

**K193602 LV Liberty Vision Model 1 90Yttrium Brachytherapy Source**May 29, 2020  
158 days to decisionK193602 · Product code: **KXK** · Radiology  
Source: <https://www.510kdatabase.net/k193602/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Source, Brachytherapy, Radionuclide (KXK)
Date received	Dec 23, 2019
Decision date	May 29, 2020
Days to decision	158 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lv Liberty Vision Corporation</b>
Location	Portsmouth, NH, US
Contact	Paul T. Finger
510(k) history	1 submissions · 1 cleared · 2020-2020

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k193602/> 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 June 18, 2026