FDA 510(k) Premarket Notification

K031891 VIRTUOSAPH

CLEARED

Jul 29, 2003
46 days to decision

K031891 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k031891/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jun 13, 2003
Decision date	Jul 29, 2003
Days to decision	46 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Terumo Cardiovascular Systems Corp.**
Location	Elkton, MD, US
Contact	KIM AVES
510(k) history	43 submissions · 43 cleared · 2000-2015