FDA 510(k) Premarket Notification

K832774 REPAIR KITS FOR RIGHT ATRIAL CATH

CLEARED

Jan 3, 1984
140 days to decision

K832774 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k832774/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Aug 16, 1983
Decision date	Jan 3, 1984
Days to decision	140 days
Third-party review	No

APPLICANT

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