FDA 510(k) Premarket Notification

K820123 SUNFLOWER

CLEARED

Mar 1, 1982
42 days to decision

K820123 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k820123/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Jan 18, 1982
Decision date	Mar 1, 1982
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Solana, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1982-1982