

**K030221 MYTECH / HAPPY LIFE BLOOD PRESSURE MONITOR**Jul 25, 2003  
185 days to decisionK030221 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k030221/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jan 21, 2003
Decision date	Jul 25, 2003
Days to decision	185 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mytech Technology Co., Ltd.</b>
Location	Hsin-Chu City, TW
Contact	JEN KE-MIN
510(k) history	4 submissions · 4 cleared · 2003-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030221/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026