

K840144 YAG LASER 8000 GASTROINTESTINAL BLEEMay 18, 1984
126 days to decisionK840144 · Product code: **FCG** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k840144/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	Jan 13, 1984
Decision date	May 18, 1984
Days to decision	126 days
Third-party review	No

APPLICANT

Company	Cooper Medical Corp.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k840144/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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