

**K041399 HYPODERMIC NEEDLE-PRO EDGE NEEDLE
PROTECTION DEVICE**Jul 28, 2004
63 days to decisionK041399 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k041399/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 26, 2004
Decision date	Jul 28, 2004
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smiths Medical Asd, Inc.
Location	Rockland, MA, US
Contact	BRIAN D FARIAS
510(k) history	52 submissions · 52 cleared · 2004-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041399/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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