FDA 510(k) Premarket Notification

K970255 KIPP MED I.V. MANIFOLD

CLEARED

Apr 14, 1997
82 days to decision

K970255 · 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