

K013800 INSYTE, ANGIOCATH, INSYTE AUTOGUARD, ANGIOCATH AUTOGUARD, AUTOGUARD PRO, INTIMA, AND SAF-T-INTIMA INTRAVASCULAR CATHETE

Dec 21, 2001
36 days to decision

K013800 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k013800/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Nov 15, 2001
Decision date	Dec 21, 2001
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Becton Dickinson Infusion Therapy Systems, Inc.
Location	Sandy, UT, US
Contact	LESLIE WOOD
510(k) history	36 submissions · 36 cleared · 1997-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k013800/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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