

**K840424 HEMOCATH REPAIR KIT**Mar 5, 1984  
33 days to decisionK840424 · Product code: **KNZ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k840424/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, A-v Shunt (KNZ)
Date received	Feb 1, 1984
Decision date	Mar 5, 1984
Days to decision	33 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/k840424/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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