

K243176 BLESSING SystemMay 16, 2025
228 days to decisionK243176 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k243176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 30, 2024
Decision date	May 16, 2025
Days to decision	228 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cellah Medical Co., Ltd.
Location	Seoul, KR
Contact	Priscilla Chung
510(k) history	1 submissions · 1 cleared · 2025-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k243176/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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