

**K061959 MYTECH / HAPPY LIFE / MAXIAIDS ARM BLOOD
PRESSURE MONITOR, HPL-301, HPL-302, HPL-310, 180310**Aug 22, 2006
42 days to decisionK061959 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k061959/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jul 11, 2006
Decision date	Aug 22, 2006
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mytech Technology Co., Ltd.
Location	Hsin-Chu City, TW
Contact	Ke-Min Jen
510(k) history	4 submissions · 4 cleared · 2003-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061959/> 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 28, 2026