FDA 510(k) Premarket Notification

K902790 NAPA

CLEARED

Sep 19, 1990
85 days to decision

K902790 · Product code: **LQZ** · Dental
Source: https://www.510kdatabase.net/k902790/

SUBMISSION DETAILS

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

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