FDA 510(k) Premarket Notification

K801908 SI-14

CLEARED

Nov 12, 1980
93 days to decision

K801908 · Product code: **KDC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801908/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Aug 11, 1980
Decision date	Nov 12, 1980
Days to decision	93 days
Third-party review	No

APPLICANT

Company	**Kmi Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980