

K970255 KIPP MED I.V. MANIFOLDApr 14, 1997
82 days to decisionK970255 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k970255/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 22, 1997
Decision date	Apr 14, 1997
Days to decision	82 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	The Kipp Group
Location	Ontario, CA, US
Contact	J EDWARD GUILMETTE
510(k) history	6 submissions · 6 cleared · 1997-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970255/> 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 4, 2026