

By Roy L. Baerter

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By ROYAL NETHERLANDS

EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I in III

MDR 77022 R000

Certificate History

(Reference to applicable Commission Specific Regulations, Harmonized Standards, etc. apply with the full text of the applicable parts thereof, if applicable, in the following certificate history symbol request form Certificate Verification (BSI reference))

Date	Reference Number	Action
2020-03-07-01	3674 5	Issue
Current	30054665	Supplemente – Addition of device category, MDN1 0- Non-ctive non-implantable gastrointestinal endoscopy instruments. Amendement – Remove "Also traded as US Endoscopy" from manufacturer name.

First Issue Date: **2020-07-21**

Current Issue Date: **2020-12-11**

Starting Validity Date: **2020-12-11**

Expiry Date: **2022-07-20**

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation's demonstration through the required surveillance activities of the Notified Body.

This certificate will be issued electronically and is bound by the conditions of the contract.

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