

K800741 CENTRAL VEIN CATHETERIZATION KITApr 16, 1980
13 days to decisionK800741 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k800741/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Apr 3, 1980
Decision date	Apr 16, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Medical Components, Inc.
Location	Mchenry, IL, US
510(k) history	63 submissions · 55 cleared · 1980-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800741/> 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 26, 2026