FDA 510(k) Premarket Notification

K833030 INTRAVASCULAR ADMINISTRATION SET

CLEARED

Nov 3, 1983
57 days to decision

K833030 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k833030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 7, 1983
Decision date	Nov 3, 1983
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Gish Biomedical, Inc.**
Location	Mchenry, IL, US
510(k) history	75 submissions · 75 cleared · 1983-2009