

# MEDICAL DEVICE GUIDANCE DOCUMENT

## HARMONISED CLASSIFICATION OF MEDICAL DEVICES IN ASEAN



**Medical Device Authority**  
MINISTRY OF HEALTH MALAYSIA

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## **Preface**

This Guidance Document was prepared by the Medical Device Authority (MDA) to help the industry and healthcare professionals in their quest to comply with the Medical Device Act (Act 737) and the regulations under it.

This Guidance Document shall be read in conjunction with the current laws and regulations used in Malaysia, which include but not limited to the following-

- a) Medical Device Act 2012 (Act 737);
- b) Medical Device Regulations 2012.

In this Guidance Document, the following verbal forms are used:

- “shall” indicates a requirement;
- “should” indicates a recommendation;
- “may” indicates a permission; and
- “can” indicates a possibility or a capability.

Irrespective of the requirements of this Guidance Document, MDA has the right to request for information or material, or define conditions not specifically described in this document that is deemed necessary for the purpose of regulatory control.

MDA has put much effort to ensure the accuracy and completeness of this guidance document. In the event of any contradiction between the contents of this document and any written law, the latter should take precedence.

MDA reserves the right to amend any part of the guidance document from time to time.

## **CONTACT INFORMATION**

For further information, please contact:

**MEDICAL DEVICE AUTHORITY**  
Ministry of Health Malaysia  
Level 6, Prima 9, Prima Avenue II  
Block 3547, Persiaran APEC 63000  
Cyberjaya, Selangor MALAYSIA  
Email: [mdb@mda.gov.my](mailto:mdb@mda.gov.my)  
Website: <http://www.mda.gov.my>

# HARMONISED CLASSIFICATION OF MEDICAL DEVICES IN ASEAN

## 1 Introduction

ASEAN Medical Device Directive (AMDD) is the Agreement established for the purpose of harmonising medical device regulations in ASEAN countries. This Agreement is entered into force on 1 January 2015. Malaysia has signed this Agreement on 23 April 2020.

The classification of medical devices is one of the requirements in AMDD which is stipulated in Annex 2 and 3 of Article 4 of the AMDD. Malaysia has implemented the classification of medical devices according to this Article. The requirement has also been specified in First Schedule of Medical Device Regulations 2012. Medical devices shall be classified into the following four classes; Class A, B, C or D in accordance with the risk-based classification.

This document is intended to provide guidance to manufacturers and authorised representatives on the risk classification of medical devices including in-vitro diagnostic (IVD) medical devices which has been harmonised in ASEAN.

This guidance document will be updated on a yearly basis to reflect the latest decisions of the ASEAN Medical Device Committee (AMDC).

## 2 Scope

This guidance document provides the harmonised list of risk classification of medical devices including in-vitro diagnostic (IVD) medical devices based on their intended use/purpose as claimed by the manufacturer.

This document shall be read in conjunction with the following documents:

- a) Medical Device Act 2012 (Act 737); and
- b) Medical Device Regulation 2012.

## 3 Terms and definitions

For the purposes of this document, the terms and definitions in Act 737, the regulations under it and the following terms and definitions apply.

### 3.1 ASEAN

Association of Southeast Asian Nations, or ASEAN, is a political and economic union of 10 members states in Southeast Asia, which are Brunei, Cambodia, Indonesia, Laos, Malaysia, Myanmar, Philippines, Singapore, Thailand and Vietnam.

[Source: asean.org]

### 3.2 intended use/ purpose

The objective intent of the manufacturer regarding the use of a product, process or service as reflected in the specifications, instructions and information provided by the manufacturer.

### **3.3 manufacturer**

As defined in Section 2 of Act 737.

### **3.4 medical device**

As defined in Section 2 of Act 737.

## **4 Harmonised Classification of Medical Devices in ASEAN**

### **4.1 General requirements**

The risk classification of medical devices listed are harmonised in ASEAN.

The list of harmonised risk classification of identified medical devices are as listed in Annex A.

