

**K990041 UROSTYM BIOFEEDBACK AND STIMULATION PROBES**Aug 5, 1999  
211 days to decisionK990041 · Product code: **KPI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k990041/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Non-implantable, For Incontinence (KPI)
Date received	Jan 6, 1999
Decision date	Aug 5, 1999
Days to decision	211 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Laborie Medical Tech Corp.</b>
Location	Mississauga, CA
Contact	DALE COLEMAN
510(k) history	5 submissions · 5 cleared · 1995-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990041/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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