

PIHAK BERKUASA PERANTI PERUBATAN KEMENTERIAN KESIHATAN MALAYSIA

TRAINING FEE



THE TRAINING OF IMPORTAND EXPORT OF MEDICAL DEVICE REQUIREMENTS SESSION 1

OVERVIEW

This training program is designed to participants in-depth provide with knowledge and hands-on experience in managing the regulatory processes for the import and export of medical devices under the Medical Device Act 2012 (Act 737) and its regulations. It offers a comprehensive guide to comply with relevant laws, documentation requirements, and the use of digital platforms streamline to application submissions. The training aims to strengthen operational efficiency and regulatory compliance, minimizing delays non-compliance risks in trade and processes.





REGISTER NOW!

INTENDED AUDIENCE

Parties involved in the import and export of medical devices, including:

Medical device establishments





MALAYSIA

Management

'ds **2024**

Excellence

REGISTRATION CLOSE: 12 MAY 2025

Upon acceptance of the registration, an invoice (for payment purposes) together with details of the payment methods will be issued within 2-3 working days)

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(**J**) (**D**) Medical Device Authority www.mda.gov.my



Training Outline

08.30 AM - 09.00 AM	Registration & Briefing
09.00 AM - 10:30 AM	 Import Permit/ Verification Slip Hands-on session
10.30 AM - 10:45 AM	Morning Break
10.45 AM - 12.00 PM	 Medical Device Exemption: Import for Re-export Hands-on session
12.00 PM - 01.00 PM	 Medical Device Exemption: Export Only Hands-on session
01.00 PM - 02.30 PM	Lunch Break
02.30 PM - 03.30 PM	 Certificate Free Sale (CFS) & Manufacturing Certificate (MC) Hands-on session
03.30 PM 03.45 PM	Tea Break
03.45 PM - 05.00 PM	 Export Certificate (EC) Hands-on session
05.00 PM	End of Program
** This training Outline is Subject to Change	



(f) (b) Medical Device Authority www.mda.gov.my