

**K163572 LV Liberty Vision Model 1 90 Yttrium Brachytherapy Source**Mar 15, 2017  
86 days to decisionK163572 · Product code: **KXK** · Radiology  
Source: <https://www.510kdatabase.net/k163572/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Source, Brachytherapy, Radionuclide (KXK)
Date received	Dec 19, 2016
Decision date	Mar 15, 2017
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Lv Liberty Vision Corp.</b>
Location	Portsmouth, NH, US
Contact	Paul T. Finger
510(k) history	1 submissions · 1 cleared · 2017-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k163572/> Data sourced from FDA 510(k) public records (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. - Generated May 16, 2026