**Annex A**  
(normative)

**LIST OF HARMONISED CLASIFICATION OF MEDICAL DEVICES IN ASEAN**

MEDICAL DEVICE NAME	CLASS	REMARK
<b>CLINICAL CHEMISTRY AND CLINICAL TOXICOLOGY DEVICES</b>		
Artificial Pancreas Device System, Threshold Suspend	C	
<b>HEMATOLOGY AND PATHOLOGY DEVICES</b>		
System, Test, Blood Typing Test	C	Except for ABO system [A (ABO1), B (ABO2), AB (ABO3)], rhesus system [RH1(D), RH2 (C), RH3 (E), RH4 (c), RH5 (e)], Kell system [Kel1 (K)], Kidd system [JK1 (Jka), JK2 (Jkb)] and Duffy system [FY1 (Fya), FY2 (Fyb)] determination which are classified as Class D.
<b>IMMUNOLOGY AND MICROBIOLOGY DEVICES</b>		
Assay, Genotype, HIV Drug Resistance, In Vitro	C	
Enzyme Linked Immunosorbent Assay, T. Cruzi	C	
Test, Equipment, Automated Bloodborne Pathogen	A	
<b>CARDIOVASCULAR DEVICES</b>		
Barrier, Adhesion, Cardiovascular	D	
Cardiovascular Catheter Sheath Introducer Kit	B	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Catheter, Peripheral, Atherectomy	B	
Clip, Vascular	B	Unless they are long-term surgically invasive medical devices, are in Class C; Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Compressor, Cardiac, External	A	Unless they are active medical device, in which case they are in Class B; Unless they are specifically intended for therapy the vital physiological parameters, where the nature of variations is such that it could result in immediate danger to the patient, for instance variations in cardiac performance, respiration, activity of central nervous system, in which case they are in Class C.
DC-Defibrillator, High Energy, (Including Paddles)	C	Unless they are surgically invasive, in which case they are in Class D.
Device, Vascular, For Promoting Embolization	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Device. Laser Peripheral Angioplasty	B	Unless this is done in a manner that is potentially hazardous with higher power, in which case they are in Class C; Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Endovascular Suturing System	C	Unless they are intended to be used in

**Annex A**  
(normative)

		direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Fibrillator, Ac	C	Unless they are surgically invasive, in which case they are in Class D.
Glue,Surgical,Arteries	C	Unless they are absorbable, in which case they are in Class D.
Graft, Vascular, Synthetic/Biologic Composite	C	Unless they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin are Class D.
Graft,Vascular,Stainless Steel Tunneler	B	
Kit, Balloon Repair, Catheter	B	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Material, Embolization, Neurovascular, Polymerizing or Precipitating	D	
Polymerizing, Neurovascular Embolization Material	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Programmer, Pacemaker	C	
Prosthesis, Vascular Graft, Of 6mm And Greater Diameter	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, and/or they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
Prosthesis, Vascular Graft, Of Less Than 6mm Diameter	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, and/or they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
Pulse-Generator, Pacemaker, External	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Pulse-Generator, Single Chamber, Single	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Shunt, Portosystemic, Endoprosthesis	C	
Stent, Iliac	D	
Stent, Renal	C	
Stent, Superficial Femoral Artery	C	
Suture, Cardiovascular	D	

**Annex A**  
(normative)

System, Catheter or Guidewire, Steerable (Magnetic)	B	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
System, Esophageal Pacing	B	
Tissue Graft Of 6mm And Greater	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, and/or they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
Tissue Graft of Less Than 6mm	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, and/or they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
Transducer, Pressure, Catheter Tip	B	
Transducer, Vessel Occlusion	B	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
<b>DENTAL DEVICES</b>		
Adhesive, Denture, Acacia and Karaya With Sodium Borate > 12% By Weight	A	
Adhesive, Denture, Karaya With Sodium Borate	A	
Cord, Retraction	A	
Resin, Root Canal Filling Containing Chloroform	B	
<b>EAR, NOSE, AND THROAT DEVICES</b>		
Device, Antichoke, Suction	A	Unless they are active medical device, in which case they are in Class B.
Paste, Injectable for Vocal Cord Augmentation	C	Unless they are absorbable, and/or they are manufactured from or incorporating with animal cells, tissues and/or derivatives thereof, rendered nonviable, or cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
<b>CLINICAL CHEMISTRY AND CLINICAL TOXICOLOGY DEVICES</b>		
Artificial Pancreas Device System, Bihormonal Control	C	
Artificial Pancreas Device System, Single Hormonal Control	C	
<b>IMMUNOLOGY AND MICROBIOLOGY DEVICES</b>		

