

**K982929 MAXAIR NASAL DILATOR SYSTEM**Sep 9, 1998  
20 days to decisionK982929 · Product code: **LWF** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k982929/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Nasal (LWF)
Date received	Aug 20, 1998
Decision date	Sep 9, 1998
Days to decision	20 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Hnl Technologies</b>
Location	San Mateo, CA, US
Contact	HANFORD N LOCKWOOD, JR.
510(k) history	1 submissions · 1 cleared · 1998-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982929/> 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 5, 2026