

**K902790 NAPA**Sep 19, 1990  
85 days to decisionK902790 · Product code: **LQZ** · DentalSource: <https://www.510kdatabase.net/k902790/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	Jun 26, 1990
Decision date	Sep 19, 1990
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>G-Ortho Lab</b>
Location	Honolulu, HI, US
Contact	PETER GEORGE
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902790/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 4, 2026