FDA 510(k) Premarket Notification

K001219 MEMOGRAPH

CLEARED

Jun 21, 2000
65 days to decision

K001219 · Product code: **JEY** · Dental
Source: https://www.510kdatabase.net/k001219/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Apr 17, 2000
Decision date	Jun 21, 2000
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary
Other names	OSSTAPLE - ORAL AND MAXILLIOFACIAL PROCEDURES FOR THE MEMOGRAPH STAPLE SYSTEM

APPLICANT

Company	**Biomedical Ent., Inc.**
Location	San Antonio, TX, US
Contact	W. CASEY FOX
510(k) history	13 submissions · 13 cleared · 1996-2014