

**K043531 MODIFICATION TO ARM BLOOD PRESSURE  
MONITOR, MODELS KP-6821, KP-6822**Apr 21, 2005  
121 days to decisionK043531 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k043531/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Dec 21, 2004
Decision date	Apr 21, 2005
Days to decision	121 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>K-Jump Health Co., Ltd.</b>
Location	Mission Viejo, CA, US
Contact	DANIEL TSENG
510(k) history	26 submissions · 26 cleared · 1990-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k043531/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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