FDA 510(k) Premarket Notification

K860347 SUNLAMP MODELS 24N, 30R, 30RE, 37RE

CLEARED

Feb 12, 1986
14 days to decision

K860347 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k860347/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Jan 29, 1986
Decision date	Feb 12, 1986
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Gti, Inc.**
Location	East Lyme, CT, US
Contact	GERHARD FRANZ
510(k) history	2 submissions · 2 cleared · 1986-1992