

**K023810 MODIFICATION TO BIOPLATE RIGID FIXATION BONE  
PLATING SYSTEM FOR CRANIOMAXILLOFACIAL SURGERY**Dec 4, 2002  
19 days to decisionK023810 · Product code: JEY · Dental  
Source: <https://www.510kdatabase.net/k023810/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Bone (JEY)
Date received	Nov 15, 2002
Decision date	Dec 4, 2002
Days to decision	19 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bioplate, Inc.</b>
Location	Los Angeles, CA, US
Contact	JUDY SOKUA
510(k) history	23 submissions · 23 cleared · 2000-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023810/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 20, 2026