FDA 510(k) Premarket Notification

K801004 FLOW MED 160

CLEARED

May 2, 1980
7 days to decision

K801004 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k801004/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 25, 1980
Decision date	May 2, 1980
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**D.R.O. Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1985