FDA 510(k) Premarket Notification

K093144 LOUSEBUSTER

CLEARED

Dec 16, 2009
72 days to decision

K093144 · Product code: **LJL** · General Hospital
Source: https://www.510kdatabase.net/k093144/

SUBMISSION DETAILS

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

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