FDA 510(k) Premarket Notification

CLEARED

K122092 MEDRANGE ELECTROSURGICAL EXPANSION SYSTEM

Oct 23, 2012
99 days to decision

K122092 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k122092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 16, 2012
Decision date	Oct 23, 2012
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medrange Corporation**
Location	Los Angeles, CA, US
Contact	KUOFANG HUANG
510(k) history	2 submissions · 2 cleared · 2010-2012