

K820516 ARTERIAL CANALIZATION DEVICEApr 15, 1982
49 days to decisionK820516 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k820516/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 25, 1982
Decision date	Apr 15, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

Company	Alimed, Inc.
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1979-1996

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820516/> 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