FDA 510(k) Premarket Notification

K012632 TOURNIQUET CUFF

CLEARED

Oct 26, 2001
74 days to decision

K012632 · Product code: **KCY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k012632/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tourniquet, Pneumatic (KCY)
Date received	Aug 13, 2001
Decision date	Oct 26, 2001
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary
Other names	PNEUMATIC TOURNIQUET

APPLICANT

Company	**Medical Instruments Technology, Inc.**
Location	Alexandria, VA, US
Contact	JACK SPEER
510(k) history	5 submissions · 5 cleared · 1989-2002