

**K933050 CHANGES TO PROTECTIV(TM) & JELCO(TM) WINGED
CATH**Dec 14, 1993
208 days to decisionK933050 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k933050/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 20, 1993
Decision date	Dec 14, 1993
Days to decision	208 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Critikon Company, LLC
Location	Mchenry, IL, US
Contact	JOHN S SAMORAJCZYK
510(k) history	51 submissions · 51 cleared · 1979-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933050/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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