

K821502 STERILE SUTURE REMOVAL KITJul 9, 1982
51 days to decisionK821502 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k821502/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	May 19, 1982
Decision date	Jul 9, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

Company	Pro-Pak
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k821502/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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