The medical device must first be registered on the electronic system of the Control Department of Medical Devices and Supplies at the Directorate General of Pharmacy and Drug Control.Moreover, the medical device will be tested so that it meets safety and security requirements according to the World Health Organization to ensure the safety of patients and device operators.It is provided by the manufacturer to support and assist the device users in its safe and appropriate use.Warning: Statement ,that alerts users about a situation that, if not avoided

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