FDA 510(k) Premarket Notification

K810008 ALP-KII

CLEARED

Jan 16, 1981
11 days to decision

K810008 · Product code: **DXQ** · Cardiovascular
Source: 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'S Maine Exchange**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1981