

K883957 ARTERIAL ANCHOR BANDAGE

Oct 18, 1988
29 days to decision

K883957 · Product code: **FMK** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k883957/>

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	Sep 19, 1988
Decision date	Oct 18, 1988
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Guidelines, Inc.
Location	Miami, FL, US
Contact	COHEN, PHD
510(k) history	1 submissions · 1 cleared · 1988-1988

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Device record: <https://www.510kdatabase.net/k883957/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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