

K830010 SUPERJET LANCET

Jan 26, 1983

K830010 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k830010/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Jan 26, 1983
Decision date	Jan 26, 1983
Third-party review	No

APPLICANT

Company	Medprobe Laboratories
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k830010/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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