

K230281 PICOCLAMPOct 23, 2023
264 days to decisionK230281 · Product code: **DXC** · CardiovascularSource: <https://www.510kdatabase.net/k230281/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Feb 1, 2023
Decision date	Oct 23, 2023
Days to decision	264 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Kono Seisakusho Co., Ltd.
Location	Bunkyo-Ku, JP
Contact	Aya Takagi
510(k) history	2 submissions · 2 cleared · 2020-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230281/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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