FDA 510(k) Premarket Notification

K861447 ALISUN -TYPE 1200, TYPE 1000/800 AND TYPE 700/900

CLEARED

May 13, 1986
26 days to decision

K861447 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k861447/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Apr 17, 1986
Decision date	May 13, 1986
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Southern Exposure, Inc.**
Location	Wauconda, IL, US
Contact	THOMAS G MASON
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k861447/ 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 8, 2026