FDA 510(k) Premarket Notification

K932446 INJECTION SITE

CLEARED

Nov 29, 1993
193 days to decision

K932446 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k932446/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 20, 1993
Decision date	Nov 29, 1993
Days to decision	193 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**3M Health Care, Ltd.**
Location	Irvine, CA, US
Contact	VON BUSCH
510(k) history	49 submissions · 49 cleared · 1990-1996