

**K030456 VASC-ALERT**Jul 11, 2003  
150 days to decisionK030456 · Product code: **KOC** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k030456/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Feb 11, 2003
Decision date	Jul 11, 2003
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vasc-Alert, LLC</b>
Location	Chicago, IL, US
Contact	JOHN KENNEDY
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030456/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026