

**K840660 POSTERIOR CAPSULOTOMY CYSTITOME**Mar 16, 1984  
30 days to decisionK840660 · Product code: **HNY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k840660/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystotome (HNY)
Date received	Feb 15, 1984
Decision date	Mar 16, 1984
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Sharpoint, Inc.</b>
Location	Walker, MI, US
510(k) history	9 submissions · 9 cleared · 1982-1986

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Device record: <https://www.510kdatabase.net/k840660/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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