

**K953551 KNOW-RECIRC**Jun 21, 1996  
329 days to decisionK953551 · Product code: **MQS** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k953551/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Hemodialysis, Access Recirculation Monitoring (MQS)
Date received	Jul 28, 1995
Decision date	Jun 21, 1996
Days to decision	329 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>H&amp;H Control Systems</b>
Location	Jackson, MS, US
Contact	ROBERT HESTER
510(k) history	1 submissions · 1 cleared · 1996-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953551/> 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 July 1, 2026