FDA 510(k) Premarket Notification

K061323 MODIFICATION TO RELIADENT DENTAL IMPLANT SYSTEM

CLEARED

May 23, 2007
377 days to decision

K061323 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k061323/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	May 11, 2006
Decision date	May 23, 2007
Days to decision	377 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bioinfera, Inc.**
Location	Beachwood, OH, US
Contact	CHAN Q WANG
510(k) history	2 submissions · 2 cleared · 2005-2007