FDA 510(k) Premarket Notification

K131696 VESALIUS

CLEARED

Jan 31, 2014
235 days to decision

K131696 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k131696/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 10, 2013
Decision date	Jan 31, 2014
Days to decision	235 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Telea Electronic Engineering Srl**
Location	Imola (Bo), IT
Contact	TESTA MARISA
510(k) history	2 submissions · 2 cleared · 2014-2016