

**K894248 CONVERTER SYSTEM**Feb 27, 1990  
251 days to decisionK894248 · Product code: **HQE** · Ophthalmic  
Source: <https://www.510kdatabase.net/k894248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Jun 21, 1989
Decision date	Feb 27, 1990
Days to decision	251 days
Third-party review	No

**APPLICANT**

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Company	<b>Mira, Inc.</b>
Location	Waltham, MA, US
Contact	ROGER O'BRLEN
510(k) history	21 submissions · 21 cleared · 1984-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894248/>; 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 May 22, 2026