

**K092195 CUTERA MULTIMODAL CELLULITE PRODUCT
(CMMCD)**Oct 1, 2009
72 days to decisionK092195 · Product code: **NUV** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k092195/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Massager, Vacuum, Light Induced Heating (NUV)
Date received	Jul 21, 2009
Decision date	Oct 1, 2009
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cutera, Inc.
Location	Brisbane, CA, US
Contact	KATHY MAYNOR
Website	http://www.cutera.com/
510(k) history	31 submissions · 31 cleared · 2004-2025

Cutera, Inc. is a medical device manufacturer specializing in aesthetic and surgical laser systems. The company operates with a manufacturing facility in Brisbane, US, and maintains a global presence across North America, Europe, and Australia. Cutera has established a strong regulatory track record with the FDA. The company has received FDA 510(k) clearances from total submissions since its first clearance in 2004. The vast majority of its submissions focus on General & Plastic Surgery devices, reflecting the company's core expertise in this category. The most recent cle...