

OmniNet: A Multi-Modality Neural Network for Robust Remote Respiratory Rate Measurement from Facial Video

Tsai-Ni Lin^{1,2}

TIFFANY10022000@GMAIL.COM

An-Sheng Liu³

D00921006@NTU.EDU.TW

Li-Chen Fu^{*3}

LICHEN@NTU.EDU.TW

¹ *Department of Mechanical Engineering, National Taiwan University, Taiwan*

² *Department of Bioengineering, Hong Kong Science and Technology University, Hong Kong*

³ *Department of Electrical Engineering, National Taiwan University, Taiwan*

Editors: Under Review for MIDL 2026

Abstract

Remote respiratory rate (RR) measurement has gained traction in recent studies due to its ability to reduce healthcare professionals’ workload and patient discomfort. Recent studies have targeted this problem through remote photoplethysmography (rPPG) to capture subtle facial color changes. However, this technique is sensitive to lighting and motion variations. To this end, we propose OmniNet, a multimodal neural network that integrates image data processed through 3D convolutional neural networks (3D CNNs) with point of interest (POI) motion data and passes the fused features to Bidirectional Long Short-Term Memory (BiLSTM) to model long-term temporal dependencies. OmniNet achieves state-of-the-art performance by effectively capturing comprehensive spatial and temporal information while reducing illumination variation and motion-induced artifacts. It also requires fewer computational resources and enables faster inference compared to Transformer networks. The code has been released on GitHub: <https://anonymous.4open.science/r/spiro-7AFD>.

Keywords: Remote Respiratory Rate Measurement, Multimodal Learning

1. Introduction

The respiratory rate (RR) is a critical vital sign that offers predictive insight into a variety of health conditions, including obstructive sleep apnea, asthma, and other respiratory disorders. Traditional RR monitoring has relied on contact sensors (*e.g.* adhesive electrocardiogram (ECG) patches and respiratory belts) that continuously record cardiac electrical activity or thoracic expansion. However, contact devices not only increase clinical workload but also cause discomfort or skin irritation during prolonged monitoring, especially in infants (Ethawi et al., 2018) and patients with dermatological conditions or burns.

Recent research in remote photoplethysmography (rPPG), a non-contact technique that detects subtle skin color variations through a RGB camera, has proposed various algorithms to obtain physiological signals. This method has been applied to estimate RR (Chen et al., 2019), heart rate (HR) (Malasinghe et al., 2022; Premkumar and Hemanth, 2022), heart rate variability (HRV) (Poh et al., 2011), and other physiological indicators. Although rPPG shows great potential, its reliance on skin color makes it sensitive to lighting variation (Tarassenko et al., 2014) and prone to motion artifacts caused by head movements (Qiu et al., 2022; Bousefsaf et al., 2013).

* Corresponding author

Pixel-based RR techniques offer an alternative approach to estimate RR by directly tracking periodic pixel motion induced by thoracoabdominal respiration, providing greater robustness under varying illumination conditions. Among the representative techniques, optical flow detects motion by computing flow vectors of moving regions over time (Koolen et al., 2015; Mateu-Mateus et al., 2020), while temporal differencing calculates pixel-wise intensity changes between consecutive video frames (Liu et al., 2020; Bai et al., 2010). These techniques are hampered by background noise in complex environments, often hindering the accurate separation of respiratory motion from unrelated movements (Bai et al., 2019). Careful selection of the region of interest (ROI) has also been proven to raise the quality of respiratory signal. Commonly selected ROIs include the chest, abdomen (Janssen et al., 2015), and facial areas such as the forehead, cheeks, or nose (Mehta and Sharma, 2020).

Deep learning (DL) methods have emerged as powerful techniques to measure physiological signals. Convolutional neural networks (CNNs) have shown strong capabilities both in image understanding (Girshick et al., 2014; Redmon et al., 2016) and extracting accurate meaningful rPPG signals from low-quality facial videos. While 2D CNNs focus on spatial information (Liu et al., 2020; Chen and McDuff, 2018), 3D CNNs effectively model both global spatial and temporal information across frames (Yu et al., 2019; Ghezzi et al., 2024).

