FDA 510(k) Premarket Notification

CLEARED

K081593 POWER ASPIRATOR, MODEL LS2 OR LS2DP

Jul 23, 2008
47 days to decision

K081593 · Product code: **QPB** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081593/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty For Removal (QPB)
Date received	Jun 6, 2008
Decision date	Jul 23, 2008
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medical Device Resource Corp.**
Location	Hayward, CA, US
Contact	MEL KIMSEY
510(k) history	2 submissions · 2 cleared · 2008-2009

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k081593/ 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 June 22, 2026