

Continuous Vital Signs Monitoring

When Early Detection and Intervention Matter in the General Ward





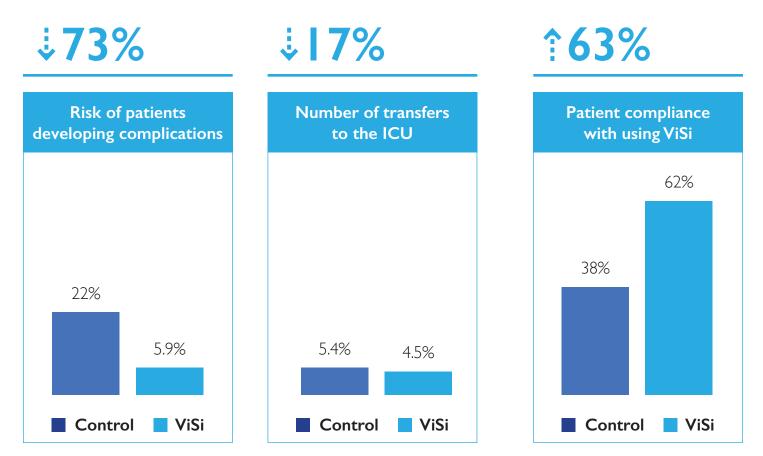
Current Monitoring Protocol

Advanced monitoring is standard in ICUs, yet nearly half of all adverse events in hospitalized patients occur on the general care ward^{1,2,3} where current monitoring protocols typically consist of intermittent spot checks by a nurse about every 4-8 hours. This leaves patients unmonitored for most of the time⁴ where subtle abnormalities in vital signs go undetected until complications arise. An alternative approach, studies have shown that continuous vital signs monitoring is associated with significant improvements in key clinical outcomes in patients treated in the general care ward.^{5,6}

ViSi Mobile bridges the gap between intermittent spot checking and ICU intensive monitoring. With continuous surveillance monitoring of key vital signs, early recognition and detection of patient deterioration enables clinicians to make timely and effective interventions.

See what ViSi Mobile did for a Mid-Atlantic medical center⁺

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An **"all-in-one"** vital signs Surveillance Monitoring for the General Ward



I – HR from ECG 2 – PR from SpO2 3 – PR from NIBP

4 – Posture

5 – NIBP Spot Check 6 – NIBP after 30 seco 7 – Continuous NIBP

6 – NIBP after 30 seconds

Flexible Configuration for any Level of Care

Basic Continuous Monitoring Continuous Vital Signs Monitoring

B

Basic monitoring includes:

• SpO2/Pulse Rate

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Proprietary SpO2 thumb sensor technology

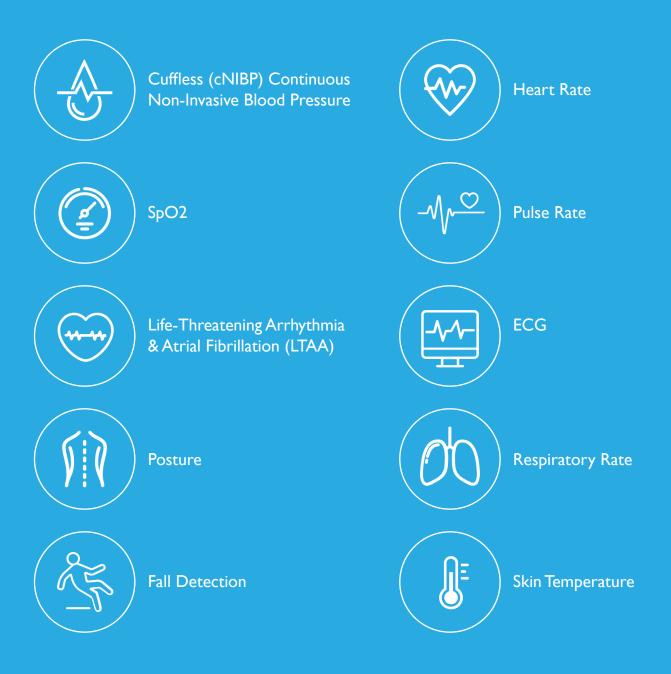
Continuous vital signs includes:

- Cuffless cNIBP (Continuous Non-Invasive Blood Pressure)
- LTAA
- SpO2/Pulse Rate
- RR and Skin Temperature
- ECG 3 and 5 Lead
- NIBP (Cuff-Based)
- Motion/Posture

All Vitals Matter. All the time.



FDA-cleared ViSi Mobile offers an "all-in-one," wearable, continuous multi-parameter monitoring designed for patients in the general care ward. With ViSi Mobile, early signs of deterioration can be detected, enabling timely interventions and preventing complications and adverse events.



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