

**K001106 MASTERSITE NEEDLELESS LUER ACTIVATED IV
CONNECTOR, MODEL MS 1000**Aug 16, 2000
133 days to decisionK001106 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k001106/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 5, 2000
Decision date	Aug 16, 2000
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Creative Plastic Technology
Location	Dana Point, CA, US
Contact	ROBERT J BARD
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001106/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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