

**K897021 MARTIN-HENSLEY SEGMENT LENS**Jun 22, 1990  
185 days to decisionK897021 · Product code: **HPX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k897021/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (polymethylmethacrylate) (HPX)
Date received	Dec 19, 1989
Decision date	Jun 22, 1990
Days to decision	185 days
Third-party review	No

**APPLICANT**

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Company	<b>Bruce W. Martin O.D. and Edward Hensley</b>
Location	Gladwin, MI, US
Contact	MARTIN, OD
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k897021/>, 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 30, 2026