

# Smart Ambulance Time-Series Data Generation and Artifact Detection

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## 1 Overview

This report documents the synthetic data generation and explicit artifact handling pipeline developed for the Smart Ambulance platform. The objective is to create realistic multi-signal physiological time-series data suitable for downstream anomaly detection and machine learning model development, while incorporating clinically plausible sensor artifacts encountered during ambulance transport.

The work addresses:

- **Task 1A:** Synthetic data generation for at least 30 minutes per patient.
- **Task 1B:** Explicit artifact detection and correction prior to anomaly detection.

## 2 Task 1A: Synthetic Data Generation

### 2.1 Signals Generated

The following physiological and contextual signals were generated at 1 Hz sampling:

- **SpO<sub>2</sub> (%)**: Peripheral oxygen saturation.
- **Heart Rate (HR, bpm)**: Beats per minute.
- **Systolic Blood Pressure (SBP, mmHg)**
- **Diastolic Blood Pressure (DBP, mmHg)**
- **Motion Signal (unitless)**: Proxy for vehicle vibration and patient movement.
- **Clinical Phase Labels**: NORMAL, DISTRESS, ACUTE.

### 2.2 Scenario Modeling

Each patient time-series covers approximately 30–35 minutes and includes:

- Normal transport phase with stable vitals.
- Distress phase with gradual physiological deterioration.
- Acute phase with clinically significant hypoxia and instability.
- Realistic transitions between phases.

### 2.3 Physiological Assumptions

- SpO<sub>2</sub> baseline: 96–100% in normal phase.
- Distress phase: gradual SpO<sub>2</sub> decline toward 90–93%.
- Acute phase: SpO<sub>2</sub> may reach 85–90%.
- HR increases with distress and acute phases.
- BP shows increased variability under stress.
- Slow physiological trends are modeled using low-frequency components and noise.

### 2.4 Artifact Injection

Explicit sensor artifacts were injected to simulate real-world EMS conditions:

- Motion-induced SpO<sub>2</sub> dropouts.
- Short-duration HR spikes from vehicle bumps.
- BP transient jumps.
- Short missing or corrupted signal segments.

Each injected artifact is labeled using ground-truth flags (e.g., `artifact_spo2`) to enable quantitative evaluation.

### 2.5 Limitations of Synthetic Data

- Does not fully capture complex patient-device interactions.
- Does not model probe repositioning dynamics.
- Noise statistics are approximate.
- Physiological variability across demographics is not explicitly modeled.

## 3 Task 1B: Explicit Artifact Detection and Cleaning

Artifact handling is implemented prior to anomaly detection to reduce false alarms caused by sensor errors.

### 3.1 Primary Cleaning Method (Final)

The final deployed method uses:

- Motion threshold gating.
- Sudden SpO<sub>2</sub> drop detection.
- Short-duration segment constraints.
- Rolling median physiological baselines.

A segment is classified as artifact if:

- Motion exceeds a defined threshold.
- SpO<sub>2</sub> drops rapidly below baseline.
- Segment duration is short (consistent with probe artifact).

Affected segments are replaced with rolling baseline estimates.

### 3.2 Additional Techniques Evaluated

Multiple techniques were tested during development:

- Rate-of-change thresholding
- Second derivative (acceleration) based detection
- Rebound-to-baseline checks
- Segment morphology constraints
- Window-tolerant evaluation buffers

These techniques were evaluated and compared quantitatively.

### 3.3 Evaluation Metrics

Artifact cleaning is evaluated against injected ground truth using:

- True Positives (TP)
- False Positives (FP)
- False Negatives (FN)
- Precision =  $TP / (TP + FP)$
- Recall =  $TP / (TP + FN)$

### 3.4 Evaluation Results

Technique	Precision	Recall	Notes
Motion + Drop + Segment (Final)	0.57	0.95	Balanced, robust
Second Derivative Method	~0.40	~0.50	Over-sensitive
Rebound-to-Baseline Check	~0.10	~0.08	Over-constrained
Window-Tolerant (2s buffer)	0.57	0.95	Allows near-artifact tolerance
Strict Pointwise (0s buffer)	1.00	0.95	Very strict matching

Table 1: Summary of artifact detection techniques and performance.

