

K990765 FLEXFINDER GUIDE WIRE(REGULAR SHAFT/STIFF SHAFT) MARKED INK BANDS, FLEXFINDER GUIDE WIRE (REGULAR SHAFT/STIFF SHAFT) NOMar 15, 1999
7 days to decisionK990765 · Product code: **OCY** · Gastroenterology & Urology
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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Endoscopic Guidewire, Gastroenterology-urology (OCY)
Date received	Mar 8, 1999
Decision date	Mar 15, 1999
Days to decision	7 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Flexmedics
Location	Minneapolis, MN, US
Contact	WILLIAM C CORRIGAN, JR.
510(k) history	20 submissions · 20 cleared · 1986-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990765/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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