

K812143 INTRODUCER, PERCUTANEOUSAug 18, 1981
21 days to decisionK812143 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k812143/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jul 28, 1981
Decision date	Aug 18, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Willson
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1980-1981

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Device record: <https://www.510kdatabase.net/k812143/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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