FDA 510(k) Premarket Notification

K830420 POLYETHYLENE MONITORING CONNECTORS

CLEARED

Feb 28, 1983
20 days to decision

K830420 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k830420/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Feb 8, 1983
Decision date	Feb 28, 1983
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Vertex Medical Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1982-1984