FDA 510(k) Premarket Notification

K831801 HEMOCLAV VENOUS CATHETER

CLEARED

Jul 7, 1983
34 days to decision

K831801 · Product code: **FIQ** · Gastroenterology & Urology
Source: 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