

K990693 FLEXNEEDLE PLUSMay 14, 1999
72 days to decisionK990693 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k990693/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 3, 1999
Decision date	May 14, 1999
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Act Medical, Inc.
Location	Waltham, MA, US
Contact	WENDY V SHOTTS
510(k) history	2 submissions · 2 cleared · 1996-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990693/> 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 July 1, 2026