

K832580 RAAF CATHETER REPAIR KITJan 3, 1984
154 days to decisionK832580 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k832580/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Aug 2, 1983
Decision date	Jan 3, 1984
Days to decision	154 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/k832580/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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