

**K874555 KEELER ACU 22XT OPHTHALMIC CRYO UNIT**Jan 29, 1988  
86 days to decisionK874555 · Product code: **HPS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k874555/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryophthalmic (HPS)
Date received	Nov 4, 1987
Decision date	Jan 29, 1988
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Keeler Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874555/>; 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 20, 2026