

**K885017 HEMOSORBA CH-500, CH-350 AND CH-180**Mar 17, 1989  
102 days to decisionK885017 · Product code: **FLD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k885017/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Hemoperfusion, Sorbent (FLD)
Date received	Dec 5, 1988
Decision date	Mar 17, 1989
Days to decision	102 days
Third-party review	No

**APPLICANT**

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Company	<b>Asahi Medical Co., Ltd.</b>
Location	Walker, MI, US
Contact	ANDERSON
510(k) history	16 submissions · 16 cleared · 1983-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k885017/>; 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 June 30, 2026