

K223332 Task Force COREApr 12, 2023
163 days to decisionK223332 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k223332/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Oct 31, 2022
Decision date	Apr 12, 2023
Days to decision	163 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cnsystems Medizintechnik GmbH
Location	Europe, AT
Contact	Raphael Gunacker
510(k) history	3 submissions · 3 cleared · 2002-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223332/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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