FDA 510(k) Premarket Notification

K803142 CONPHAR VELCRO TOUNIGUER

CLEARED

Jan 13, 1981
32 days to decision

K803142 · Product code: **GAX** · General & Plastic Surgery
Source: 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