

K962065 3M VITREBOND LIGHT CURE GLASS IONOMER/BASEJul 18, 1996
51 days to decisionK962065 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962065/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026