

K761193 SUTURE REMOVAL KITJan 12, 1977
37 days to decisionK761193 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k761193/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Dec 6, 1976
Decision date	Jan 12, 1977
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Superior Plastic Products Corp.
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1976-1985

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Device record: <https://www.510kdatabase.net/k761193/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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