FDA 510(k) Premarket Notification

K811083 ESOPHAGEAL MOTILITY PROBE

CLEARED

Aug 13, 1981
114 days to decision

K811083 · 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