FDA 510(k) Premarket Notification

K801017 TANIT COMMERCIAL SUNTAN BOOTH

CLEARED

Jun 17, 1980
48 days to decision

K801017 · Product code: **FTC** · Radiology
Source: https://www.510kdatabase.net/k801017/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Apr 30, 1980
Decision date	Jun 17, 1980
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Tanit, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980