

**K894466 HEPCON(R) CV WHOLE BLOOD COAGULATION CONTROLS**Sep 5, 1989  
49 days to decisionK894466 · Product code: **JPA** · Hematology  
Source: <https://www.510kdatabase.net/k894466/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Jul 18, 1989
Decision date	Sep 5, 1989
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Hemotec, Inc.</b>
Location	Mchenry, IL, US
Contact	BAUGH, PHD
510(k) history	15 submissions · 15 cleared · 1980-1990

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894466/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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