

**K012632 TOURNIQUET CUFF**Oct 26, 2001  
74 days to decisionK012632 · Product code: **KCY** · General & Plastic SurgerySource: <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	<b>Medical Instruments Technology, Inc.</b>
Location	Alexandria, VA, US
Contact	JACK SPEER
510(k) history	5 submissions · 5 cleared · 1989-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k012632/> 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 20, 2026