

NecroMax 6.0™

The Future of Necrosectomy

EndoRotor is the first-and-only FDA-cleared device to address challenges of endoscopic necrosectomy.

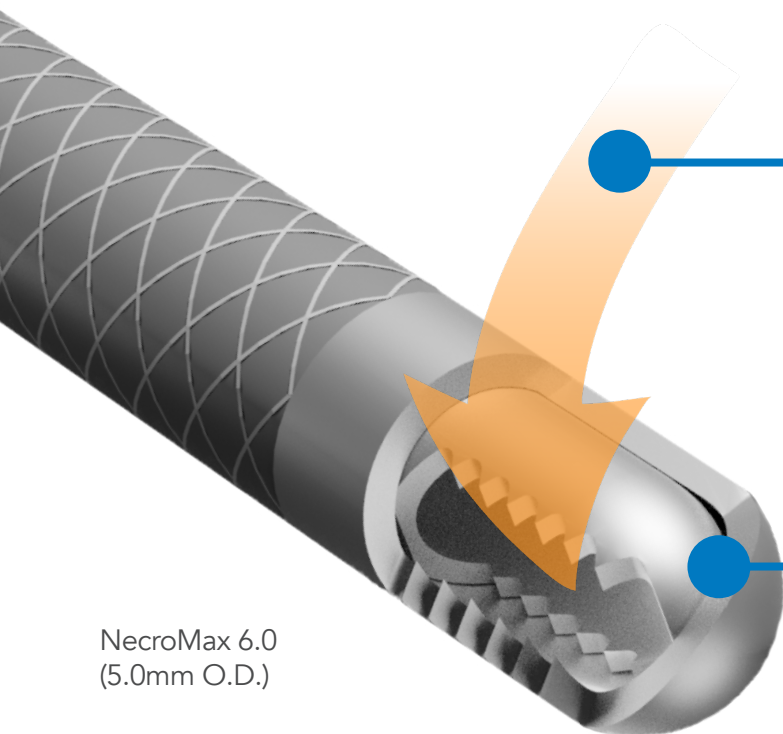
With the endoscopist and patient in mind, we built a larger catheter to maximize necrosis throughput.



NecroMax 6.0



3.2 PED



NecroMax 6.0
(5.0mm O.D.)

8x
Greater
Throughput*

3x
Larger Cutting
Window*

*Compared to EndoRotor 3.2 PED
in benchtop testing

Intended Use: The EndoRotor device is intended for resection and removal of necrotic tissue in symptomatic WOPN/WON after patients have undergone endoscopic ultrasound (EUS) guided drainage.

Warning: The EndoRotor device should not be used in patients with known or suspected pancreatic cancer as per the assessment of the treating physician. Refer to "Instructions for Use" for additional information.

For more information visit www.interscopemed.com
or contact Customer Service:

USA

Tel: 800-461-4289 (toll-free in U.S.)
Email: info@interscopemed.com



Interscope, Inc.
200 Commerce Drive
Northbridge, MA 01534
USA
TEL: +1-800-461-4289
0031 (0) 73-3032541
www.interscopemed.com



EMERGO EUROPE
Prinsessegracht 20
2514 AP The Hague
The Netherlands
EMERGObyUL.com

