

**K990672 LACRIMAL INTUBATION SET,  
DACRYOCYSTORHINOSTOMY (DCR) SET**May 18, 1999  
77 days to decisionK990672 · Product code: **OKS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k990672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Mar 2, 1999
Decision date	May 18, 1999
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hurricane Medical</b>
Location	Bradenton, FL, US
Contact	DAVID A CLAPP
510(k) history	3 submissions · 3 cleared · 1999-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990672/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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