FDA 510(k) Premarket Notification

K041651 STRYKER LEIBINGER SKELETAL ANCHORING SYSTEM

CLEARED

Sep 30, 2004
105 days to decision

K041651 · 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	**Stryker Leibinger**
Location	Kalamazoo, MI, US
Contact	WADE T RUTKOSKIE
510(k) history	12 submissions · 12 cleared · 1999-2005