FDA 510(k) Premarket Notification

K852355 SINGLE USE ARTHROSCOPY ELECTRODE

CLEARED

Sep 3, 1985
92 days to decision

K852355 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k852355/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 3, 1985
Decision date	Sep 3, 1985
Days to decision	92 days
Third-party review	No

APPLICANT

Company	**Aspen Laboratories, Inc.**
Location	Mchenry, IL, US
Contact	JOHNSON
510(k) history	55 submissions · 55 cleared · 1976-1998