FDA 510(k) Premarket Notification

K780977 SINGLE USE BLADE ELECTRODE

CLEARED

Jun 22, 1978
13 days to decision

K780977 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k780977/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 9, 1978
Decision date	Jun 22, 1978
Days to decision	13 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k780977/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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