

**K033335 LITHOSTAR MODULARIS WITH SHOCKWAVE  
SYSTEM CPLUS**Nov 25, 2003  
40 days to decisionK033335 · Product code: **MMZ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k033335/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Table, Cystometric, Electric (MMZ)
Date received	Oct 16, 2003
Decision date	Nov 25, 2003
Days to decision	40 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	DEBBIE PEACOCK
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033335/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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