

**K810531 PERI-PATCH PERITONEAL CATHETER EXTENSION**Mar 24, 1981  
26 days to decisionK810531 · Product code: **KGZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k810531/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter (KGZ)
Date received	Feb 26, 1981
Decision date	Mar 24, 1981
Days to decision	26 days
Third-party review	No

**APPLICANT**

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Company	<b>Quinton, Inc.</b>
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003

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

Device record: <https://www.510kdatabase.net/k810531/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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