FDA 510(k) Premarket Notification

K780975 SINGLE USE NEEDLE ELECTRODE

CLEARED

Jun 22, 1978
13 days to decision

K780975 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k780975/

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