**Annex A**  
(normative)

Assay, Enzyme Linked Immunosorbent, Hepatitis C Virus	C	Unless they are intended to be used to detect the presence of, or exposure to, a transmissible agent that causes a life-threatening, often incurable, disease with a high risk of propagation, in which case they are in Class D.
Assay, Hybridization and/or Nucleic Acid Amplification for Detection of Hepatitis C RNA, Hepatitis C Virus	C	Unless they are intended to be used to detect the presence of, or exposure to, a transmissible agent that causes a life-threatening, often incurable, disease with a high risk of propagation, in which case they are in Class D.
<b>CARDIOVASCULAR DEVICES</b>		
Aid, Cardiopulmonary Resuscitation	B	Unless their characteristics are such that they may administer or exchange energy to or from the human body in a potentially hazardous way, including ionising radiation, taking account of the nature, the density and site of application of the energy, in which case they are in Class C.
Automated External Defibrillators (Non-Wearable)	C	
Over-The-Counter Automated External Defibrillator	C	
Reprocessed Intravascular Ultrasound Catheter	D	
Wearable Automated External Defibrillator	C	
<b>GASTROENTEROLOGY-UROLOGY DEVICES</b>		
Agent, Bulking, Injectable for Gastro-Urology Use	C	Unless they are intended to have a biological effect or to be wholly or mainly absorbed, or manufactured from or incorporating animal cells, tissues and/or derivatives thereof, rendered nonviable, cells, tissues and/or derivatives of microbial or recombinant origin, in which case they are in Class D.
Analyzer, Diagnostic, Fiber Optic (Colon)	B	
Apparatus, Hemoperfusion, Sorbent	C	
Cannula, A-V Shunt	B/C	For short-term use are in Class B, and long-term surgically invasive medical devices, are in Class C.
Catheter, Hemodialysis, Implanted	C	Unless they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system, in which case they are in Class D.
Cooler, Esophageal and Gastric	B	
Cooler, Prostatic	B	
Device, Ultrasonic, Thermal Ablation	C	
Graft, Vascular, Hemodialysis Access, Synthetic/Biological Composite	C/D	For synthetic non-absorbable are in Class C, and animal origin absorbable, are in Class D.
Implant, Anti-Gastroesophageal Reflux	C	
Implant, Intra-gastric For Morbid Obesity	C	
Implanted Fecal Incontinence Device	C	
Light Source System, Diagnostic Endoscopic	A	
Lithotripter, Shockwave (For Treating Gallbladder	C	

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(normative)

