

K242688 Finiff SystemDec 20, 2024
105 days to decisionK242688 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k242688/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 6, 2024
Decision date	Dec 20, 2024
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Skigrab Co., Ltd.
Location	Seoul, KR
Contact	Oskar Lee
510(k) history	1 submissions · 1 cleared · 2024-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242688/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026