

K903329 CUI ANATOMICAL SILICONE MALARSep 10, 1990
47 days to decisionK903329 · Product code: **LZK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k903329/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Malar (LZK)
Date received	Jul 25, 1990
Decision date	Sep 10, 1990
Days to decision	47 days
Third-party review	No

APPLICANT

Company	Cui Corp.
Location	Santa Barbara, CA, US
Contact	MORRIS SHERWOOD
510(k) history	7 submissions · 7 cleared · 1990-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k903329/>; 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 7, 2026