

K832047 STERILE SUTURE REMOVAL SETSep 12, 1983
80 days to decisionK832047 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k832047/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Jun 24, 1983
Decision date	Sep 12, 1983
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Ormed Mfg., Inc.
Location	Mchenry, IL, US
510(k) history	23 submissions · 23 cleared · 1983-1994

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Device record: <https://www.510kdatabase.net/k832047/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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