FDA 510(k) Premarket Notification

K800622 EXTENSION LINE

CLEARED

Apr 29, 1980
42 days to decision

K800622 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k800622/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 18, 1980
Decision date	Apr 29, 1980
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Norton Performance Plastics Corp.**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1977-1985