

K831801 HEMOCLAV VENOUS CATHETERJul 7, 1983
34 days to decisionK831801 · Product code: **FIQ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k831801/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, A-v Shunt (FIQ)
Date received	Jun 3, 1983
Decision date	Jul 7, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Exo, Inc.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-2025

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Device record: <https://www.510kdatabase.net/k831801/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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