FDA 510(k) Premarket Notification

K240598 Maestro System (REF100)

CLEARED

Jun 3, 2024
91 days to decision

K240598 · Product code: **FQO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k240598/

SUBMISSION DETAILS

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

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

REGULATORY CONSULTANT

Consulting firm	**Daniel & Daniel Consulting**
Contact	Michael Daniel

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