FDA 510(k) Premarket Notification

K170899 27HJ713S

CLEARED

May 30, 2017
64 days to decision

K170899 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k170899/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Mar 27, 2017
Decision date	May 30, 2017
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lg Electronics**
Location	Austin, TX, US
Contact	Jinhwan Jun
Website	http://www.lg.com/
510(k) history	5 submissions · 5 cleared · 2014-2019

LG Electronics is a global consumer electronics manufacturer with a manufacturing facility in Austin, US. The company produces televisions, audio equipment, home appliances, computing devices, and climate control systems. LG Electronics has received FDA 510(k) clearances from total submissions. The company's cleared devices span radiology and general and plastic surgery categories, with additional activity in cardiovascular devices. The regulatory record spans from 2014 to 2019. This company is inactive in the FDA 510(k) clearance process, with no submissions recorded in ...