FDA 510(k) Premarket Notification

K170610 PlasmaBlade T

CLEARED

Apr 13, 2017
43 days to decision

K170610 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k170610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 1, 2017
Decision date	Apr 13, 2017
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Advanced Energy**
Location	Portsmouth, NH, US
Contact	Lydia Sakakeeny
510(k) history	8 submissions · 8 cleared · 2014-2019