# An Interpretable CNN-Transformer for Pediatric Sleep Apnea Diagnosis using ECG

Clara García-Vicente<sup>1,\*</sup>, Gonzalo C. Gutiérrez-Tobal<sup>1,2</sup>, Adrián Martín-Montero<sup>1,2</sup>, Jorge Jiménez-García<sup>1,2</sup>, Verónica Barroso-García<sup>1,2</sup>, David Gozal<sup>3</sup>, Roberto Hornero<sup>1,2</sup>

Abstract— Pediatric obstructive sleep apnea (OSA) is a common respiratory disorder that has been associated with increased cardiovascular risk. The standard diagnosis is polysomnography, but its complexity, cost, and inconvenience lead to underdiagnosis. To address this situation, we propose for the first time a simplified alternative using the overnight electrocardiogram (ECG) and a hybrid model based on a convolutional neural network and a transformer network to estimate the severity of pediatric OSA. In addition, the Gradient-weighted Class Activation Mapping (GradCAM) method is proposed to interpret the model results. For the development of the study, 2,591 recordings from the Childhood Adenotonsillectomy Trial (CHAT) and University of Chicago (UofC) databases were used. The model achieved a 4-class Cohen's Kappa of 0.392 in CHAT and 0.346 in UofC. GradCAM highlighted bradycardia-tachycardia patterns, and PQ and QT segments, as well as identified U waves. Therefore, this approach may improve the diagnosis of pediatric OSA and provide new related cardiac information, thus encouraging the adoption of automated systems in clinical settings.

**Keywords**— Pediatric obstructive sleep apnea, Transformer, eXplainable Artificial Intelligence, electrocardiogram.

### I. INTRODUCTION

Pediatric obstructive sleep apnea (OSA) is a common respiratory disorder that leads to changes in the cardiovascular (CV) system and increases CV risk [1]. The standard diagnostic method, polysomnography (PSG), is costly, complex, and uncomfortable, leading to under-diagnosis of children [1]. To address these complications, several studies have developed simplified strategies using a reduced number of PSG signals along with artificial intelligence techniques [1]. Focusing on pediatric OSA and cardiac signals, most studies have developed automated diagnostic alternatives using feature-engineering techniques [2]. Additionally, there is only one study that used a deep learning (DL) approach, which was based on a convolutional neural network (CNN) [3]. Despite the effectiveness shown in that study, CNNs are designed specifically to capture spatial features within the input data, disregarding the time dependencies present in sequences, such as those found in a nightly recording [4]. Another important limitation in all previous related literature is the lack of interpretability of the models used [5], thus hindering a higher acceptance of automatic models in clinical practice.

For these reasons, this study aims to assess the effectiveness of an interpretable hybrid DL network to identify spatial patterns through a CNN and temporal dependencies with a transformer (TF). Electrocardiogram (ECG) signals fed the CNN-TF model to establish OSA severity and uncover relevant ECG patterns associated with the disease.

## \*Corresponding Author: clara.garciav@uva.es

#### II. METHODS

#### A. Subjects and signals preprocessing

Signals were acquired from the semipublic Childhood Adenotonsillectomy Trial (CHAT, n=1610) [6] and a private University of Chicago (UofC, n=981) databases [7]. CHAT was randomly partitioned into training (60%), validation (20%), and testing (20%) sets, while UofC was reserved for independent model evaluation (only test). The databases presented annotations according to OSA severity (no OSA: AHI<1 event/hour; mild OSA: AHI<5 e/h; moderate OSA: AHI<5 e/h; and severe OSA: AHI>10 e/h). Table 1 presents clinical and demographic data from children.

The ECG-II lead from both databases underwent uniform preprocessing procedures. The signals were resampled at 100 Hz and the continuous component was corrected by subtracting the signal mean within 30-second windows. Subsequently, a high-pass filter with a cut-off frequency of 0.5 Hz was applied to mitigate noise. Finally, the amplitude of the signal was standardized using z-score normalization.

