FDA 510(k) Premarket Notification

K260747 YP-710T Series NIBP Cuff

CLEARED

May 6, 2026
61 days to decision

K260747 · Product code: **DXQ** · Cardiovascular
Source: https://www.510kdatabase.net/k260747/

SUBMISSION DETAILS

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

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

REGULATORY CONSULTANT

Consulting firm	**Nihon Kohden America, LLC**
Contact	Charlemagne Chua

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