FDA 510(k) Premarket Notification

K882551 AMBULATORY PH DATA RECORDER SYSTEM, APH-2000

CLEARED

Oct 12, 1988
113 days to decision

K882551 · Product code: **KLA** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k882551/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Esophageal Motility, Anorectal Motility, And Tube (KLA)
Date received	Jun 21, 1988
Decision date	Oct 12, 1988
Days to decision	113 days
Third-party review	No

APPLICANT

Company	**Intl. Biomedics, Inc.**
Location	Cleburne, TX, US
Contact	BENNO L DUNN
510(k) history	2 submissions · 2 cleared · 1986-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k882551/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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