FDA 510(k) Premarket Notification

K833178 SOLUTION ADMINISTRATION LINE W/SPIKE

CLEARED

Nov 30, 1983
72 days to decision

K833178 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k833178/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 19, 1983
Decision date	Nov 30, 1983
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Intramed, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983