Transformers, first used in Natural Language Processing (NLP) for sequence modeling, better capture long-term dependencies than convolutional networks and are well suited for modeling the periodic nature of respiratory signals due to their self-attention mechanism. Although Transformers achieve high accuracy, they suffer from relatively high complexity (Chen et al., 2024), and their temporal attention can be inaccurate, leading to phase shifts and irrelevant attention (Yu et al., 2022, 2023). In camera-based physiological measurement, where data are more limited than in other vision tasks, transformers often perform worse than CNNs despite their strong modeling capabilities (Liu et al., 2023). Furthermore, Transformers require large-scale annotated facial videos to perform well, but such datasets are scarce and difficult to collect (Yue et al., 2023). CNNs are more effective in small-data settings but struggle to model long-term temporal dependencies.

Recurrent neural networks (RNNs) are widely used DL models well-suited for time series and sequential data. Long Short-Term Memory (LSTM) (Hochreiter and Schmidhuber, 1997) networks, a variant of RNNs, are effectively capture temporal dependencies and are often combined with CNNs for physiological signal estimation. Transformers require substantial training data, whereas LSTMs generally provide more efficient performance in environments with limited resources. While LSTMs are suitable for sequential modeling, their application to contactless RR estimation remains limited, as recent studies (Kumar et al., 2022; Rodrigues et al., 2024) have relied exclusively on contact-based datasets, without data derived from images or videos. Moreover, existing contactless methods often rely on single-modality input and fail to integrate complementary information sources.

To address these limitations, we propose a multimodal network named OmniNet. By incorporating multimodal input, our model minimizes the effect of noise from a single source, which improves the accuracy and generalizability of rPPG methods. OmniNet utilizes two types of information: frame differencing with a 3D CNN and POI motion trajectories. In this paper, we use a 3D CNN to capture temporal information, which is crucial for sequential signal extraction. Furthermore, the POI motion trajectories, inspired by OPOIRES (Deo Mehta and Sharma, 2023), is proposed to track meaningful points on

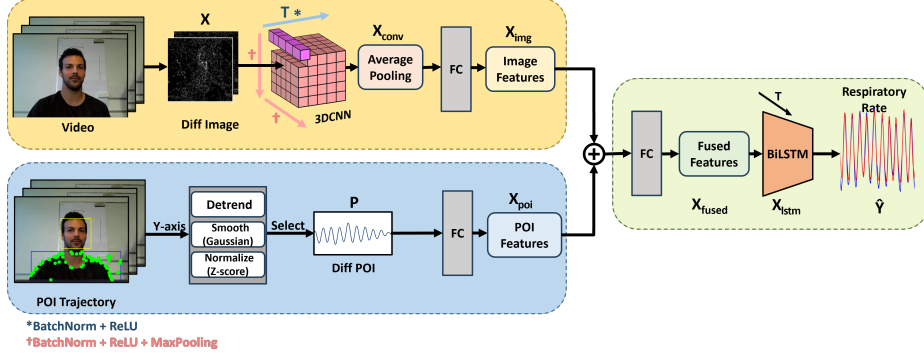


Figure 1: OmniNet framework.

the human torso rather than the fixed ROIs. Then we fuse these two types of features and feed them into a bidirectional LSTM (BiLSTM) to leverage its strength in modeling long-range temporal dependencies. We validate our approach on the COHFACE (Heusch et al., 2016) dataset against state-of-the-art methods, where it achieves superior performance. Moreover, the lightweight architecture of OmniNet ensures efficient deployment on mobile and embedded devices, emphasizing its relevance for practical real-world usage, especially in mobile healthcare and telemedicine.

2. Methodology

2.1. Overview of OmniNet Architecture

Illustrated in Figure 1, OmniNet is a multimodal DL model designed for robust respiratory signal estimation from facial videos under challenging conditions, *e.g.* illumination changes, head movements, and varying skin tones. Given a batch of input videos \mathbf{X} and POI-based motion signals \mathbf{P} , OmniNet adopts a dual-branch architecture consisting of a video stream and a POI stream. The video stream processes \mathbf{X} with a lightweight 3D CNN to extract motion-aware spatiotemporal features while preserving temporal resolution. These features are globally averaged across spatial dimensions and linearly projected to form the image embedding \mathbf{X}_{img} . In parallel, the POI stream passes \mathbf{P} through a linear projection layer to obtain \mathbf{X}_{poi} . The two modality-specific features are concatenated along the last dimension and fused via a ReLU-activated fully connected layer, yielding $\mathbf{X}_{\text{fused}}$. Temporal modeling is then performed using a single-layer BiLSTM. The resulting sequence \mathbf{X}_{lstn} is passed through a dropout layer and a final linear projection to estimate the respiratory signal $\hat{\mathbf{Y}}$.

