

**K953973 ANGIOVISION 1000**Nov 17, 1995  
86 days to decisionK953973 · Product code: **HKI** · Ophthalmic  
Source: <https://www.510kdatabase.net/k953973/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Aug 23, 1995
Decision date	Nov 17, 1995
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medivision Medical Imaging, Ltd.</b>
Location	Hopkinton, MA, US
Contact	RICHARD A DOMAINGUE
510(k) history	4 submissions · 4 cleared · 1995-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953973/> 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 May 20, 2026