FDA 510(k) Premarket Notification

K896252 MODIFIED QUINTON SINGLE LUMEN CATHETER

CLEARED

Jan 29, 1990
91 days to decision

K896252 · Product code: **MPB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k896252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Non-implanted (MPB)
Date received	Oct 30, 1989
Decision date	Jan 29, 1990
Days to decision	91 days
Third-party review	No

APPLICANT

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