FDA 510(k) Premarket Notification

K802684 HELIOMAT

CLEARED

Nov 24, 1980
27 days to decision

K802684 · Product code: **EAY** · Dental
Source: https://www.510kdatabase.net/k802684/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Fiber Optic, Dental (EAY)
Date received	Oct 28, 1980
Decision date	Nov 24, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Vivadent (Usa), Inc.**
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1978-1986