

**K905046 DERMABRADER**Dec 19, 1990  
41 days to decisionK905046 · Product code: **GFE** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k905046/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Brush, Dermabrasion, Powered (GFE)
Date received	Nov 8, 1990
Decision date	Dec 19, 1990
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Aaron Medical Industries</b>
Location	St. Petersburg, FL, US
Contact	ROBERT SARON
510(k) history	46 submissions · 46 cleared · 1990-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905046/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026