

K830467 TOURNIQUET/LATEX TOURNIQUETMar 31, 1983
44 days to decisionK830467 · Product code: **GAX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k830467/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tourniquet, Nonpneumatic (GAX)
Date received	Feb 15, 1983
Decision date	Mar 31, 1983
Days to decision	44 days
Third-party review	No

APPLICANT

Company	Primeline Industries, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k830467/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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