

**K905463 GENERAL MEDICAL PROCEDURE KIT/SKIN  
PREPARATION**Jan 22, 1991  
48 days to decisionK905463 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k905463/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Dec 5, 1990
Decision date	Jan 22, 1991
Days to decision	48 days
Third-party review	No

**APPLICANT**

---

Company	<b>Angiosystems, Inc.</b>
Location	Mchenry, IL, US
Contact	ALAN P SCHWARTZ
510(k) history	14 submissions · 11 cleared · 1984-1991

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905463/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026