FDA 510(k) Premarket Notification

K884243 BLOOD TUBING SET

CLEARED

Dec 8, 1988
58 days to decision

K884243 · Product code: **FJK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k884243/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Tubing, Blood, With And Without Anti-regurgitation Valve (FJK)
Date received	Oct 11, 1988
Decision date	Dec 8, 1988
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Mediflex Intl.**
Location	Mchenry, IL, US
Contact	UDINE
510(k) history	9 submissions · 8 cleared · 1984-1992