

K810101 WATER RESISTANT FOOTSWITCH W/CABLEJan 29, 1981
13 days to decisionK810101 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k810101/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 16, 1981
Decision date	Jan 29, 1981
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

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Device record: <https://www.510kdatabase.net/k810101/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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