FDA 510(k) Premarket Notification

CLEARED

K173107 Vios Central Station Monitor Software, Vios Central Server Software

Jul 26, 2018
300 days to decision

K173107 · 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