FDA 510(k) Premarket Notification

K904433 DUAL LUMEN NEEDLES

CLEARED

Nov 23, 1990
56 days to decision

K904433 · Product code: **LBW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k904433/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialysis, Single Needle (co-axial Flow) (LBW)
Date received	Sep 28, 1990
Decision date	Nov 23, 1990
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Akcess Medical Products, Inc.**
Location	New Brunswick, NJ, US
Contact	BALBIR KAPANY
510(k) history	36 submissions · 28 cleared · 1989-1994