

REGISTRATION PROCESS FLOW (CLASS A MEDICAL DEVICE)

01	Determine whether the product is a medical device	(Jo
02	Accurately determine the risk class and classification rules	.d
03	Appropriately group the medical devices	
04	Submit application via MeDC@St 2.0+ system	
05	Pay application fee	
06	Assessment by MDA	=>>
07	Regulatory decision	<u> </u>
08	Issuance of electronic registration certificate	1
09	Registered device is maintained in the MDA Register (MDAR) for 5 years	产