FDA 510(k) Premarket Notification

K821502 STERILE SUTURE REMOVAL KIT

CLEARED

Jul 9, 1982
51 days to decision

K821502 · Product code: **KDD** · General & Plastic Surgery
Source: 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