2.2. POI Selection

Following the previous literature (Deo Mehta and Sharma, 2023), we adopt the Viola-Jones algorithm for face detection due to its lightweight and real-time performance. As shown in Figure 2, after the face detection, a rectangular region below the face is defined to approximate the chest area, where respiratory motion is most prominent.

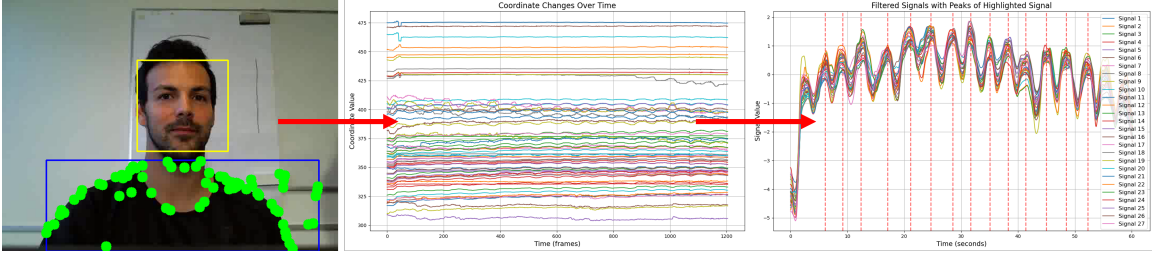


Figure 2: Visualization of POI selection: The system detects the face and defines an ROI below it, within which POIs are selected. The y -axis trajectories of these POIs are tracked over time, with each line representing the positional change of a single coordinate. Candidate signals are filtered after preprocessing, including detrending, smoothing, and normalization. Detected peaks are marked with red dashed lines indicate the estimated breathing cycles.

To enhance contrast, Contrast Limited Adaptive Histogram Equalization (CLAHE) and Gaussian smoothing are applied to the selected region. Harris corner detection is then performed to identify POIs, followed by sub-pixel refinement. A non-maximum suppression step with a minimum distance constraint enforces spatial diversity, retaining up to 30 points. These POIs are subsequently mapped to full-frame coordinates.

Only the y -axis displacement of each POI is preserved (Keall et al., 2006) after tracking POIs through frames using optical flow, resulting in one-dimensional temporal signals. Each signal is detrended, smoothed and standardized via z-score normalization:

$$\tilde{y}(t) = \frac{y(t) - \mu}{\sigma}, \quad (1)$$

where μ and σ denote the mean and standard deviation of $y(t)$. The normalized autocorrelation function is then computed and scaled by its maximum to ensure consistency across signals, where N denotes the length of the standardized signal $\tilde{y}(t)$:

$$R(\tau) = \sum_{t=\tau-N+1}^N \tilde{y}(t) \cdot \tilde{y}(t+N-\tau), \quad N \leq \tau \leq 2N-1. \quad (2)$$

Signals with fewer than 2 or more than 30 autocorrelation peaks and with a mean difference of peak intervals exceeding 70 are discarded. To further enhance robustness, a Pearson correlation matrix $C \in \mathbb{R}^{S \times S}$ is computed among the autocorrelation functions of S candidate signals, and the mutual similarity score for each is defined as:

$$L_j = \sum_{i=1}^S C(i, j), \quad 1 \leq j \leq S. \quad (3)$$

The top $\lceil S/2 \rceil$ signals with the highest L_j scores are retained as valid candidate signals.

2.3. Frame Differencing and Model Training

To extract RIM from video sequences, we employ a frame differencing strategy followed by a 3D CNN to capture both spatial and temporal dynamics of subtle respiratory movements.

