

K813231 TOURNIQUETDec 14, 1981
27 days to decisionK813231 · Product code: **KCY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k813231/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tourniquet, Pneumatic (KCY)
Date received	Nov 17, 1981
Decision date	Dec 14, 1981
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Aspen Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1976-1998

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Device record: <https://www.510kdatabase.net/k813231/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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