

K812543 TANARAMAOct 8, 1981
34 days to decisionK812543 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k812543/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Sep 4, 1981
Decision date	Oct 8, 1981
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Marketing Consultants of America
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k812543/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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