FDA 510(k) Premarket Notification

K820423 SUTURE REMOVAL TRAY

CLEARED

Mar 10, 1982
22 days to decision

K820423 · Product code: **KDD** · General & Plastic Surgery
Source: 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