FDA 510(k) Premarket Notification

K842975 BROWNIE BOOTH BB1

CLEARED

Aug 17, 1984
18 days to decision

K842975 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k842975/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Jul 30, 1984
Decision date	Aug 17, 1984
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Weingart, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1984-1986