FDA 510(k) Premarket Notification

K090902 SPIDER SINGLE PORT SURGICAL DEVICE, SUPPORT ARM ACCESSORY, MODEL 90001, 90002, 90003

CLEARED

Jul 22, 2009
112 days to decision

K090902 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k090902/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Apr 1, 2009
Decision date	Jul 22, 2009
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Transenterix, Inc.**
Location	Durham, NC, US
Contact	TAMMY CARREA
510(k) history	15 submissions · 15 cleared · 2009-2021