

**K022099 SAFE-T-PEEL SAFETY NEEDLE/INTRODUCER,
MODELS 350-300 S,
360-300'S;S,380-300'S;S,390-300'S**

Sep 20, 2002
85 days to decision

K022099 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k022099/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jun 27, 2002
Decision date	Sep 20, 2002
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hdc Corp.
Location	Walker, MI, US
Contact	JONATHAN S KAHAN
510(k) history	30 submissions · 29 cleared · 1983-2007

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k022099/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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