FDA 510(k) Premarket Notification

K120510 TRANSCEND

CLEARED

Jul 19, 2013
514 days to decision

K120510 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k120510/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 21, 2012
Decision date	Jul 19, 2013
Days to decision	514 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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