FDA 510(k) Premarket Notification

K070092 MODIFICATION TO VELASMOOTH, SHAPER

CLEARED

Jul 27, 2007
198 days to decision

K070092 · Product code: **NUV** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k070092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Vacuum, Light Induced Heating (NUV)
Date received	Jan 10, 2007
Decision date	Jul 27, 2007
Days to decision	198 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Syneron Medical, Ltd.**
Location	Yokneam Elite, IL
Contact	YONI IGER
510(k) history	35 submissions · 35 cleared · 2002-2017