FDA 510(k) Premarket Notification

K070569 SM INTERNAL./EXTERNAL IMPLANT SYSTEM

CLEARED

Nov 8, 2007
253 days to decision

K070569 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k070569/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 28, 2007
Decision date	Nov 8, 2007
Days to decision	253 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dio Department Dsi, Inc.**
Location	Santa Fe Springs, CA, US
Contact	KENNY LIM
510(k) history	8 submissions · 8 cleared · 2006-2008