FDA 510(k) Premarket Notification

K894842 IMED GEMINI 12 NONVENTED Y-TYPE BLOOD/SOLUTION ADM

CLEARED

Sep 12, 1989
43 days to decision

K894842 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k894842/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 31, 1989
Decision date	Sep 12, 1989
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Imed Corp.**
Location	Mchenry, IL, US
Contact	JARBOE, PHD
510(k) history	43 submissions · 43 cleared · 1977-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k894842/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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