FDA 510(k) Premarket Notification

K830541 INJECTION SEALING CAP 13791

CLEARED

Aug 8, 1983
167 days to decision

K830541 · 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