

**K003978 AFX MICROWAVE GENERATOR, FLEX ABLATION WAND, LYNX ABLATION WAND, MODEL SERIES 1000, P/N 102006, P/N 102007**May 22, 2001  
151 days to decisionK003978 · Product code: **NEY** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k003978/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Ablation, Microwave And Accessories (NEY)
Date received	Dec 22, 2000
Decision date	May 22, 2001
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Afx, Inc.</b>
Location	Fremont, CA, US
Contact	NANCY NORRIS
510(k) history	2 submissions · 2 cleared · 2001-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k003978/>; 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 July 3, 2026