

K231073 SkinStylus SteriLock® MicroSystem, Model Number MP1209SLJul 28, 2023
105 days to decisionK231073 · Product code: **QAI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k231073/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Microneedle Device (QAI)
Date received	Apr 14, 2023
Decision date	Jul 28, 2023
Days to decision	105 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Esthetic Medical, Inc.
Location	Long Beach, CA, US
Contact	Kristin Groop
510(k) history	2 submissions · 2 cleared · 2023-2026

REGULATORY CONSULTANT

Consulting firm	Contract In-House Counsel and Consultants,
Contact	Marc C Sanchez

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k231073/> 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 26, 2026