FDA 510(k) Premarket Notification

K831900 MACLEAN-FOGG, DISP. BASE MOLD

CLEARED

Jul 28, 1983
44 days to decision

K831900 · Product code: **KER** · Pathology
Source: https://www.510kdatabase.net/k831900/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Embedding (KER)
Date received	Jun 14, 1983
Decision date	Jul 28, 1983
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**Polymer Technology Corp.**
Location	Mchenry, IL, US
510(k) history	23 submissions · 23 cleared · 1983-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k831900/ 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 May 14, 2026