

K813143 DESILETNov 27, 1981
22 days to decisionK813143 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k813143/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Nov 5, 1981
Decision date	Nov 27, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Exco, Inc.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k813143/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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