

**K830836 Q.C. TESTER**Apr 18, 1983  
27 days to decisionK830836 · Product code: **LHI** · Microbiology  
Source: <https://www.510kdatabase.net/k830836/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Mar 22, 1983
Decision date	Apr 18, 1983
Days to decision	27 days
Third-party review	No

**APPLICANT**

---

Company	<b>Alltek Filtration, Inc.</b>
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830836/> 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