FDA 510(k) Premarket Notification

K831401 SINGLE NEEDLE SUBCLAVIAN VEIN ACCESS

CLEARED

Jun 3, 1983
32 days to decision

K831401 · Product code: **FIQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k831401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, A-v Shunt (FIQ)
Date received	May 2, 1983
Decision date	Jun 3, 1983
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Med-West, Inc.**
Location	Mchenry, IL, US
510(k) history	11 submissions · 8 cleared · 1983-1988