

**K860122 HEMOTEC ACT HIGH RANGE ACTIVATED CLOTTING TIME**Apr 4, 1986  
80 days to decisionK860122 · Product code: **JBP** · Hematology  
Source: <https://www.510kdatabase.net/k860122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Activated Whole Blood Clotting Time (JBP)
Date received	Jan 14, 1986
Decision date	Apr 4, 1986
Days to decision	80 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Device Consultants, Inc.</b>
Location	N. Attleboro, MA, US
Contact	WILLIAM MORTON
510(k) history	14 submissions · 14 cleared · 1982-2002

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860122/> 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 May 19, 2026