

K190795 Tetragraph Neuromuscular Transmission MonitorOct 18, 2019
204 days to decisionK190795 · Product code: **KOI** · Anesthesiology
Source: <https://www.510kdatabase.net/k190795/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Peripheral, Electric (KOI)
Date received	Mar 28, 2019
Decision date	Oct 18, 2019
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Senzime AB
Location	Uppsala, SE
Contact	Johanna Faris
510(k) history	2 submissions · 2 cleared · 2019-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k190795/> 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 June 1, 2026