

K260142 StimeIMD (SSMD) systemMay 27, 2026
127 days to decisionK260142 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k260142/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Stimulator, Muscle, Powered (IPF) |
| Date received | Jan 20, 2026 |
| Decision date | May 27, 2026 |
| Days to decision | 127 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Motion Informatics Ltd. |
| Location | Yarka, IL |
| Contact | Liora Levi |
| 510(k) history | 1 submissions · 1 cleared · 2026-2026 |

REGULATORY CONSULTANT

| | |
|-----------------|----------------------|
| Consulting firm | RA Regulation |
| Contact | Orly Maor |

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.netDevice record: <https://www.510kdatabase.net/k260142/> 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 6, 2026