

K811083 ESOPHAGEAL MOTILITY PROBEAug 13, 1981
114 days to decisionK811083 · Product code: **FFT** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k811083/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Ph, Stomach (FFT)
Date received	Apr 21, 1981
Decision date	Aug 13, 1981
Days to decision	114 days
Third-party review	No

APPLICANT

Company	Konigsberg Instruments, Inc.
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1981-1998

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Device record: <https://www.510kdatabase.net/k811083/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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