FDA 510(k) Premarket Notification

K823923 MF 360B ELECTROSURGICAL UNIT

CLEARED

Feb 15, 1983
49 days to decision

K823923 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k823923/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 28, 1982
Decision date	Feb 15, 1983
Days to decision	49 days
Third-party review	No

APPLICANT

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