FDA 510(k) Premarket Notification

K000889 SENOJ IMPLANT SYSTEM

CLEARED

Apr 27, 2001
403 days to decision

K000889 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k000889/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 20, 2000
Decision date	Apr 27, 2001
Days to decision	403 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Senoj Biocare, Inc.**
Location	San Diego, CA, US
Contact	PENNY WOLLUM
510(k) history	1 submissions · 1 cleared · 2001-2001