Devices for Modification of Appearance or Anatomy (aesthetic-related purposes)

Product categories:

- Any implant for the modification of any body part
- Any injectable derma filler or mucous membrane filler
- Any instrument, apparatus, implement, machine or appliance intended to be used for the removal or degradation of fat by invasive means
- Any instrument, apparatus, appliance, software, material or any other article, whether used single or in combination, including the software necessary for its proper application intended by the manufacturer used in or on human beings for the purpose of Laser procedures, Micro needling, Ultrasonic Cavitation, High intensity ultrasound tightening and Fraxel radio frequency technology for skin tightening

Product Category	Applicable Rule	Classification
Implants for modification	Rule 8	Class III
Injectable dermal/mucosal membrane fillers	Rule 18	Class III
Fat removal (invasive)	Rule 6 / Rule 7 (surgically invasive devices) and Rule 9 if energy-based	Class IIa / IIb
Laser / RF / HIFU / Cavitation	Rule 9	Class IIb
Microneedling devices	Rule 6	Class IIa

A one year grace period is hereby granted, until 23rd July 2027 for the registration of the above mentioned devices.

22/7/25

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Chief Executive Officer/NMRA

Dr. Prabhath Werawatte
Chief Executive Officer
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Colombo - 02