

K973572 KELOCOTE SCAR GEL AND KELOCOTE LASER GELOct 21, 1997
32 days to decisionK973572 · Product code: **MDA** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k973572/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Elastomer, Silicone, For Scar Management (MDA)
Date received	Sep 19, 1997
Decision date	Oct 21, 1997
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hanson Medical, Inc.
Location	Goleta, CA, US
Contact	ERIK HANSON
510(k) history	14 submissions · 14 cleared · 1997-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973572/> 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 6, 2026