

K840175 DOUBLE LUMEN INFUSION CATHETERSFeb 10, 1984
24 days to decisionK840175 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k840175/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 17, 1984
Decision date	Feb 10, 1984
Days to decision	24 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/k840175/> 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