

**K982668 PHORESOR II,MODEL PM900**Mar 2, 1999  
214 days to decisionK982668 · Product code: **EGJ** · Physical Medicine  
Source: <https://www.510kdatabase.net/k982668/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Iontophoresis, Other Uses (EGJ)
Date received	Jul 31, 1998
Decision date	Mar 2, 1999
Days to decision	214 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>lomed, Inc.</b>
Location	Salt Lake City, UT, US
Contact	W.TIM MILLER
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/k982668/> 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 28, 2026