FDA 510(k) Premarket Notification

# K963812 LM BONE ANCHOR (ORTHOPEDICS)

**CLEARED**

Nov 12, 1996
64 days to decision

K963812 · Product code: **MBI** · Orthopedic
Source: https://www.510kdatabase.net/k963812/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Fastener, Fixation, Nondegradable, Soft Tissue (MBI) |
| Date received | Sep 9, 1996 |
| Decision date | Nov 12, 1996 |
| Days to decision | 64 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Li Medical Technologies, Inc.** |
| Location | Greenwich, CT, US |
| Contact | RHODEMAN LI |
| 510(k) history | 17 submissions · 17 cleared · 1993-2000 |