FDA 510(k) Premarket Notification

K831647 DENTURE RELINING, REPAIR/REBASING RESIN

CLEARED

Jun 30, 1983
38 days to decision

K831647 · Product code: **EBI** · Dental
Source: https://www.510kdatabase.net/k831647/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	May 23, 1983
Decision date	Jun 30, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Dentsply Intl.**
Location	Walker, MI, US
510(k) history	279 submissions · 279 cleared · 1976-2013