

**K911996 SURGICUTT-BLEEDING TIME DEVICE, MODIFICATION**Jul 17, 1991  
86 days to decisionK911996 · Product code: **JCA** · Hematology  
Source: <https://www.510kdatabase.net/k911996/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Bleeding Time (JCA)
Date received	Apr 22, 1991
Decision date	Jul 17, 1991
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>International Technidyne Corp.</b>
Location	Walker, MI, US
Contact	LES HEIMANN
510(k) history	47 submissions · 47 cleared · 1983-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911996/> 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 8, 2026