Schedule

Issue date: 01 March 2024 Valid Until: 29 November 2025



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LABORATORY LOCATION/ CENTRAL OFFICE:	Malaysian Institute of Pharmaceuticals and Nutraceuticals (IPharm), National Institutes of Biotechnology Malaysia Block 5-A, Halaman Bukit Gambir 11700 Gelugor Pulau Pinang,
	11700, PULAU PINANG MALAYSIA
ACCREDITED SINCE :	24 MARCH 2025
FIELD(S) OF TESTING:	TOXICITY

This laboratory has demonstrated its technical competence to operate in accordance with MS ISO/IEC 17025:2017 (ISO/IEC 17025:2017).

This laboratory's fulfillment of the requirements of ISO/IEC 17025 means the laboratory meets both the technical competence requirements and management system requirements that are necessary for it to consistently deliver technically valid test results and calibrations. The management system requirements in ISO/IEC 17025 are written in language relevant to laboratory operations and operate generally in accordance with the principles of ISO 9001 (see Joint ISO-ILAC-IAF Communiqué dated April 2017).

CENTRAL LOCATION:	Malaysian Institute of Pharmaceuticals and Nutraceuticals (IPharm), National Institutes of Biotechnology Malaysia Block 5-A, Halaman Bukit Gambir 11700 Gelugor Pulau Pinang , 11700, Pulau Pinang
FIELD(S) OF TESTING:	TOXICITY,

SCOPE OF TESTING: TOXICITY

Material / Product Tested	Type Of Test / Properties Measured / Range Of Measurement	Standard Test Methods / Equipment / Techniques
Medical Devices	Cytotoxicity	ISO 10993-5 (2009): Biological evaluation of medical devices – Part 5: Tests for in vitro cytotoxicity – Annex C: MTT cytotoxicity test
Health Supplements Ingredients For Traditional Medicine industrial Chemicals ingredients For Cosmetics	Cytotoxicity	In-house method: IP/LTM/002 based on ISO 10993-5 (2009): Biological evaluation of medical devices – Part 5: Tests for in vitro cytotoxicity – Annex C: MTT cytotoxicity test

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