FDA 510(k) Premarket Notification

K891176 INJECTION SITE PLUG (INTRAVASCULAR ADMINISTRATION)

CLEARED

May 18, 1989
73 days to decision

K891176 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k891176/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 6, 1989
Decision date	May 18, 1989
Days to decision	73 days
Third-party review	No

APPLICANT

Company	**Multi-Med, Inc.**
Location	West Swanzey, NH, US
Contact	ALAN P REID
510(k) history	7 submissions · 7 cleared · 1989-1989