

K061089 MODULAR THUMB IMPLANT, MODELS 17600, 17601, 17602, 17603, 17596, 17597, 17598, 17599Jun 20, 2006
63 days to decisionK061089 · Product code: **KYI** · Orthopedic
Source: <https://www.510kdatabase.net/k061089/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Wrist, Carpal Trapezium (KYI)
Date received	Apr 18, 2006
Decision date	Jun 20, 2006
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Biopro, Inc.
Location	Port Huron, MI, US
Contact	DAVID MRAK
510(k) history	41 submissions · 35 cleared · 1987-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061089/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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