

K833308 DUAL LUMEN NEEDLEDec 27, 1983
92 days to decisionK833308 · Product code: **KOC** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k833308/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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