

K820423 SUTURE REMOVAL TRAYMar 10, 1982
22 days to decisionK820423 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820423/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Feb 16, 1982
Decision date	Mar 10, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Megaplast, Inc.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1982

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820423/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026