FDA 510(k) Premarket Notification

CLEARED

K241411 Welch Allyn Connex® Spot Monitor

Dec 20, 2024
217 days to decision

K241411 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k241411/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	May 17, 2024
Decision date	Dec 20, 2024
Days to decision	217 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	901058 Vital Signs Monitor Core (CSM)

APPLICANT

Company	**Welch Allyn, Inc.**
Location	Mchenry, IL, US
Contact	Beth Rice
Website	http://www.welchallyn.com/
510(k) history	111 submissions · 111 cleared · 1977-2025

Welch Allyn, Inc. is a medical device manufacturer based in McHenry, US. The company specializes in patient monitoring and diagnostic equipment for healthcare settings. Welch Allyn has maintained a strong FDA 510(k) regulatory record since 1977. The company has received FDA 510(k) clearances from total submissions. Cardiovascular monitoring devices represent the dominant category in recent clearances, including the Connex vital signs monitor series and central station systems. The company's latest clearance in 2025 demonstrates continued regulatory activity and product in...