

K081004 PROTECTISCOPE CS COLONOSCOPE, NON-INTRAPULLMay 2, 2008
24 days to decisionK081004 · Product code: **FDF** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k081004/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	Apr 8, 2008
Decision date	May 2, 2008
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker GI
Location	Flower Mound, TX, US
Contact	TIFFANI ROGERS
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081004/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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