

**K904113 OPERATIVE RECORDING CAMERA**Oct 30, 1990  
54 days to decisionK904113 · Product code: **IKO** · Physical MedicineSource: <https://www.510kdatabase.net/k904113/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hammer, Reflex, Powered (IKO)
Date received	Sep 6, 1990
Decision date	Oct 30, 1990
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Dynamics, Inc.</b>
Location	Englewood, CO, US
Contact	JO BREHM
510(k) history	18 submissions · 17 cleared · 1988-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904113/>; 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 22, 2026