FDA 510(k) Premarket Notification

K833308 DUAL LUMEN NEEDLE

CLEARED

Dec 27, 1983
92 days to decision

K833308 · Product code: **KOC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k833308/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Sep 26, 1983
Decision date	Dec 27, 1983
Days to decision	92 days
Third-party review	No

APPLICANT

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