

REGISTRATION PROCESS FLOW (CLASS B, C and D MEDICAL DEVICE)

01	Determine whether the product is a medical device	ပြံ
02	Accurately determine the risk class and classification rules	al
03	Appropriately group the medical devices	
04	Conduct conformity assessment by a registered Conformity Assessment Body (CAB)	¥= *=
05	Submit application via MeDC@St 2.0+ system	
06	Pay application fee	
07	Assessment by MDA	¥=
08	Regulatory decision	<u>×</u>
09	Pay registration fee	
10	Issuance of electronic registration certificate	Ł
11	Registered device is maintained in the MDA Register (MDAR) for 5 years	櫽