

K051703 CLEVER TD-3018A BLOOD PRESSURE MONITOR

Aug 12, 2005
49 days to decision

K051703 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k051703/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jun 24, 2005
Decision date	Aug 12, 2005
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Taidoc Technology Corporation
Location	Taipei, County, TW
Contact	SHU-MEI WU
510(k) history	123 submissions · 123 cleared · 2004-2024

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Device record: <https://www.510kdatabase.net/k051703/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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