

K021684 THE BIOPLATE RIGID FIXATION BONE PLATING SYSTEM FOR CRANIOMAXILLOFACIAL SURGERY

Jun 26, 2002
35 days to decision

K021684 · Product code: JEY · Orthopedic
Source: <https://www.510kdatabase.net/k021684/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	May 22, 2002
Decision date	Jun 26, 2002
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bioplate, Inc.
Location	Los Angeles, CA, US
Contact	CAROL E JONES
510(k) history	23 submissions · 23 cleared · 2000-2008

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k021684/> 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 20, 2026