

K902135 PERCUTANEOUS INTRODUCER KITAug 1, 1990
83 days to decisionK902135 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k902135/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 10, 1990
Decision date	Aug 1, 1990
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Device Labs, Inc.
Location	Medway, MA, US
Contact	ELTON M TUCKER
510(k) history	6 submissions · 5 cleared · 1988-1993

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