

K001219 MEMOGRAPHJun 21, 2000
65 days to decisionK001219 · Product code: **JEY** · DentalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001219/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026