

**K914621 IONTOPHORESIS DEVICE, MODIFICATION**Dec 20, 1991  
100 days to decisionK914621 · Product code: **EGJ** · Physical MedicineSource: <https://www.510kdatabase.net/k914621/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Device, Iontophoresis, Other Uses (EGJ)
Date received	Sep 11, 1991
Decision date	Dec 20, 1991
Days to decision	100 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>lomed, Inc.</b>
Location	Salt Lake City, UT, US
Contact	ANNE T CARTER
510(k) history	17 submissions · 12 cleared · 1990-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914621/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026