FDA 510(k) Premarket Notification

K864629 MODIFIED DUAL LUMEN NEEDLE

CLEARED

Dec 16, 1986
21 days to decision

K864629 · Product code: **LBW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k864629/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialysis, Single Needle (co-axial Flow) (LBW)
Date received	Nov 25, 1986
Decision date	Dec 16, 1986
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
Contact	MARY-LEE DONOGHUE
510(k) history	164 submissions · 160 cleared · 1976-2003