

**K964812 NIMA NEEDLELESS INJECTION SITE MASTER
ADAPTER AND I.V. SET**Feb 10, 1997
73 days to decisionK964812 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k964812/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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