

K830637 VERTEX DIGITAL SUBSTRATION KITMar 24, 1983
23 days to decisionK830637 · Product code: **DQO** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830637/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026