FDA 510(k) Premarket Notification

K903541 SAFETY INJECTION PORT

CLEARED

Sep 14, 1990
38 days to decision

K903541 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k903541/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 7, 1990
Decision date	Sep 14, 1990
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Du-Call Miller Plastics, Inc.**
Location	Lisle, IL, US
Contact	W. C MILLER
510(k) history	1 submissions · 1 cleared · 1990-1990