

**K010705 OPTIDOS**Mar 11, 2003  
732 days to decisionK010705 · Product code: **KXK** · Radiology  
Source: <https://www.510kdatabase.net/k010705/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Source, Brachytherapy, Radionuclide (KXK)
Date received	Mar 9, 2001
Decision date	Mar 11, 2003
Days to decision	732 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ptw-New York Corp.</b>
Location	Bellmore, NY, US
Contact	STEPHEN R SZEGLIN
510(k) history	8 submissions · 8 cleared · 1995-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010705/> 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 18, 2026