

**K872003 KINDERWRAP**Jun 23, 1987  
28 days to decisionK872003 · Product code: **GAX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k872003/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tourniquet, Nonpneumatic (GAX)
Date received	May 26, 1987
Decision date	Jun 23, 1987
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>The Kinder Co.</b>
Location	Ypsilanti, MI, US
Contact	TIM KINDER
510(k) history	1 submissions · 1 cleared · 1987-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872003/>; 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 8, 2026