Frame Differencing Given a facial video sequence $\mathbf{I}_1, \mathbf{I}_2, \dots, \mathbf{I}_T$ of T grayscale frames, we compute consecutive frame differences to highlight temporal motion: $\mathbf{X}_t = |\mathbf{I}_{t+1} - \mathbf{I}_t|$. This operation emphasizes motion features while reducing the impact of illumination changes. The resulting difference sequence \mathbf{X} serves as an intermediate representation of potential respiratory activity. To preserve the temporal structure, we stack the differences as a new input volume to the subsequent 3D CNN encoder. Similarly for POI, we compute consecutive sample differences: $\mathbf{P}_t = \tilde{y}(t+1) - \tilde{y}(t)$. Afterward, we sum up the differences of different POIs and perform z-score normalization on the temporal dimension, producing a difference sequence \mathbf{P} , followed by the same process as the video stream.

3D CNN Architecture To process the stacked difference volume, we adopt a lightweight 3D CNN encoder to model spatial and short-term temporal patterns. Specifically, the architecture consists of three convolutional blocks, each of which contains a 3D convolution layer with kernel size $3 \times 3 \times 3$, followed by BatchNorm3D and ReLU activation. The first two blocks also has a 3D max pooling with kernel size $1 \times 2 \times 2$ to downsample spatial dimensions while preserving the temporal axis. The output feature maps are then averaged across the spatial dimensions and applied a linear projection to obtain \mathbf{X}_{img} for temporal alignment with \mathbf{X}_{poi} from the POI stream. This design allows the model to capture fine-grained temporal cues from motion-only input, making it more robust to appearance variations, head pose changes, and illumination artifacts compared to directly analyzing RGB frames.

BiLSTM Architecture Following the fusion of image and POI features, we apply a single-layer BiLSTM. This allows the model to incorporate both past and future contexts for each time step. A dropout layer is applied to the LSTM output before a final fully connected layer maps each temporal feature vector to a scalar prediction.

Optimization Strategy We refer the reader to Appendix A for more details.

2.4. Peak Detection and RR Estimation

To estimate the RR, we first preprocess the predicted signal using Savitzky-Golay smoothing, followed by a fourth-order Butterworth band-pass filter with cutoff frequencies of 0.1–0.4 Hz to suppress noise and baseline drift. Peak detection is performed using the `find_peaks` signal processing functions build-in in the SciPy library, where a dynamic threshold is applied to exclude spurious local maxima. The minimum peak distance is set to 7 frames to avoid detecting peaks caused by motion artifacts.

If multiple valid peaks are found, we compute the pairwise inter-peak intervals and calculate the average breathing cycle. The RR (in breaths per minute) is then given by: $\text{RR} = 60/\bar{\Delta}$, where $\bar{\Delta}$ denotes the average peak interval in seconds. If fewer than two peaks are detected, the signal is excluded from the RR computation.

3. Experiments

3.1. Dataset and Experimental Setup

Due to the limited availability of RR datasets, we rely solely on the COHFACE (Heusch et al., 2016) dataset for our experiments.

COHFACE: The COHFACE dataset contains 160 one-minute RGB videos of 40 subjects (12 women, 28 men), recorded at 640×480 resolution and 20 Hz, with synchronized respiration and blood volume pulse signals at 256 Hz. Each subject contributed two videos under studio lighting and two under natural light. Official experimental splits are provided under three protocols: all, clean, and natural. Each protocol offers standard train/dev/test partitions for evaluating model robustness across varied illumination conditions.

We set the input window size to 240 frames, which corresponds to the length of both the differential image sequences and the respiration belt readings. The original 20 Hz video is temporally downsampled to 4 Hz by selecting one frame every five frames. This method aligns the visual input with the respiration belt signal, which is also downsampled to 4 Hz, enabling a more efficient calculation and facilitating synchronization between modalities.

We evaluated the performance of the model using three standard metrics: mean absolute error (MAE), root mean square error (RMSE), and Pearson correlation coefficient (PCC). Standard deviation of absolute error (STD) is also used when comparing with OPOIRES (Deo Mehta and Sharma, 2023). These metrics are described in Appendix B.

3.2. Intra-Database Tests

Table 1: Performance comparison between existing methods and our method on the COHFACE dataset.

