

**K093144 LOUSEBUSTER**Dec 16, 2009  
72 days to decisionK093144 · Product code: **LJL** · General Hospital  
Source: <https://www.510kdatabase.net/k093144/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detectors And Removers, Lice, (including Combs) (LJL)
Date received	Oct 5, 2009
Decision date	Dec 16, 2009
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Larada Sciences</b>
Location	Salt Lake City, UT, US
Contact	PHIL TRIOLO
510(k) history	2 submissions · 2 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093144/> 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 21, 2026