#### B. Deep learning architecture

A hybrid CNN-TF architecture was developed to capture both spatial structure and long-range relationships. The model received overnight ECG signals as input. The convolutional part corresponded to the previously presented CNN model [3]. Subsequently, TF was implemented using attentional mechanisms to capture dependencies and contextual information throughout the night sequence [8]. The output of the model was the apnea-hypopnea index (AHI) per subject.

#### C. Model interpretability using GradCAM

Gradient-weighted Class Activation Mapping (GradCAM) method was used as an eXplainable Artificial Intelligence (XAI) technique. Its function was to comprehend the intrinsic mechanisms of the model concerning the identification of apneic events and discerning cardiac patterns linked with pediatric OSA [9]. The GradCAM computation entailed using gradients derived from individual convolutional layers to compute layer-specific heatmaps. The final heatmap was obtained by averaging all the generated heatmaps [10].

#### **III. RESULTS AND DISCUSSION**

#### A. Diagnostic capability

Figure 1 shows the confusion matrices after the classification of the OSA severity. In the CHAT test set, the 4-class metrics obtained were  $Acc_4=61.2\%$  and kappa=0.392, whereas, in the UofC test set, the model obtained  $Acc_4=55.1\%$ 

Table 1: Demogra	aphic and	clinical	data from	CHAT	and U	JofC (	datasets
• /							

	CHA	АT		UofC
	Training	Validation	Test	Test
Subjects	988	323	299	981
Age (years)	7.00 [2.00]	7.00 [2.00]	6.90 [2.00]	6.0 [6.0]
Males (%)	51.72	49.23	46.15	61.37
BMI(kg/m <sup>2</sup> )	17.31[5.92]	17.12[6.25]	17.43[6.04]	18.02[5.86]
AHI (e/h)	2.64[4.77]	2.45[4.77]	2.32 [5.11]	3.8[7.76]
AHI≥1(e/h)	488	167	144	401
AHI≥5(e/h)	159	44	49	178
AHI≥10(e/h)	129	45	41	229

<sup>&</sup>lt;sup>1</sup>GIB Group, University of Valladolid, Spain

<sup>&</sup>lt;sup>2</sup>CIBER-BBN, Instituto de Salud Carlos III, Madrid, Spain

<sup>&</sup>lt;sup>3</sup>Joan C. Edwards, School of Medicine, Marshall University, Huntington, WV.

and *kappa*=0.346. In addition, Table 2 shows that the highest *Acc* is obtained for identifying the most severe children in both databases. Our approach demonstrated higher diagnostic performance in detecting pediatric OSA compared to prior studies, especially in severe children [7].

### B. Identification of ECG patterns

In Figure 2, GradCAM reveals heart rate alterations in the form of bradycardia-tachycardia patterns. In addition, heatmaps highlight U waves and PQ and QT segments. The CNN model utilizes information from bradycardia-tachycardia patterns, consistent with the physiological response of the heart to apneic events [11]. The inclusion of P, Q, and T-wave data also aligns with the observed dispersion of the P-wave and QT interval in severe pediatric OSA patients, indicating potential consequences for the onset of atrial fibrillation and elevated risk of sudden death [12]. The U-wave presence may be associated with long QT syndrome and bradycardia, both clinical factors linked to pediatric OSA [11]. Thus, heatmaps revealed ECG patterns related to OSA and potential indicators of CV risk.

#### **IV. CONCLUSIONS**

The integration of an ECG-interpretable CNN-TF model enables reliable diagnosis of pediatric OSA. In addition, GradCAM facilitates the identification of disease-related cardiac patterns. This approach may be a useful starting point when assessing the risk of CV comorbidities, a clinically relevant issue in pediatric OSA. In conclusion, this approach is an attractive alternative for PSG, as it offers a faster, more objective, and cost-effective method to diagnose OSA.

#### ACKNOWLEDGMENT

This research was funded by MCIN/AEI10.13039/501100011033/', ERDF, and 'NextGenerationEU/PRTR'(PID2020-115468RB-I00, PDC2021-120775-I00, CPP2022-009735) and by 'CIBER-BBN (CB19/01/00012)'. C. García-Vicente received an "Ayudas para contratos predoctorales para la Formación de Doctores" grant from the MCIN (PRE2021-100792). D. Gozal is supported in part by NIH grants G061824 and HL16617.



