FDA 510(k) Premarket Notification

K033767 DUAL TOP ANCHOR SYSTEM SCREWS

CLEARED

Feb 24, 2004
83 days to decision

K033767 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k033767/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 3, 2003
Decision date	Feb 24, 2004
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Jeil Medical Corporation**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	53 submissions · 53 cleared · 2002-2026