FDA 510(k) Premarket Notification

K770569 CATHETER, EMBOLECTOMY, UMI

CLEARED

Apr 5, 1977
11 days to decision

K770569 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k770569/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Mar 25, 1977
Decision date	Apr 5, 1977
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Universal Medical Instrument Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1977-1996