FDA 510(k) Premarket Notification

K150824 TD-Wand

CLEARED

May 28, 2015
57 days to decision

K150824 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k150824/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 1, 2015
Decision date	May 28, 2015
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Tdm Surgitech, Inc.**
Location	Palm Harbor, FL, US
Contact	MICHAEL WEBER
510(k) history	1 submissions · 1 cleared · 2015-2015