FDA 510(k) Premarket Notification

K831317 BIOPSY ATTACHMENT

CLEARED

May 18, 1983
26 days to decision

K831317 · Product code: **FCI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k831317/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Punch, Biopsy (FCI)
Date received	Apr 22, 1983
Decision date	May 18, 1983
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Diasonics, Inc.**
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1978-1997