FDA 510(k) Premarket Notification

K812621 DELTA-A.V. FISTULA CANNULATION SET

CLEARED

Sep 24, 1981
9 days to decision

K812621 · Product code: **FIE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k812621/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Fistula (FIE)
Date received	Sep 15, 1981
Decision date	Sep 24, 1981
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Amisco Trading Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812621/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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