

K831317 BIOPSY ATTACHMENTMay 18, 1983
26 days to decisionK831317 · Product code: **FCI** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k831317/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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