### 3.5 Interpretation of Buffer Results

Two evaluation modes were used:

- **Strict (0 second buffer):** Only exact artifact-labeled points count.
- **Window-tolerant (2 second buffer):** Near-artifact points are considered acceptable.

Results:

- Buffer = 0s: Precision = 1.00, Recall = 0.95
- Buffer = 2s: Precision = 0.57, Recall = 0.95

This indicates that some cleaned points occur adjacent to true artifact regions, which is clinically realistic for probe motion artifacts that smear across neighboring samples.

## 4 Before vs After Cleaning

### 4.1 SpO<sub>2</sub> Raw vs Cleaned Overlay

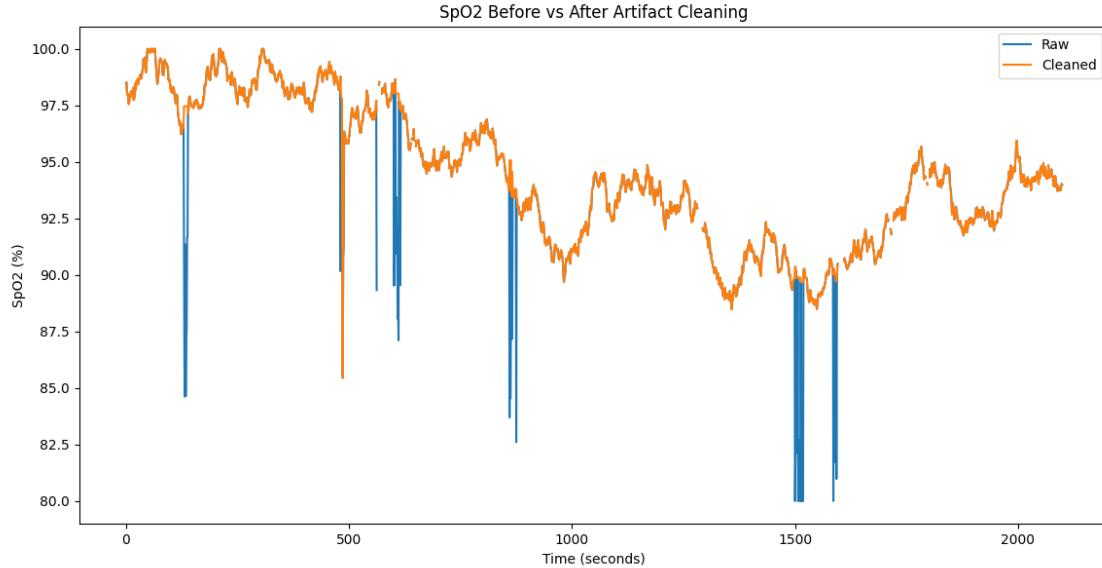


Figure 1: SpO<sub>2</sub> before and after artifact cleaning.

