FDA 510(k) Premarket Notification

K990390 HEPARIN IV FLUSH SYRINGE, 10 U/ML \

CLEARED

Dec 10, 1999
304 days to decision

K990390 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k990390/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Feb 9, 1999
Decision date	Dec 10, 1999
Days to decision	304 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Rocap Div. of Sabratek Corp.**
Location	Orlando, FL, US
Contact	AVIA TONEY
510(k) history	3 submissions · 3 cleared · 1999-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k990390/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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