FDA 510(k) Premarket Notification

K834477 DISPOS. SCOTT CANNULAS 16-1054 ETC.

CLEARED

Mar 19, 1984
90 days to decision

K834477 · Product code: **HCA** · Neurology
Source: https://www.510kdatabase.net/k834477/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Ventricular (HCA)
Date received	Dec 20, 1983
Decision date	Mar 19, 1984
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Codman & Shurtleff, Inc.**
Location	Mchenry, IL, US
510(k) history	152 submissions · 151 cleared · 1976-2020