FDA 510(k) Premarket Notification

K203710 Storz Medical MAGNETOLITH Muscle Stimulator

CLEARED

May 3, 2021
133 days to decision

K203710 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k203710/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Dec 21, 2020
Decision date	May 3, 2021
Days to decision	133 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Storz Medical AG**
Location	Tagerwilen, CH
Contact	Pavel Novak
510(k) history	3 submissions · 3 cleared · 2018-2021

REGULATORY CONSULTANT

Consulting firm	**Biomed Research, Inc.**
Contact	Michael Dayton

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