FDA 510(k) Premarket Notification

K973914 SENSE TECHNOLOGY INC. FRAS, SENSE TECHNOLOGY INC. PULSTAR

CLEARED

Jun 30, 1998
259 days to decision

K973914 · Product code: **LXM** · Physical Medicine
Source: 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