

K810796 SUNTAN BOOTHAug 3, 1981
133 days to decisionK810796 · Product code: **FTC** · General & Plastic SurgerySource: <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&apos;S Mfg. Corp.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k810796/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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