

K833676 SURGICAL VESSEL DILATORS-CATHERIZA-Mar 19, 1984
157 days to decisionK833676 · Product code: **DWP** · CardiovascularSource: <https://www.510kdatabase.net/k833676/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, Surgical (DWP)
Date received	Oct 14, 1983
Decision date	Mar 19, 1984
Days to decision	157 days
Third-party review	No

APPLICANT

Company	Cimed Intl., Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1984-1984

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833676/> 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 July 8, 2026