

**K991435 AII-PRO 100 PLUS AND ALL-PRO 200**May 20, 1999  
24 days to decisionK991435 · Product code: **IXW** · Radiology  
Source: <https://www.510kdatabase.net/k991435/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Processor, Radiographic-film, Automatic (IXW)
Date received	Apr 26, 1999
Decision date	May 20, 1999
Days to decision	24 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>All-Pro Imaging Corp.</b>
Location	Hicksville, NY, US
Contact	FREDERICK R FISCHER
510(k) history	4 submissions · 4 cleared · 1991-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991435/> 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 19, 2026