FDA 510(k) Premarket Notification

K962065 3M VITREBOND LIGHT CURE GLASS IONOMER/BASE

CLEARED

Jul 18, 1996
51 days to decision

K962065 · Product code: **EMA** · Dental
Source: https://www.510kdatabase.net/k962065/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Dental (EMA)
Date received	May 28, 1996
Decision date	Jul 18, 1996
Days to decision	51 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	AMY FOWLER
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025