

**K810690 CURITY 100% SILICONE SAFETY INDICATOR**Apr 23, 1981  
41 days to decisionK810690 · Product code: **KOD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k810690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Mar 13, 1981
Decision date	Apr 23, 1981
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>The Kendal Co.</b>
Location	Mchenry, IL, US
510(k) history	63 submissions · 60 cleared · 1980-2000

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

Device record: <https://www.510kdatabase.net/k810690/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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