FDA 510(k) Premarket Notification

K771330 DISP. GROUND PLATE

CLEARED

Sep 6, 1977
50 days to decision

K771330 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k771330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 18, 1977
Decision date	Sep 6, 1977
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Edward Weck, Inc.**
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992