Stones)		
Prosthesis, Vas Deferans	C	
Replacer, Ureteral	C	
Stent, Urethral, Bulbous, Long term	C	
Stent, Urethral, Prostatic, Long term	C	
Stent, Urethral, Prostatic, Short term	B	
Stent,Urethral,External Sphincter, Long Term	C	
Transurethral Occlusion Insert, Urinary Incontinence- Control, Female, short term use	B	
Valve, Ureterovesicle	C	
<b>GENERAL AND PLASTIC SURGERY DEVICES</b>		
Bronchial Thermoplasty System	C	
Pump, Vacuum, Electric, Suction-Type Electrode	B	
Sealant,Polymerizing, Absorbable	D	
Surgical Device, For Ablation of Cardiac Tissue	D	
Suture, Absorbable, Natural	D	
System, Laser, Photodynamic Therapy, Activation of injected therapeutic compounds or photosensitizing chemical substances for treatment	B	
<b>GENERAL HOSPITAL AND PERSONAL USE DEVICES</b>		
Controller, Closed-Loop Blood Glucose	C	
Dressing, Wound and Burn, Interactive	B	Unless they are intended to be used principally with wounds which have breached the dermis and can only heal by secondary intent, in which case they are in Class C.
Monitor, Skin Resistance/ Skin Temperature, for Insulin Reactions	C	
Pump, Drug Administration, Closed Loop	C	
<b>NEUROLOGICAL DEVICES</b>		
Adhesive, Tissue for Aneurysmorrhaphy	D	
Stimulator, Functional Walking Neuromuscular, Non- Invasive	B	
<b>OBSTETRICAL AND GYNECOLOGICAL DEVICES</b>		
Band, Tubal Occlusion	D	
Barrier, Absorbable, Adhesion	D	
Cap, Cervical, Contraceptive	C	Unless they are incorporating, as an integral part, a substance which, if used separately, can be considered to be a medicinal product (e.g. spermicide), and which is liable to act on the human body with action ancillary to that of the medical devices, are in Class D.
Condom, Female, Animal Tissue	D	
Condom, Female, Single-Use	C	
Device, Occlusion, Tubal, Contraceptive	D	
Analyzer, Semen Analysis	B	
Device, Thermal Ablation, Endometrial	C	
Dilator, Cervical, Expandable	A	Unless they are intended for short-term use, are in Class B.

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(normative)

Dilator, Cervical, Vibratory	A	Unless they are intended for short-term use, are in Class B.
Endoscope, Fetal Blood Sampling (And Accessories)	B	
Prosthesis, Suture, Cerclage, synthetic, non-absorbable	C	
Sensor, Electro-Optical (For Cervical Cancer)	B	
Stimulator, Fetal, Acoustic	B	
Stimulator, Vaginal, Muscle, Powered, For Therapeutic Use	B	
System, Abortion, Metreurynter-Balloon	B	
System, Telethermographic, Infrared	B	
System, Thermographic, Liquid Crystal, Powered (Adjunctive Use)	B	
<b>PHYSICAL MEDICINE DEVICES</b>		
Diathermy, Shortwave, For Use Other Than Applying Therapeutic Deep Heat	B	Unless their characteristics are such that they may administer or exchange energy to or from the human body in a potentially hazardous way, including ionising radiation, taking account of the nature, the density and site of application of the energy, in which case they are in Class C.
Diathermy, Ultrasonic, For Use Other Than Applying Therapeutic Deep Heat	B	Unless their characteristics are such that they may administer or exchange energy to or from the human body in a potentially hazardous way, including ionising radiation, taking account of the nature, the density and site of application of the energy, in which case they are in Class C.
Extracorporeal Shock Wave Treatment, for physical therapy purpose	B	
Orthosis, Pneumatic Structure, Rigid	B	
Peripheral Electromagnetic Field (Pemf) To Aid Wound Healing	B	
<b>IMMUNOLOGY AND MICROBIOLOGY DEVICES</b>		
Kit, Immunochromatographic, Bacillus Anthracis Differential Antibody	C	
Reagents, 2009 H1n1 Influenza Virus (Swine Origin), Nucleic Acid or Antigen, Detection and Identification	C	
<b>CARDIOVASCULAR DEVICES</b>		
Batteries, Rechargeable	A	
Balloon Aortic Valvuloplasty	D	
Cannula, Arterial, Cardiopulmonary Bypass (CPB), Embolism Protection	D	
Device, Biopsy, Endomyocardial	D	
Probe, Blood-Flow, Extravascular	B	
Pulmonary (Pulmonic) Valvuloplasty Catheters/ Percutaneous Valvuloplasty Catheter	D	
<b>DENTAL DEVICES</b>		
Barrier, Animal Source, Intraoral	D	
Bone Grafting Material, Animal Source	D	
Oral Wound Dressing W/Drug and/or Biologic	D	

