

K973914 SENSE TECHNOLOGY INC. FRAS, SENSE TECHNOLOGY INC. PULSTARJun 30, 1998
259 days to decisionK973914 · Product code: **LXM** · Physical MedicineSource: <https://www.510kdatabase.net/k973914/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Oct 14, 1997
Decision date	Jun 30, 1998
Days to decision	259 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Sense Technology, Inc.
Location	Pittsburgh, PA, US
510(k) history	2 submissions · 2 cleared · 1994-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973914/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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