FDA 510(k) Premarket Notification

K980062 MONOJECTT BLUNTIP I.V. ACCESS CANNULA WITH VIAL ACCESS PIN DEVICE

CLEARED

Mar 20, 1998
72 days to decision

K980062 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k980062/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 7, 1998
Decision date	Mar 20, 1998
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Davis & Geck, Inc.**
Location	Mchenry, IL, US
Contact	STEPHEN J TARNSETT
510(k) history	45 submissions · 42 cleared · 1981-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k980062/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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