyn Ruby as documented in
	Automated Differential Count /	CRCSGH-QWI-LAB-3-017
	Parameter:	
	1. Neutrophil (NEUT)	
	2. Lymphocyte (LYMPH)	
	3. Monocyte (MONO)	
	4. Eosinophil (EO)	
	5. Basophil (BASO)	
	Platelet Count (PLT)	
	Haematocrit (HCT)	
	Haemoglobin (Hb)	
	Red Cell Counts (RCC)	
	White Cell Count (WCC)	
	Mean Corpuscular Volume (MCV)	

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	Mean Corpuscular Hemoglobin (MCH)	Cell Count by using Abbott Cell Dyn Ruby as documented in CRCSGH-QWI-LAB-3-017
	Mean Corpuscular Hemoglobin Concentration (MCHC)	
	Red Blood Cell Distribution Width (RDW-CV)	
	Mean Platelet Volume (MPV)	
	Reticulocyte (Absolute Count) (%)	New Methylene Blue NCCLS method, Supravital Staining Technique by using Abbott Cell Dyn Ruby as documented in CRCSGH-QWI-LAB-3-017
Coagulation Plasma	Prothrombin Time (PT)	Clotting method using SYSMEX CA-600
Гіазіна	Activated Partial Thromboplastin Time (APTT)	series as Documented in CRCSGH-QWI-LAB-3-005

	1. MOLECULAR LABORATORY LEVEL 1, PATHOLOGY DEPARTMENT SARAWAK HEART CENTER,MALAYSIA
FIELD(S) OF MEDICAL TESTING:	MOLECULAR TESTING

# FIELD(S) OF MEDICAL TESTING: MOLECULAR GENETIC

Specimen Tested	Type Of Test / Properties	Test Methods, Specifications/
	Measured	Equipment/Techniques Used
Whole Blood	Detection of Familial	DNA sequencing is conducted
	Hypercholesterolemia	using NGS
	Genetic Variants for:	(MiSeq) as described in
	1) LDLR	CRCSGH-QWI-7-018
	2) APOB	
	3) PCSK9	
	4) LDLRAP1	

NOTE: