FDA 510(k) Premarket Notification

K830637 VERTEX DIGITAL SUBSTRATION KIT

CLEARED

Mar 24, 1983
23 days to decision

K830637 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k830637/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Mar 1, 1983
Decision date	Mar 24, 1983
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Professional Medical Services**
Location	Mchenry, IL, US
510(k) history	19 submissions · 18 cleared · 1983-1992