

**K041651 STRYKER LEIBINGER SKELETAL ANCHORING SYSTEM**Sep 30, 2004  
105 days to decisionK041651 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k041651/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jun 17, 2004
Decision date	Sep 30, 2004
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Stryker Leibinger</b>
Location	Kalamazoo, MI, US
Contact	WADE T RUTKOSKIE
510(k) history	12 submissions · 12 cleared · 1999-2005

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041651/> 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 19, 2026