FDA 510(k) Premarket Notification

K812999 PET POLYETHYLENE TORQUE CATHETER

CLEARED

Nov 16, 1981
21 days to decision

K812999 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k812999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Oct 26, 1981
Decision date	Nov 16, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

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