

**K852683 ELECTROSURGICAL LATERAL RELEASE PROBE  
MOD ALR-100**

Aug 12, 1985  
48 days to decision

K852683 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k852683/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 25, 1985
Decision date	Aug 12, 1985
Days to decision	48 days
Third-party review	No

**APPLICANT**

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Company	<b>American Bentley</b>
Location	Mchenry, IL, US
Contact	DONALD A RAIBLE
510(k) history	59 submissions · 59 cleared · 1983-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k852683/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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