FDA 510(k) Premarket Notification

K812543 TANARAMA

CLEARED

Oct 8, 1981
34 days to decision

K812543 · Product code: **FTC** · General & Plastic Surgery
Source: 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