FDA 510(k) Premarket Notification

K905302 DENTHESIVE

CLEARED

Feb 26, 1991
90 days to decision

K905302 · Product code: **KLE** · Dental
Source: https://www.510kdatabase.net/k905302/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Nov 28, 1990
Decision date	Feb 26, 1991
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Kulzer, Inc.**
Location	Mchenry, IL, US
Contact	SHARON PARKER
510(k) history	31 submissions · 31 cleared · 1981-1991