FDA 510(k) Premarket Notification

K821032 MODEL SR-1020LF

CLEARED

May 3, 1982
20 days to decision

K821032 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k821032/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 13, 1982
Decision date	May 3, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Stewart-Riess Labs**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982