FDA 510(k) Premarket Notification

K810796 SUNTAN BOOTH

CLEARED

Aug 3, 1981
133 days to decision

K810796 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810796/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Mar 23, 1981
Decision date	Aug 3, 1981
Days to decision	133 days
Third-party review	No

APPLICANT

Company	**Gibson'S Mfg. Corp.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1981-1981