

**K890066 STAR 36**Mar 28, 1989  
78 days to decisionK890066 · Product code: **IZO** · Radiology  
Source: <https://www.510kdatabase.net/k890066/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, High-voltage, X-ray, Diagnostic (IZO)
Date received	Jan 9, 1989
Decision date	Mar 28, 1989
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Transworld X-Ray Corp.</b>
Location	Charlotte, NC, US
Contact	J. A PRINCEHORN
510(k) history	7 submissions · 7 cleared · 1982-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890066/>; 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