### 4.2 Clinical Safety Audit View

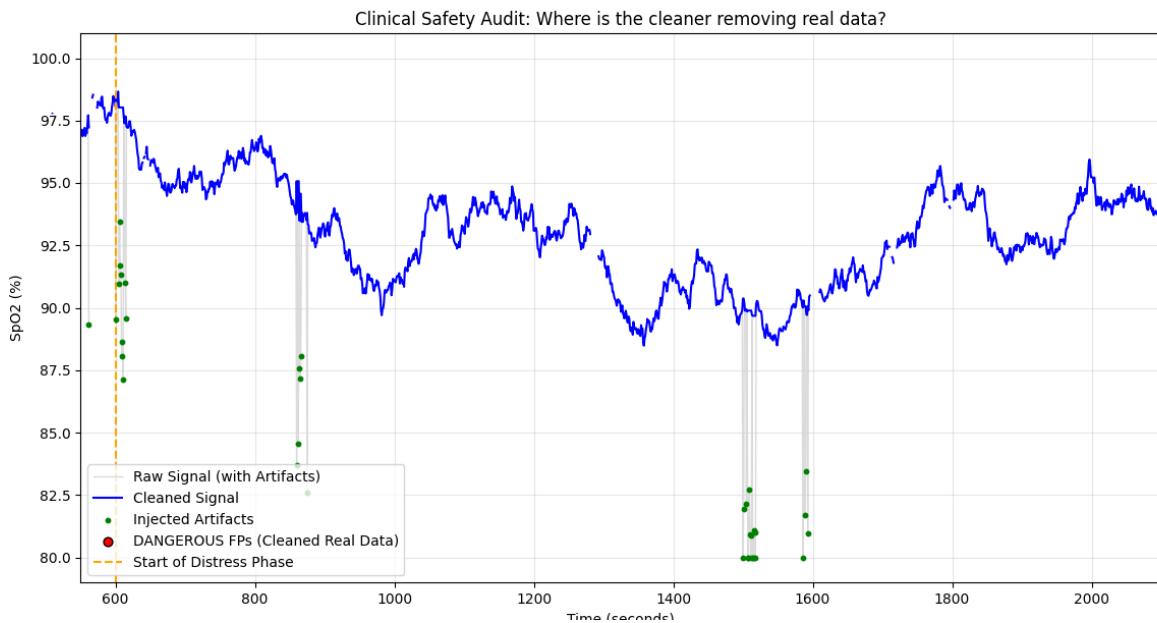


Figure 2: Clinical safety audit showing injected artifacts, cleaned points, and distress phase.

### 4.3 Raw and Cleaned SpO<sub>2</sub> (Separate)

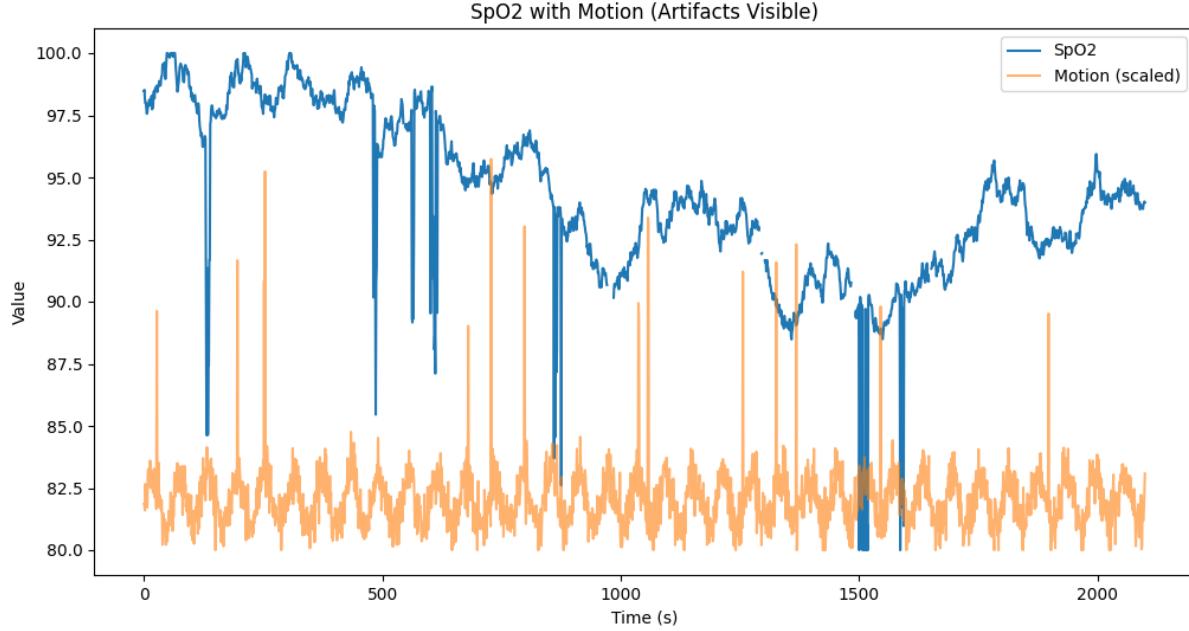


Figure 3: Raw SpO<sub>2</sub> with motion artifacts

## 5 Conclusion

A realistic synthetic physiological dataset was generated with labeled sensor artifacts. An explicit artifact handling pipeline was implemented using motion gating, physiological baselines, and segment-level logic.

The final method achieves high recall (0.95), ensuring most injected artifacts are detected, while maintaining moderate precision under window-tolerant evaluation. This balance is appropriate for preparing training data for downstream anomaly detection models in mobile EMS environments.

The generated dataset and artifact-cleaned signals are suitable for subsequent machine learning and clinical analytics development.