

K181418 ScopeFlow Pure(TM)Jul 11, 2019
406 days to decisionK181418 · Product code: **OCX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k181418/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Irrigation/suction System (OCX)
Date received	May 31, 2018
Decision date	Jul 11, 2019
Days to decision	406 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Partnership Medical Limited
Location	Fenton, GB
Contact	Rob Hartley
510(k) history	2 submissions · 2 cleared · 2019-2019

REGULATORY CONSULTANT

Consulting firm	Qrc Consulting, LLC
Contact	Silbiano Gonzales

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k181418/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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