

K803142 CONPHAR VELCRO TOUNIGUERJan 13, 1981
32 days to decisionK803142 · Product code: **GAX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k803142/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tourniquet, Nonpneumatic (GAX)
Date received	Dec 12, 1980
Decision date	Jan 13, 1981
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Conphar, Inc.
Location	Mchenry, IL, US
510(k) history	122 submissions · 122 cleared · 1979-1982

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k803142/> Data sourced from FDA 510(k) public records (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. - Generated May 23, 2026