

K831795 FIRST AID KITSSep 1, 1983
90 days to decisionK831795 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k831795/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Jun 3, 1983
Decision date	Sep 1, 1983
Days to decision	90 days
Third-party review	No

APPLICANT

Company	I M, Inc.
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1976-1992

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Device record: <https://www.510kdatabase.net/k831795/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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