

**K260747 YP-710T Series NIBP Cuff**May 6, 2026  
61 days to decisionK260747 · Product code: **DXQ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k260747/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Mar 6, 2026
Decision date	May 6, 2026
Days to decision	61 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	YP-840T Series Disposable Cuff

**APPLICANT**

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Company	<b>Nihon Kohden Corporation</b>
Location	Tokyo, JP
Contact	Charlemagne Chua
510(k) history	19 submissions · 19 cleared · 2015-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Nihon Kohden America, LLC</b>
Contact	Charlemagne Chua

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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**510k Database** - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k260747/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 6, 2026