Methods	MAE (bpm) ↓	RMSE (bpm) ↓	PCC ↑
DeepPhys (Chen and McDuff, 2018)	3.21	6.56	0.52
TS-CAN (Liu et al., 2020)	2.97	5.49	0.63
TS-DAN (Ren et al., 2021)	2.83	5.72	0.59
PhysNet (Yu et al., 2019)	2.31	4.77	0.67
EVM-MPP (Alnaggar et al., 2023)	1.63	2.10	0.45
PhysFormer (Yu et al., 2022)	1.44	2.29	0.62
ACTNet (Chen et al., 2024)	1.08	1.57	0.81
CliffPhys (Ghezzi et al., 2024)	0.83	1.97	0.86
OmniNet	0.24	0.42	0.99

All results are reported on the COHFACE test set. The CliffPhys model is pre-trained on the SCAMPS dataset and fine-tuned using the COHFACE training set. All other results, including TS-CAN, TS-DAN, PhysFormer, and ACTNet, are cited from ACTNet.

The results on the COHFACE dataset are shown in Table 1. We follow the official training and testing protocols provided by the dataset. OmniNet achieves the lowest MAE (0.24) and RMSE (0.42) and the highest PCC (0.99) on the COHFACE test set. These

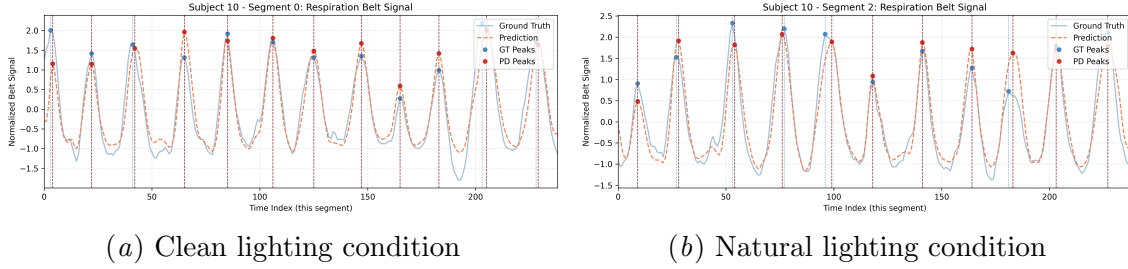


Figure 3: Comparison of predicted and ground truth respiration belt signals for Subject 10 using the model trained on the whole COHFACE dataset. Dots indicate peak positions of the ground truth and predicted signals.

results surpass all compared baselines, demonstrating the effectiveness of our framework for accurate RR estimation.

Table 2: Performance comparison of our method on the COHFACE dataset.

Train Set	Test Set	MAE (bpm) ↓	RMSE (bpm) ↓	PCC ↑
All	All	0.242	0.421	0.991
All	Clean	0.223	0.404	0.993
All	Natural	0.261	0.437	0.991
Clean	Clean	0.234	0.511	0.989
Natural	Natural	0.307	0.588	0.985

We further assess illumination robustness by training and testing OmniNet on the clean and natural subsets separately, reporting all results to three-decimal precision. As shown in Table 2, training on the full dataset (All) and testing on the clean subset achieves the lowest MAE (0.223) and RMSE (0.404) and the highest PCC (0.993). Although the clean subset yields slightly better performance, the differences across illumination conditions remain small, indicating that OmniNet is robust to lighting variations and performs well even under natural lighting.

Moreover, training on the whole dataset consistently outperforms training solely on a single subset, likely due to the increased diversity and volume of training data that improve the model’s generalizability. For example, the MAE reduces from 0.307 to 0.261 and the RMSE from 0.588 to 0.437 with the PCC increasing from 0.985 to 0.991 when switching from natural-subset training and testing to whole-set training and natural-subset testing. The consistent trend across MAE, RMSE, and PCC reinforces the effectiveness of using more comprehensive training data.

