

**K952573 RUSCH URODYNAMIC RECTAL BALLOON CATHETER**

Jun 15, 1995  
 10 days to decision

K952573 · Product code: **FEN** · Gastroenterology & Urology  
 Source: <https://www.510kdatabase.net/k952573/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Cystometric, Hydraulic (FEN)
Date received	Jun 5, 1995
Decision date	Jun 15, 1995
Days to decision	10 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rusch Intl.</b>
Location	Jeffrey, NH, US
Contact	NEIL R ARMSTRONG
510(k) history	43 submissions · 43 cleared · 1995-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k952573/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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