

K173107 Vios Central Station Monitor Software, Vios Central Server SoftwareJul 26, 2018
300 days to decisionK173107 · Product code: **DXJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k173107/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Sep 29, 2017
Decision date	Jul 26, 2018
Days to decision	300 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vios Medical, Inc.
Location	St. Paul, MN, US
Contact	Amit Patel
510(k) history	3 submissions · 3 cleared · 2015-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173107/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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