

**K933161 ULTRA-VISION SUPER RAPID INTENSIFYING SCREEN**Oct 7, 1993  
100 days to decisionK933161 · Product code: **EAM** · Radiology  
Source: <https://www.510kdatabase.net/k933161/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Jun 29, 1993
Decision date	Oct 7, 1993
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dupont Medical Products</b>
Location	Wilmington, DE, US
Contact	JEAN BARTLETT
510(k) history	28 submissions · 28 cleared · 1991-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933161/> 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