

**K884590 HEPCON/SYSTEM B-10 HEPARIN  
ASSAY/SCREEN/CARTRIDGES**Jan 10, 1989  
68 days to decisionK884590 · Product code: **KFF** · Hematology  
Source: <https://www.510kdatabase.net/k884590/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assay, Heparin (KFF)
Date received	Nov 3, 1988
Decision date	Jan 10, 1989
Days to decision	68 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/k884590/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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