FDA 510(k) Premarket Notification

K841878 SHEATH REMOVER

CLEARED

Jul 13, 1984
67 days to decision

K841878 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k841878/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	May 7, 1984
Decision date	Jul 13, 1984
Days to decision	67 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019