FDA 510(k) Premarket Notification

K832542 I.V. ADMINISTRATION SET

CLEARED

Oct 31, 1983
94 days to decision

K832542 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k832542/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 29, 1983
Decision date	Oct 31, 1983
Days to decision	94 days
Third-party review	No

APPLICANT

Company	**Research Industries Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1983-1995