

K252146 Multi-functional Facial Beauty Device (SKB-1703,SKB-1803,SKB-1909, SKB-2003 SKB-2109,SKB-2129,SKB-2208PRO, SKB-2209)Dec 23, 2025
167 days to decisionK252146 · Product code: **NFO** · Neurology
Source: <https://www.510kdatabase.net/k252146/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Jul 9, 2025
Decision date	Dec 23, 2025
Days to decision	167 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Shenzhen Siken 3D Technology Development Co., Ltd.
Location	Shenzhen, CN
Contact	Jing Quan Liu
510(k) history	6 submissions · 6 cleared · 2017-2025

REGULATORY CONSULTANT

Consulting firm	Feiyong Drug & Medical Consulting Technical Service Group
Contact	Bing Huang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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