FDA 510(k) Premarket Notification

K000778 ENTEC REFLEX WAND 55, MODEL E4055-01, E4045-01

CLEARED

May 3, 2000
54 days to decision

K000778 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k000778/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 10, 2000
Decision date	May 3, 2000
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Arthrocare Corp.**
Location	Mountain View, CA, US
Contact	BRUCE PROTHRO
Website	http://www.arthrocare.com/
510(k) history	112 submissions · 112 cleared · 1995-2016