

**K020088 MODIFICATION OF THE BIOPLATE ZIP CRANIOTOMY
FIXATION SYSTEM**

Feb 8, 2002
29 days to decision

K020088 · Product code: **GXN** · Neurology
Source: <https://www.510kdatabase.net/k020088/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Cranioplasty, Preformed, Non-alterable (GXN)
Date received	Jan 10, 2002
Decision date	Feb 8, 2002
Days to decision	29 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/k020088/> 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