

K964827 PROD. NO. 01-132 - SUTURE REMOVAL KITFeb 14, 1997
74 days to decisionK964827 · Product code: **MCZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k964827/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture Removal Kit (MCZ)
Date received	Dec 2, 1996
Decision date	Feb 14, 1997
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Continental Medical Laboratories, Inc.
Location	Waterford, WI, US
Contact	JOSEPH DUNN
510(k) history	8 submissions · 6 cleared · 1991-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k964827/> 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 25, 2026