

**K882481 HEMOCHRON CONTROL ACT TEST TUBE  
NORMAL/ABNOR. I/II**Aug 30, 1988  
75 days to decisionK882481 · Product code: **GGN** · Hematology  
Source: <https://www.510kdatabase.net/k882481/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasma, Coagulation Control (GGN)
Date received	Jun 16, 1988
Decision date	Aug 30, 1988
Days to decision	75 days
Third-party review	No

**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/k882481/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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