FDA 510(k) Premarket Notification

K944320 VENTED/NON-VENTED GEMINI ADMINISTRATION SET

CLEARED

Feb 14, 1995
169 days to decision

K944320 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k944320/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 29, 1994
Decision date	Feb 14, 1995
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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