MDA CORE
CENTRE OF EXCELLENCE

Medical Device AUTHORITY MALAYSIA
PIHAK BERKUASA PERANTI PERUBATAN
KEMENTERIAN KESIHATAN MALAYSIA

HRD CORP CLAIMABLE

GDPMD SESSION 1:
REGULATORY REQUIREMENTS, COMPLIANCE ASSESSMENT AND IMPLEMENTATION

TUESDAY
10-11 FEBRUARY 2026

START AT
08.30 AM - 17.00 PM

VENUE
PUTRAJAYA/CYBERJAYA (TBC)

TRAINING FEE
RM 1,000
PER PARTICIPANT

Training Overview

This training provides a comprehensive understanding of the Good Distribution Practice for Medical Devices (GDPMD), a mandatory requirement under the Medical Device Authority (MDA) for Authorized Representatives, Importers, and Distributors. GDPMD ensures that medical devices are consistently handled, stored, transported, and distributed in a way that safeguards their safety, quality, and performance throughout the supply chain. Participants will learn how GDPMD supports patient safety by maintaining device integrity from procurement to post-market surveillance.

The program focuses on the core principles of GDPMD, including establishing a quality management system, maintaining suitable premises and equipment, ensuring accurate record-keeping, storage, and distribution, managing outsourced activities responsibly, conducting internal audits and management reviews, and implementing effective vigilance systems to monitor device safety and performance after placement in the market.

By attending this training, participants will gain practical knowledge to meet regulatory expectations, strengthen their organization's compliance framework, and prepare for GDPMD certification. This training supports organizations in achieving certification, enhancing credibility, and protecting patients by ensuring medical devices remain safe and effective throughout their lifecycle"

Who Sould Attend

- Authorized Representatives of medical device companies
- Importers of medical devices
- Distributors of medical devices
- Participants who intend to pursue the GDPMD certification

SCAN QR CODE TO REGISTER

REGISTER BEFORE :
30 JANUARY 2026

Upon submission of your registration, an invoice for payment, along with the payment method details, will be issued within 2–3 working days.
This program is claimable under the SBL Scheme.
Please refer here for the **SBL Scheme terms and conditions**

MDA CORE
CENTRE OF EXCELLENCE

Medical Device AUTHORITY MALAYSIA

PIHAK BERKUASA PERANTI PERUBATAN
KEMENTERIAN KESIHATAN MALAYSIA

HRDCORP CLAIMABLE

TRAINING OUTLINE

Day 1- 10 February 2026

Time	Agenda
08.30 AM– 08.55 AM	**Registration and Training Briefing**
08.55 AM – 09.00AM	Opening Remark
09.00 AM– 09.30 AM	Part 1: Preliminary: Objective, Scope and Application, and Definition
09.30 AM - 10.00 AM	Part 2: Organization and GDPMD Regulatory Compliance System
10.00 AM - 10.30 AM	Part 3: Establishment Responsibilities
10.30 AM – 11.00 AM	Part 4: Resource Management
11.00 AM - 11.30 AM	**Morning Break**
11.30 AM - 01.00 PM	Part 5: Supply Chain and Device Specific
01.00 PM - 02.30 PM	**Lunch Break**
02.30 PM - 03.30 PM	(Cont.) Part 5: Supply Chain and Device Specific
03.30 PM - 03.45 PM	**Tea Break**
03.45 PM - 05.00 PM	Part 6: Surveillance and Vigilance
05.00 PM	End of Day 1

** This Training Outline is Subject to Change

MDA CORE
CENTRE OF EXCELLENCE

Medical Device AUTHORITY MALAYSIA

PIHAK BERKUASA PERANTI PERUBATAN
KEMENTERIAN KESIHATAN MALAYSIA

HRDCORP CLAIMABLE

TRAINING OUTLINE

Day 2 - 11 February 2026

Time	Agenda
08.30 AM– 09.00 AM	**Registration**
09.00 AM– 10.30 AM	Documentation Preparation for GDPMD: Regulatory Compliance Manual and SOPs
10.30 AM – 10.45 AM	**Morning Break**
10.45 AM – 01.00 PM	(Cont.) Documentation Preparation for GDPMD: Regulatory Compliance Manual and SOPs
01.00 PM – 02.30 PM	**Lunch Break**
02.30 PM – 03.40 PM	Audit Preparation and Evidence Expected
03.40 PM – 04.00 PM	**Tea Break**
04.00 PM – 05.00 PM	(Cont.) Audit Preparation and Evidence Expected
05.00 PM	**End of the training**

** This Training Outline is Subject to Change