

**K030036 TRIPLE-CURE PASTE/PASTE RESIN-REINFORCED
GLASS IONOMER CEMENT**Mar 12, 2003
68 days to decisionK030036 · Product code: **DYH** · Dental
Source: <https://www.510kdatabase.net/k030036/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Adhesive, Bracket And Tooth Conditioner, Resin (DYH)
Date received	Jan 3, 2003
Decision date	Mar 12, 2003
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Puldent Corporation
Location	Watertown, MA, US
Contact	KENNETH J BERK
510(k) history	32 submissions · 32 cleared · 2002-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030036/> 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 July 1, 2026