FDA 510(k) Premarket Notification

K090174 TSI

CLEARED

Sep 14, 2009
234 days to decision

K090174 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k090174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 23, 2009
Decision date	Sep 14, 2009
Days to decision	234 days
Third-party review	No
Summary / Statement	Summary
Other names	ERI

APPLICANT

Company	**Oco Biomedical**
Location	Sterling, VA, US
Contact	JACK BLOOM
510(k) history	4 submissions · 4 cleared · 2008-2013