

**K970077 LIBERTY PLUS SYSTEM(PFS-300)/VAGINAL PROBE,  
LIBERTY PLUS(PFS-044)**May 23, 1997  
134 days to decisionK970077 · Product code: **KPI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k970077/>**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 9, 1997
Decision date	May 23, 1997
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Utah Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	NADEEM AHMED
510(k) history	38 submissions · 38 cleared · 1979-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970077/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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