

**K180155 Arm Type Blood Pressure Monitor**Aug 3, 2018  
196 days to decisionK180155 · Product code: **DXN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k180155/>**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 19, 2018
Decision date	Aug 3, 2018
Days to decision	196 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Avita Corporation</b>
Location	New Taipei City, CN
Contact	Steven Jhou
Website	<a href="https://www.avita.com.tw">https://www.avita.com.tw</a>
510(k) history	26 submissions · 26 cleared · 2000-2026

Avita Corporation is a medical device manufacturer based in New Taipei City, China. The company specializes in home care and clinical monitoring devices. Avita has received FDA 510(k) clearances from total submissions since 2000. The company's cleared devices span cardiovascular monitoring, anesthesiology, and general hospital applications. The latest clearance was in 2026, demonstrating continued regulatory activity and product innovation. The company's product portfolio includes blood pressure monitors, pulse oximeters, infrared thermometers, nasal aspirators, nebulizer...

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