FDA 510(k) Premarket Notification

K761193 SUTURE REMOVAL KIT

CLEARED

Jan 12, 1977
37 days to decision

K761193 · Product code: **KDD** · General & Plastic Surgery
Source: 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