FDA 510(k) Premarket Notification

K905219 4' LOW VOLUME IV EXTENSION SET

CLEARED

Jun 4, 1991
196 days to decision

K905219 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k905219/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 20, 1990
Decision date	Jun 4, 1991
Days to decision	196 days
Third-party review	No

APPLICANT

Company	**Marquette Medical, Inc.**
Location	Crofton, MD, US
Contact	MARQUETTE JR
510(k) history	17 submissions · 17 cleared · 1987-2003