

K810008 ALP-KIIJan 16, 1981
11 days to decisionK810008 · Product code: **DXQ** · CardiovascularSource: <https://www.510kdatabase.net/k810008/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Jan 5, 1981
Decision date	Jan 16, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

Company	Martin&apos;S Maine Exchange
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810008/> 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 June 29, 2026