FDA 510(k) Premarket Notification

K973572 KELOCOTE SCAR GEL AND KELOCOTE LASER GEL

CLEARED

Oct 21, 1997
32 days to decision

K973572 · Product code: **MDA** · General & Plastic Surgery
Source: 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