

**K830993 AMPLIGREFFE**May 3, 1983  
35 days to decisionK830993 · Product code: **GFD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k830993/>**SUBMISSION DETAILS**

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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**

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Company	<b>Prothia USA, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830993/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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