FDA 510(k) Premarket Notification

K101274 VASOVIEW HEMOPRO 2 ENDOSCOPIC VESSEL HARVESTING SYSTEM

CLEARED

Jun 11, 2010
36 days to decision

K101274 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k101274/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 6, 2010
Decision date	Jun 11, 2010
Days to decision	36 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Maquet Cardiovascular, LLC**
Location	San Jose, CA, US
Contact	MARK H SMITH
510(k) history	14 submissions · 14 cleared · 2008-2026