

**K831829 KARICKHOFF FLYING CORPUSCLE VIEWER**Oct 19, 1983  
134 days to decisionK831829 · Product code: **HIX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k831829/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Maxwell Spot, Ac-powered (HIX)
Date received	Jun 7, 1983
Decision date	Oct 19, 1983
Days to decision	134 days
Third-party review	No

**APPLICANT**

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Company	<b>Surgidev Corp.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k831829/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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