

K822085 PRO-SCOPEMay 9, 1983
300 days to decisionK822085 · Product code: **FER** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822085/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anoscope And Accessories (FER)
Date received	Jul 13, 1982
Decision date	May 9, 1983
Days to decision	300 days
Third-party review	No

APPLICANT

Company	Exitec, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k822085/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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