

**K904909 I/A VITROPHAGE MODEL YPR 2001**Jan 28, 1991  
89 days to decisionK904909 · Product code: **HQE** · Ophthalmic  
Source: <https://www.510kdatabase.net/k904909/>**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	Oct 31, 1990
Decision date	Jan 28, 1991
Days to decision	89 days
Third-party review	No

**APPLICANT**

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Company	<b>Visioncare Devices, Inc.</b>
Location	Redding, CA, US
Contact	CHET CRACCHIOLO
510(k) history	5 submissions · 5 cleared · 1991-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904909/>; 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 14, 2026