

K023665 MODIFICATION TO BIOPLATE RIGID BONE PLATING SYSTEM FOR CRANIOMAXILLOFACIAL SURGERY

Nov 22, 2002
22 days to decision

K023665 · Product code: JEY · Dental
Source: <https://www.510kdatabase.net/k023665/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Bone (JEY)
Date received	Oct 31, 2002
Decision date	Nov 22, 2002
Days to decision	22 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/k023665/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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