

**K931974 AZ 92**Oct 8, 1993  
169 days to decisionK931974 · Product code: **IWE** · Radiology  
Source: <https://www.510kdatabase.net/k931974/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Apr 22, 1993
Decision date	Oct 8, 1993
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>A2j, Inc.</b>
Location	Houston, TX, US
Contact	JIM L.
510(k) history	4 submissions · 4 cleared · 1988-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931974/> 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