FDA 510(k) Premarket Notification

# K830993 AMPLIGREFFE

**CLEARED**

May 3, 1983
35 days to decision

K830993 · Product code: **GFD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k830993/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Dermatome (GFD) |
| Date received | Mar 29, 1983 |
| Decision date | May 3, 1983 |
| Days to decision | 35 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Prothia USA, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 5 submissions · 5 cleared · 1982-1983 |