Figure 3 illustrates the respiration belt signals predicted by models trained on the whole dataset for a representative sample under clean and natural lighting conditions. The predicted curves closely follow the ground truth in the clean condition, with accurate peak

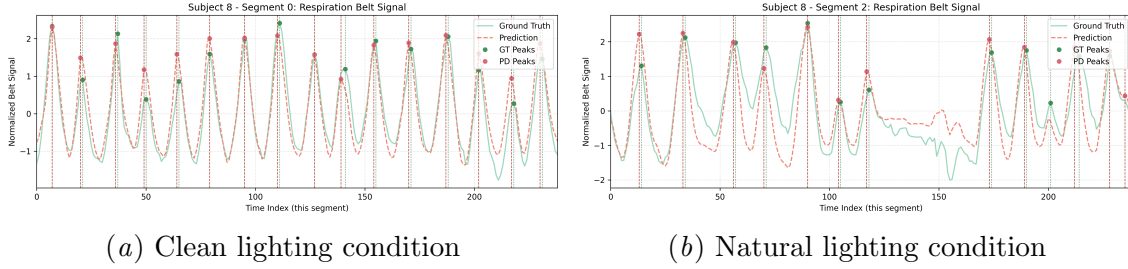


Figure 4: Comparison of predicted and ground truth respiration belt signals for Subject 8 using models trained and tested on the same condition of the COHFACE dataset under different lighting settings. Dots indicate peak positions of the ground truth and predicted signals.

detection, while predictions under natural lighting exhibit slight deviations. To further analyze the effect of training data quality, Figure 4 compares the results when training and testing solely on clean versus natural subsets, using a representative sample with notably divergent behavior. When trained on the natural subset, the model produces significant mismatches, including unstable fluctuations and incorrect peak detection. In contrast, training on the clean subset results in more accurate and stable predictions.

Since our method was inspired by the framework proposed in OPOIRES (Deo Mehta and Sharma, 2023), we conducted a direct comparison using their reported best setting (60-second input). For a fair comparison, we also adopted STD as one of the evaluation metrics, in addition to MAE and RMSE. All models are trained on the whole dataset and evaluated separately on the clean and natural test subsets.

Table 3: Performance comparison between OPOIRES and our method on the COHFACE two test set, using models trained on the whole dataset.

Test Set	Methods	MAE (bpm) ↓	RMSE (bpm) ↓	STD (bpm) ↓
Clean	OPOIRES (60s) (Deo Mehta and Sharma, 2023)	0.48	1.13	1.03
	OmniNet	0.22	0.40	0.34
Natural	OPOIRES (60s) (Deo Mehta and Sharma, 2023)	0.60	1.40	1.28
	OmniNet	0.26	0.44	0.35

As shown in Table 3, our method significantly outperforms OPOIRES. These results demonstrate the robustness and effectiveness of our multimodal design. Compared to relying solely on POI selection, incorporating multiple modalities leads to more accurate and stable RR estimation.

3.3. Ablation Studies

3.3.1. IMPACT OF 3D CNN

Table 4: Performance comparison under ablation settings on the COHFACE dataset, using models trained on the whole dataset.

Test Set	Methods	MAE (bpm) ↓	RMSE (bpm) ↓	PCC ↑
All	OmniNet	0.242	0.421	0.991
	OmniNet (<i>w/o 3D CNN</i>)	0.269	0.500	0.988
Clean	OmniNet	0.223	0.404	0.993
	OmniNet (<i>w/o 3D CNN</i>)	0.273	0.552	0.986
Natural	OmniNet	0.261	0.437	0.991
	OmniNet (<i>w/o 3D CNN</i>)	0.265	0.442	0.991

In this ablation study, we evaluated the impact of removing the 3D CNN module. As shown in the first two rows of Table 4, removing the 3D CNN results in only a slight performance drop on the whole dataset (All).

On both the natural and clean test sets, the full model consistently outperforms the model without 3D CNN. However, the changes are relatively small, indicating that the POI branch alone can still capture meaningful temporal features. These findings highlight the value of our multimodal architecture. While the POI branch offers robustness in noisy conditions, the integration of image-level features through 3D CNN provides complementary information that is especially beneficial in clean and stable environments. A detailed comparison of all POI-related hyperparameters is provided in Appendix C.

3.3.2. IMPACT OF POI SELECTION

Table 5: Performance comparison under ablation settings on the COHFACE dataset, using models trained on the whole dataset.

