

K842975 BROWNIE BOOTH BB1Aug 17, 1984
18 days to decisionK842975 · Product code: **FTC** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k842975/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026