

K122217 CERTUS 140 2.45GHZ ABLATION SYSTEMDec 19, 2012
147 days to decisionK122217 · Product code: **NEY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k122217/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Ablation, Microwave And Accessories (NEY)
Date received	Jul 25, 2012
Decision date	Dec 19, 2012
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Neuwave Medical, Inc.
Location	Madison, WI, US
Contact	DAN KOSEDNAR
510(k) history	15 submissions · 15 cleared · 2010-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122217/> 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 May 26, 2026