

**K980386 HOSPAL MULTIFLOW 100, MULTIFLOW 100 KITS A0 (A0,A0/B AND A0/0), MULTIFLOW 100 KITS B22 (B22,B22/B, & B22/0)**Feb 24, 1999  
387 days to decisionK980386 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k980386/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Feb 2, 1998
Decision date	Feb 24, 1999
Days to decision	387 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gambro Healthcare</b>
Location	Oklahoma City, OK, US
Contact	JEFFREY R SHIDEMAN
510(k) history	13 submissions · 13 cleared · 1996-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980386/> 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 29, 2026