

**K890844 MICROSEPT 60/20**Jun 7, 1989  
106 days to decisionK890844 · Product code: **CAK** · General Hospital  
Source: <https://www.510kdatabase.net/k890844/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Microfilter, Blood Transfusion (CAK)
Date received	Feb 21, 1989
Decision date	Jun 7, 1989
Days to decision	106 days
Third-party review	No

**APPLICANT**

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Company	<b>Arbor Technologies, Inc.</b>
Location	Ann Arbor, MI, US
Contact	DAWN I MOORE
510(k) history	10 submissions · 10 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890844/>; 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 22, 2026