FDA 510(k) Premarket Notification

K950153 VENTED/NONVENTED GEMINI ADMINISTRATION SET WITH CHECK VALVE AND TWO SAFSITE(TM) VALVES

CLEARED

Aug 24, 1995
219 days to decision

K950153 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k950153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 17, 1995
Decision date	Aug 24, 1995
Days to decision	219 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