Test Set	Methods	MAE (bpm) ↓	RMSE (bpm) ↓	PCC ↑
All	OmniNet	0.242	0.421	0.991
	OmniNet (<i>w/o POI</i>)	0.744	1.191	0.939
Clean	OmniNet	0.223	0.404	0.993
	OmniNet (<i>w/o POI</i>)	0.707	1.126	0.949
Natural	OmniNet	0.261	0.437	0.991
	OmniNet (<i>w/o POI</i>)	0.782	1.252	0.931

For all these ablation experiments, all CNN-related hyperparameters are fixed to the best-performing configuration to ensure a fair comparison. As shown in Table 5, removing the POI modality leads to substantial performance degradation, causing much greater impact compared to removing 3D CNN. This may be because respiratory motion is highly localized and periodic, making POI features more effective in capturing relevant temporal patterns. In contrast, 3D CNN processes broader spatiotemporal regions, which are more susceptible to noise from lighting variations or non-respiratory motion. Nonetheless, even when either image or POI modality is ablated, OmniNet still outperforms all baselines. These results highlight the effectiveness and robustness of the model’s multimodal design, as well as the complementary nature of image and POI features.

3.4. Complexity Analysis

We also compared the complexity of the proposed method with the baselines, from which we can conclude that OmniNet achieved state of the art not only in performance but also in complexity. We refer the reader to Appendix D for more details.

4. Conclusion and future work

In this paper, we proposed OmniNet, a lightweight multimodal framework for remote respiratory rate (RR) estimation. Our method integrates pixel-based POI selection with a 3D CNN module using frame differencing and a BiLSTM decoder to model temporal dynamics. Through extensive experiments on the COHFACE dataset, OmniNet consistently outperforms existing methods under all conditions, particularly in clean data scenarios. Notably, our network has the smallest parameter size and computational complexity among all compared approaches, making it suitable for deployment on portable medical devices.

Despite its strong performance, our method has limitations. RR is a semi-voluntary signal that can be consciously controlled by subjects, potentially introducing noise in measurements. Nevertheless, we believe that OmniNet remains valuable in specific clinical contexts. It is particularly suitable for unconscious patients or individuals with special skin conditions, such as burn injuries or in newborns, where contact methods are unfeasible and remote visual monitoring is preferred.

In future work, we plan to: 1) validate the generalizability of OmniNet through cross-dataset experiments using any available RR datasets, 2) improve the pipeline by developing end-to-end architectures, and 3) extend the framework to comprehensive vital sign estimation in practical clinical environments.

Acknowledgments

This research was supported by the National Science and Technology Council & National Taiwan University, Taiwan, under the grant numbers 113-2634-F-002-002- , 113-2223-E-002-006- & 113-2221-E-002-127-MY3.

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Appendix A. Optimization Strategy

Our method is implemented using PyTorch and evaluated on a machine equipped with an NVIDIA GeForce RTX 4060 GPU. The model is trained with a batch size of 8. We use the Adam optimizer, and the learning rate is tuned per model to maximize performance, with 5×10^{-3} serving as the common starting point in most configurations. The loss function is the mean squared error (MSE) between the predicted respiratory belt signals $\hat{Y}(t)$ and the ground truth values $Y(t)$, defined as:

$$\mathcal{L}_{\text{MSE}} = \frac{1}{T} \sum_{t=1}^T \left(\hat{Y}(t) - Y(t) \right)^2, \quad (4)$$

where T denotes the number of temporal samples in each sequence. To stabilize training and prevent gradient explosion at the early stage, we adopt a linear learning rate schedule with warm-up. Specifically, the learning rate increases linearly during the first $N_{\text{warmup}} = 10$ steps, followed by a linear decay until the total number of training steps $N_{\text{total}} = 100$. The learning rate at step i is computed as $\eta_i \cdot \text{lr}_{\text{init}}$, where

$$\eta_i = \begin{cases} \frac{i}{N_{\text{warmup}}} & \text{if } i < N_{\text{warmup}}, \\ \frac{N_{\text{total}} - i}{N_{\text{total}} - N_{\text{warmup}}} & \text{otherwise.} \end{cases} \quad (5)$$

This scheduling strategy facilitates stable convergence by allowing sufficient exploration in the initial phase and finer adjustment in the later training stages. Early stopping is triggered after 10 epochs of no improvement. At each epoch, we save the best model based on the loss of the development set, which is used for all subsequent evaluations.

