FDA 510(k) Premarket Notification

K801328 THE EQUALIZER

CLEARED

Jun 9, 1980
6 days to decision

K801328 · Product code: **MDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801328/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Surgical, General Use (MDM)
Date received	Jun 3, 1980
Decision date	Jun 9, 1980
Days to decision	6 days
Third-party review	No

APPLICANT

Company	**The Anspach Effort, Inc.**
Location	Mchenry, IL, US
510(k) history	60 submissions · 60 cleared · 1980-2022