

**K083206 LOUSEBUSTER**Mar 10, 2009  
130 days to decisionK083206 · Product code: **LJL** · General Hospital  
Source: <https://www.510kdatabase.net/k083206/>**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 31, 2008
Decision date	Mar 10, 2009
Days to decision	130 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/k083206/> 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 22, 2026