

K101274 VASOVIEW HEMOPRO 2 ENDOSCOPIC VESSEL HARVESTING SYSTEM

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k101274/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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