

**K031950 BAYLIS PAIN MANAGEMENT GENERATOR, MODELS  
PMG-115-TD AND PMG-230-TD**Jul 17, 2003  
23 days to decisionK031950 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k031950/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 24, 2003
Decision date	Jul 17, 2003
Days to decision	23 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Baylis Medical Co., Inc.</b>
Location	Mississauga, CA
Contact	KRIS SHAH
510(k) history	28 submissions · 28 cleared · 1998-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031950/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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