

**K982439 PROXIDERM, MODELS TN 460, TN 90 460, BK 460**Sep 8, 1998  
56 days to decisionK982439 · Product code: **MKY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k982439/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Skin Closure (MKY)
Date received	Jul 14, 1998
Decision date	Sep 8, 1998
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Progressive Surgical Products, Inc.</b>
Location	Westbury, NY, US
Contact	ROBERT ODDSEN
510(k) history	3 submissions · 3 cleared · 1994-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982439/> 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 4, 2026