

K864705 MODIFICATION (MATERIAL CHANGE) OF PEEL-AWAY SHEATHFeb 13, 1987
73 days to decisionK864705 · Product code: **DYB** · Cardiovascular
Source: <https://www.510kdatabase.net/k864705/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Dec 2, 1986
Decision date	Feb 13, 1987
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Quinton, Inc.
Location	Mchenry, IL, US
Contact	RANDY WALLS
510(k) history	164 submissions · 160 cleared · 1976-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k864705/>; 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 20, 2026