FDA 510(k) Premarket Notification

K122217 CERTUS 140 2.45GHZ ABLATION SYSTEM

CLEARED

Dec 19, 2012
147 days to decision

K122217 · Product code: **NEY** · General & Plastic Surgery
Source: 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	Madision, WI, US
Contact	DAN KOSEDNAR
510(k) history	15 submissions · 15 cleared · 2010-2024