

K830541 INJECTION SEALING CAP 13791Aug 8, 1983
167 days to decisionK830541 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k830541/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Feb 22, 1983
Decision date	Aug 8, 1983
Days to decision	167 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/k830541/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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