FDA 510(k) Premarket Notification

K083206 LOUSEBUSTER

CLEARED

Mar 10, 2009
130 days to decision

K083206 · Product code: **LJL** · General Hospital
Source: https://www.510kdatabase.net/k083206/

SUBMISSION DETAILS

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

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