FDA 510(k) Premarket Notification

K813231 TOURNIQUET

CLEARED

Dec 14, 1981
27 days to decision

K813231 · Product code: **KCY** · General & Plastic Surgery
Source: 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