

**K882358 PULSED FLUOROSCOPIC CONTROL MODEL:PF-02**Oct 19, 1988  
134 days to decisionK882358 · Product code: IZI · Radiology  
Source: <https://www.510kdatabase.net/k882358/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Jun 7, 1988
Decision date	Oct 19, 1988
Days to decision	134 days
Third-party review	No

**APPLICANT**

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Company	<b>U.S. Imaging, Inc.</b>
Location	Greensburg, OH, US
Contact	MARTIN J KULIS
510(k) history	4 submissions · 4 cleared · 1988-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882358/>; 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 21, 2026