**Annex A**  
(normative)

Point, Silver, Endodontic	B	
<b>EAR, NOSE, AND THROAT DEVICES</b>		
Collagen, Injectable for Vocal Cord Augmentation	D	
Polymer, Ent Natural-Collagen Material	D	
Replacement, Ossicular (Stapes) Using Absorbable Gelatin Material	D	
<b>GASTROENTEROLOGY-UROLOGY DEVICES</b>		
Catheter, Urinary (Excludes HIV Testing)	B	
<b>GENERAL AND PLASTIC SURGERY DEVICES</b>		
Clamp	A/B	
Composite Cultured Skin (Human Origin)	Non-MD	
Forceps	A/B	
Haemostat Clamp	A/B	
Mesh, Surgical, Absorbable, Abdominal Hernia	D	
<b>NEUROLOGICAL DEVICES</b>		
Catheter, Ventricular, General & Plastic Surgery	D	
<b>RADIOLOGY DEVICES</b>		
Ablation System, High Intensity Focused Ultrasound (Hifu), Mr-Guided	C	
Analyzer, Medical Image, for diagnostic in critical condition	C	
Analyzer, Medical Image, to analyse previously acquired radiology images to identify abnormality during interpretation by the clinician	B	
Applicator, Hyperthermia, Deep Heating, Ultrasound, non-invasive	B/C	Class C if it is High-intensity focused ultrasonic (HIFU) or for high risk application.
Applicator, Hyperthermia, Interstitial, for high risk application	C/D	Class D if they are intended to be used in direct contact with the heart, the central circulatory system or the central nervous system.
Applicator, Hyperthermia, Superficial, RF/ Microwave	B/C	Class C if used for high risk application.
Cap, Cooling (Infants), hypothermia therapy in infants to prevent brain damage cause by oxygen deprivation, active device	B	
Digital Breast Tomosynthesis	C	
Hyperthermia System, Bladder Cancer Treatment, with Chemotherapy	C	
Imager, Breast, Electrical Impedance	B	
Optical Diagnostic Device for Melanoma Detection	B	
System, Hyperthermia, Rf/Microwave (Benign Prostatic Hyperplasia), Thermotherapy	C	
Tester, Acoustic, Bone Quality	B	
Transilluminator (Diaphanoscope)	A/B	
Ultrasound, Hyperthermia, Cancer Treatment, non-invasive	C	
<b>CLINICAL CHEMISTRY AND CLINICAL TOXICOLOGY DEVICES</b>		
Automated Urinalysis System	B	
Calculator/Data Processing Module, For Clinical Use, for specific analyte	B/C/D	Follows the risk class of analytes.

**Annex A**  
(normative)

Calibrators	B/C/D	Calibrators follows the risk class of reagents or analytes.
Complete Gene Expression Profiling Accessory Reagents	C	
Counter (Beta, Gamma) For Clinical Use	A	
Hyaline Membrane Disease Assay	B/C	
Controls without assigned value	B	
Controls with quantitative or qualitative assigned value	B/C/D	Controls follows the risk class of reagents or analytes.
Real Time Nucleic Acid Amplification System, for specific analyte	B/C/D	Follows the risk class of analytes.
Turbidimetric Method, Globulin test	B	
<b>HEMATOLOGY AND PATHOLOGY DEVICES</b>		
Apparatus, Cell-Freezing and Reagents, for IVD use	A	
Kit, Quality Control for Blood Banking Reagents	C/D	
Manual Blood Grouping and Antibody Test Systems	C/D	
Reagents, Specific, Analyte	B/C/D	
Self-Contained Blood Grouping	C/D	
Substance, Blood Grouping of Non-Human Origin For In Vitro Diagnostic Use	C/D	
System, Test, Automated Blood Grouping and Antibody	C/D	
<b>GENERAL AND PLASTIC SURGERY DEVICES</b>		
Mesh, Surgical, Absorbable, Fistula	D	
Mesh, Surgical, Absorbable, Large Abdominal Wall Defects	D	
Mesh, Surgical, Absorbable, Organ Support	D	
Mesh, Surgical, Absorbable, Orthopaedics, Reinforcement of Tendon	D	
Mesh, Surgical, Absorbable, Plastic and Reconstructive Surgery	D	
Mesh, Surgical, Absorbable, Staple Line Reinforcement	D	
Mesh, Surgical, Absorbable, Thoracic, Chest Wall Reconstruction	D	
Mesh, Surgical, Acetabular, Hip, Prosthesis	C/D	If non-absorbable it will be Class C and Class D if the product is absorbable.
Mesh, Surgical, Collagen, Diaphragmatic Hernia	D	
Mesh, Surgical, Collagen, Fistula	D	
Mesh, Surgical, Collagen, Large Abdominal Wall Defects	D	
Mesh, Surgical, Collagen, Orthopaedics, Reinforcement of Tendon	D	
Mesh, Surgical, Collagen, Plastic and Reconstructive Surgery	D	
Mesh, Surgical, Collagen, Plastics, Facial Implants	D	
Mesh, Surgical, Collagen, Staple Line Reinforcement	D	
Mesh, Surgical, Collagen, Thoracic, Chest Wall Reconstruction	D	

