

**K240598 Maestro System (REF100)**Jun 3, 2024  
91 days to decisionK240598 · Product code: **FQO** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k240598/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	Mar 4, 2024
Decision date	Jun 3, 2024
Days to decision	91 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Moon Surgical</b>
Location	Paris, FR
Contact	Anne Osdoit
510(k) history	4 submissions · 4 cleared · 2022-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Daniel &amp; Daniel Consulting</b>
Contact	Michael Daniel

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.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240598/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 13, 2026