

**K771655 CURITY PEAKOMETER**Oct 14, 1977  
46 days to decisionK771655 · Product code: **FFG** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k771655/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Urine Flow Rate Measuring, Non-electrical, Disposable (FFG)
Date received	Aug 29, 1977
Decision date	Oct 14, 1977
Days to decision	46 days
Third-party review	No

**APPLICANT**

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Company	<b>Kendall Research Center</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1976-1980

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

Device record: <https://www.510kdatabase.net/k771655/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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