

MALAYSIAN STANDARD

MS 2636:2019

Halal medical device – General requirements

ICS: 11.040.01

Descriptors: halal, medical device, requirements, compliance, certification

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Committee representation

The Industry Standards Committee on Halal Standards (ISC I) under whose authority this Malaysian Standard was developed, comprises representatives from the following organisations:

Department of Chemistry Malaysia Department of Islamic Development Malaysia

Department of Standards Malaysia

Department of Veterinary Services

Federal Agricultural Marketing Authority

Federation of Malaysian Manufacturers

Halal Industry Development Corporation

Institute of Islamic Understanding Malaysia

Institute of Quality Malaysia

International Islamic University Malaysia

Malaysian Agricultural Research and Development Institute

Malaysian Association of Standards Users

Ministry of Domestic Trade and Consumer Affairs

Ministry of Health Malaysia (Food Safety and Quality Division)

Ministry of Health Malaysia (Medical Device Authority)
Ministry of Health Malaysia (National Pharmaceutical Regulatory Agency)

Ministry of International Trade and Industry

Muslim Consumers' Association of Malaysia

National Metrology Institute of Malaysia

Royal Customs Department

Universiti Sains Islam Malaysia

The Working Group on Halal Medical Devices which developed this Malaysian Standard consists of representatives from the following organisations:

Association of Malaysian Medical Industries

Department of Islamic Development Malaysia, Halal Hub Division

Department of Islamic Development Malaysia, Planning and Research Division

Department of Standards Malaysia (Secretariat)

Halal Industry Development Corporation

Malaysia Medical Device Association

Malaysian Pharmaceutical Society

Ministry of Defence Malaysia, Malaysian Armed Forces Headquarters (Health Services Division)

Ministry of Health Malaysia (Medical Device Authority)

Ministry of Health Malaysia (National Pharmaceutical Regulatory Agency)

Muslim Consumers Association of Malaysia Universiti Sains Malaysia

Co-opted members:

Academy of Contemporary Islamic Studies, Universiti Teknologi MARA

Ain Medicare Sdn Bhd

Allen Healthcare Products (M) Sdn Bhd

APS Medical Sdn Bhd

B. Braun Medical Industries Sdn Bhd

Duopharma Biotech Bhd.

Fresenius Medical Care Malaysia Sdn Bhd

GranuLab (M) Sdn Bhd

Ideal Healthcare Sdn Bhd

Industrial Biotechnology Research Centre, SIRIM Berhad

Pharmaniaga Berhad

Renal Laboratories Sdn Bhd

UWC Healthcare MFG (M) Sdn Bhd

Foreword

This Malaysian Standard was developed by the Working Group on Halal Medical Devices under the authority of the Industry Standards Committee on Halal Standards.

Compliance with a Malaysian Standard does not of itself confer immunity from legal obligations.

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Introduction

The Malaysian Standards relating to halal have been developed to meet the challenges of the growing demand for halal products and services, to complement the halal ecosystem within Malaysia. This standard was formulated based on the concept of halal built-in and used by existing competent authorities towards halal certification.

Halal built-in is a systematic approach to halal product development which begins with predefined objectives by the management. It embeds or integrates the requirements of halal as part of the overall management and control systems. It includes all aspect of manufacturing/ production, from strategy/ planning to research and development to raw material sourcing until delivery of finished product to its point of purchase. This ensures continuous compliance to the specific halal requirements and the aspects of product safety, performance, efficacy/effectiveness and quality, along with the hygienic aspects in manufacturing and handling of the halal product.

Halal requirements are as stated by *Shariah* and *fatwa* that are incorporated into halal standards which is in the form of the standards developed by the Department of Standards Malaysia. Halal should be built-in into the entire system.

Obtaining halal certification is a business strategy and is seen as a value added element. Since, the main requirements of all medical devices are there its safety and performance as such, all medical devices have to comply with the Medical Device Act 2012 (Act 737) before they can be certified halal.