

**K883233 EV MONOCANALICULAR STENT (TM)**Aug 22, 1988  
21 days to decisionK883233 · Product code: **OKS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k883233/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Aug 1, 1988
Decision date	Aug 22, 1988
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>Eagle Vision, Inc.</b>
Location	Bethesda, MD, US
Contact	DONALD L MACKEEN
510(k) history	7 submissions · 7 cleared · 1988-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883233/>; 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 29, 2026