

**K923471 PLANMED CYTOGUIDE**Nov 24, 1992  
133 days to decisionK923471 · Product code: **IZH** · Radiology  
Source: <https://www.510kdatabase.net/k923471/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Mammographic (IZH)
Date received	Jul 14, 1992
Decision date	Nov 24, 1992
Days to decision	133 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Planmed OY</b>
Location	Helsinki, FI
Contact	ISMO SEPPA
510(k) history	20 submissions · 20 cleared · 1992-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923471/> 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 14, 2026