

K230459 C.C. Life 21Oct 19, 2023
240 days to decisionK230459 · Product code: **NFO** · Neurology
Source: <https://www.510kdatabase.net/k230459/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Feb 21, 2023
Decision date	Oct 19, 2023
Days to decision	240 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	21 Equipment S.R.L.
Location	Torino, IT
Contact	Dolores Moretto
510(k) history	2 submissions · 2 cleared · 2022-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230459/> 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