**Annex A**  
(normative)

Deployment Balloon for Surgical Mesh	B	
Mesh, Surgical, Non-absorbable, For Stress Urinary Incontinence, Male,	C	
Mesh, Surgical, Non-Synthetic, Urogynecologic, For Apical Vaginal and Uterine Prolapse, Transabdominally Placed	C/D	Class D if the product is from animal origin.
Mesh, Surgical, Non-Synthetic, Urogynecologic, For Pelvic Organ Prolapse, Transvaginally Placed	C/D	Class D if the product is from animal origin.
Mesh, Surgical, Non-Synthetic, Urogynecologic, For Stress Urinary Incontinence, Retropubic Or Transobturator	C/D	Class D if the product is from animal origin.
Mesh, Surgical, Non-absorbable, Polymeric	C	
Septal Stapler/Absorbable Staples	D	
Staple, Non-absorbable, Implantable	C	
Staple, Removable (Skin), for short term	B	
Stapler, Surgical	B	
Suture, Absorbable	D	
Suture, Absorbable, Synthetic	D	
Suture, Absorbable, Synthetic, Polyglycolic Acid	D	
Suture, Knot, Non-absorbable, Mechanical	C	
Suture, Surgical, Absorbable, Polydioxanone	D	
Tape, Surgical, Internal	C/D	
<b>GENERAL HOSPITAL AND PERSONAL USE DEVICES</b>		
Burn Kit include Dressing with Drug	D	
Dressing, Wound and Burn, Hydrogel with Drug and/or Biologic	D	
Dressing, Wound, Collagen	D	
Dressing, Wound, Drug	D	
Gauze, External (With Drug/Biologic/Animal Source Material)	D	
Basic Dressing for external use	A	
<b>NEUROLOGICAL DEVICES</b>		
Balloon, Detachable, For Neurovascular Occlusion	D	
Cannula, Ventricular	D	
Clamp, Carotid Artery, Single Use	B	
Clamp, Carotid Artery, Reusable	A	
Cuff, Nerve	C/D	
Deep Brain Stimulator for Obsessive Compulsive Disorder (OCD)	D	
Instrument, Shunt System Implantation, reusable instrument	A	
Instrument, Shunt System Implantation, single use instrument	B	
Instrument, Shunt System Implantation, if direct contact with Central Nervous System (CNS)	D	
Leukotome	D	
Methyl Methacrylate for Aneurysmorrhaphy	D	
Needle, Neurosurgical Suture	D	
Rongeur, Manual	A/B/D	
Rongeur, Powered	B/D	
Stimulator, Peripheral Nerve, Implanted (Pain	D	

**Annex A**  
(normative)

Relief)		
<b>IMMUNOLOGY AND MICROBIOLOGY DEVICES</b>		
Automated Radioimmunoassay Systems, For Clinical Use include testing agent	B/C/D	

# **MEDICAL DEVICE AUTHORITY**

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## **MINISTRY OF HEALTH, MALAYSIA**

### **Contact Information:**

**Medical Device Authority**  
Ministry of Health Malaysia  
Level 6, Prima 9, Prima Avenue II  
Block 3547, Persiaran APEC  
63000 Cyberjaya, Selangor  
MALAYSIA  
T: (03) 8230 0300  
**Website:** <http://www.mda.gov.my>

