FDA 510(k) Premarket Notification

K861570 ALPHA TAN 1000 AND ALPHA TAN 1200

CLEARED

May 13, 1986
15 days to decision

K861570 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k861570/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Apr 28, 1986
Decision date	May 13, 1986
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Midwest Marketing Service**
Location	Klemme, IA, US
Contact	NOEL KATTER
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k861570/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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