FDA 510(k) Premarket Notification

K872003 KINDERWRAP

CLEARED

Jun 23, 1987
28 days to decision

K872003 · Product code: **GAX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k872003/

SUBMISSION DETAILS

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

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