FDA 510(k) Premarket Notification

K782092 ARMBOARD CVER, DISPOSABLE

CLEARED

Jan 10, 1979
27 days to decision

K782092 · Product code: **FSE** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k782092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Manual (FSE)
Date received	Dec 14, 1978
Decision date	Jan 10, 1979
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Anago, Inc.**
Location	Mchenry, IL, US
510(k) history	44 submissions · 42 cleared · 1979-1993