Appendix B. Evaluation Metrics

The four evaluation metrics are described as follows, where n is the number of samples, \hat{r}_i denotes the predicted RR, and r_i is the ground truth value for the i -th subject.

Mean absolute error (MAE) measures the average deviation between predicted and actual RR (in bpm):

$$\text{MAE} = \frac{1}{n} \sum_{i=1}^n |\hat{r}_i - r_i|. \quad (6)$$

Root mean square error (RMSE) quantifies overall prediction error magnitude:

$$\text{RMSE} = \sqrt{\frac{1}{n} \sum_{i=1}^n (\hat{r}_i - r_i)^2}. \quad (7)$$

Pearson correlation coefficient (PCC) indicates linear correlation between predicted and true values:

$$\text{PCC} = \frac{\sum_{i=1}^n (\hat{r}_i - \bar{\hat{r}})(r_i - \bar{r})}{\sqrt{\sum_{i=1}^n (\hat{r}_i - \bar{\hat{r}})^2} \cdot \sqrt{\sum_{i=1}^n (r_i - \bar{r})^2}}. \quad (8)$$

Standard deviation of absolute error (STD) measures the variability of the absolute errors:

$$\text{STD} = \sqrt{\frac{1}{n} \sum_{i=1}^n \left(|\hat{r}_i - r_i| - \frac{1}{n} \sum_{j=1}^n |\hat{r}_j - r_j| \right)^2}. \quad (9)$$

Appendix C. Analyses of POI Selection Parameters

We tuned the number of POIs (#POIs), quality level, and minimum distance (MD) in the POI stream, showing the results in Table 6. Based on these analyses, we fix the POI-related hyperparameters to their best-performing values and adopt this configuration as the default in all subsequent experiments.

Appendix D. Complexity Analysis

The number of parameters (Param.) and multiply-accumulate operations (MACs) are shown in Table 7. The parameters in the proposed method consist of two parts: the pre-trained models via AdaBoost in the Viola-Jones algorithm and the OmniNet, including 90K and 114K parameters, respectively. We omitted the MACs of the Viola-Jones and subsequent algorithms, as they are negligible compared to those of OmniNet, whose complexities are measured by `thop`. We can observe that OmniNet achieved the state of the art in both complexity and performance, demonstrating the excellence of the method.

Table 6: Analyses of POI-related hyperparameters on RR estimation performance on the whole COHFACE dataset.

#POIs	Quality level	MD	MAE (bpm) ↓	RMSE (bpm) ↓	PCC ↑
<i>Default configuration</i>					
100	10^{-4}	2	0.242	0.421	0.991
<i>Varying number of POIs, others fixed to default</i>					
50	10^{-4}	2	0.272	0.503	0.988
150	10^{-4}	2	0.255	0.518	0.987
<i>Varying quality level, others fixed to default</i>					
100	10^{-2}	2	0.355	0.723	0.976
100	10^{-3}	2	0.248	0.476	0.989
100	10^{-5}	2	0.264	0.523	0.987
<i>Varying minimum distance (MD), others fixed to default</i>					
100	10^{-4}	4	0.277	0.512	0.987
100	10^{-4}	6	0.273	0.575	0.984

Table 7: Comparison of the complexity between existing methods and our method.

Methods	Param. (M) ↓	MACs (G) ↓	MAE (bpm) ↓
DeepPhys (Chen and McDuff, 2018)	7.50	111.76	3.21
TS-CAN (Liu et al., 2020)	7.50	111.76	2.97
TS-DAN (Ren et al., 2021)	7.50	111.91	2.83
PhysNet (Yu et al., 2019)	0.73	65.19	2.31
PhysFormer (Yu et al., 2022)	7.03	47.01	1.44
ACTNet (Chen et al., 2024)	20.72	77.84	1.08
OmniNet	0.20	15.01	0.25

All results, including TS-CAN, TS-DAN, PhysFormer, and ACTNet, are cited from ACTNet.