AHI	Test	Se	Sp	PPV	NPV	$LR^+$	LR-	Acc
cutoff	set	(%)	(%)	(%)	(%)			(%)
1 e/h	CHAT	94.9	29.2	82.8	61.3	1.3	0.2	80.6
	UofC	95.7	14.5	83.9	41.7	1.1	0.3	81.4
5 e/h	CHAT	75.6	92.3	81.0	89.8	9.9	0.3	87.3
	UofC	73.5	84.7	77.3	81.8	4.8	0.3	80.0
10 e/h	CHAT	61.0	96.1	71.4	93.9	15.7	0.4	91.3
	UofC	59.4	97.3	87.2	88.7	22.3	0.4	88.5

 $LR^+$  and  $LR^-$  (positive and negative likelihood ratio).

#### References

P. Moridian *et al.*, "Automatic diagnosis of sleep apnea from biomedical signals using artificial intelligence techniques: Methods, challenges, and future works," *WIREs Data Min. Knowl. Discov.*, vol. 12, no. 6, Nov. 2022.
 A. Martín-Montero *et al.*, "Pediatric sleep apnea: Characterization of apneic events and sleep stages using heart rate variability," *Comput. Biol. Med.*, vol. 154, p. 106549, Mar. 2023.

[3] C. García-Vicente *et al.*, "ECG-based convolutional neural network in pediatric obstructive sleep apnea diagnosis," *Comput. Biol. Med.*, vol. 167, no. September, p. 107628, Dec. 2023.

[4] I. Goodfellow et al., Deep learning. MIT Press, 2016.

[5] G. C. Gutiérrez-Tobal *et al.*, "Reliability of machine learning to diagnose pediatric obstructive sleep apnea: Systematic review and meta-analysis," *Pediatr. Pulmonol.*, vol. 57, no. 8, pp. 1931–1943, Aug. 2022.

[6] C. L. Marcus *et al.*, "A Randomized Trial of Adenotonsillectomy for Childhood Sleep Apnea," *N. Engl. J. Med.*, vol. 368, no. 25, pp. 2366–2376, Jun. 2013.

[7] R. Hornero et al., "Nocturnal Oximetry-based Evaluation of Habitually Snoring Children.," Am. J. Respir. Crit. Care Med., vol. 196, no. 12, pp. 1591–1598, 2017.

[8] A. Vaswani et al., "Attention Is All You Need," Adv. Neural Inf. Process. Syst., vol. 2017-Decem, pp. 5999–6009, Jun. 2017.

[9] R. R. Selvaraju *et al.*, "Grad-CAM: Visual Explanations from Deep Networks via Gradient-Based Localization," *Int. J. Comput. Vis.*, vol. 128, no. 2, pp. 336–359, Feb. 2020.

[10] F. Vaquerizo-Villar *et al.*, "An explainable deep-learning model to stage sleep states in children and propose novel EEG-related patterns in sleep apnea," *Comput. Biol. Med.*, vol. 2, no. 2, p. 107419, Aug. 2023.

[11] C. Guilleminault *et al.*, "Cyclical variation of the heart rate in sleep apnoea syndrome. Mechanisms, and Usefulness of 24 h Electrocardiography as a Screening Technique," Lancet, vol. 323, no. 8369, pp. 126–131, Jan. 1984.

[12] C. L. Marcus, "Sleep-disordered Breathing in Children," Am. J. Respir. Crit. Care Med., vol. 164, no. 1, pp. 16–30, Jul. 2001.



Figure 1: Confusion matrices for the 4 OSA severities in the test sets.
1: No OSA (AHI<1 e/h); 2: mild OSA (1≤AHI<5 e/h);</li>
3: moderate OSA (5≤AHI<10 e/h); 4: severe OSA (AHI≥10 e/h).</li>



Figure 2: GradCAM visualization of relevant ECG regions from nocturnal recordings. ↑ HR and ↓ HR: increase and decrease in heart rate.