FDA 510(k) Premarket Notification

CLEARED

K964812 NIMA NEEDLELESS INJECTIONSITE MASTER ADAPTER AND I.V. SET

Feb 10, 1997
73 days to decision

K964812 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k964812/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 29, 1996
Decision date	Feb 10, 1997
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Critical Device Corp.**
Location	Brea, CA, US
Contact	DAN HYUN
510(k) history	6 submissions · 6 cleared · 1996-1999