

**K964918 NIMA NEEDLELESS INJECTION SITE MASTER ADAPTER WITH POSIFLOW POSITIVE DISPLACEMENT FEATURE, AND I.V. SETS**

Apr 14, 1997  
126 days to decision

K964918 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k964918/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 9, 1996
Decision date	Apr 14, 1997
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k964918/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 5, 2026