

**K102135 VECTOR EVENT GRID ARCHITECTURE (VEGA)
SYSTEM**Oct 22, 2010
85 days to decisionK102135 · Product code: **MWI** · General Hospital
Source: <https://www.510kdatabase.net/k102135/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jul 29, 2010
Decision date	Oct 22, 2010
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nuvon, Inc.
Location	Washington, Dc, DC, US
Contact	JONATHAN S KAHAN
510(k) history	2 submissions · 2 cleared · 2010-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102135/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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