

K241813 SI-TECHNOLOGY® SI-DESIS® X™ Sacroiliac Joint Fusion SystemAug 13, 2024
53 days to decisionK241813 · Product code: **OUR** · Orthopedic
Source: <https://www.510kdatabase.net/k241813/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Sacroiliac Joint Fixation (OUR) |
| Date received | Jun 21, 2024 |
| Decision date | Aug 13, 2024 |
| Days to decision | 53 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Si-Technology, LLC |
| Location | Fort Collins, CO, US |
| Contact | Donner Chris |
| 510(k) history | 3 submissions · 3 cleared · 2015-2025 |

REGULATORY CONSULTANT

| | |
|-----------------|----------------------------------|
| Consulting firm | Secure BioMed Evaluations |
| Contact | Justin Gracyalny |

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/k241813/> 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 14, 2026