

cgaroufis@mail.ntua.gr, {filby,neftymiou}@central.ntua.gr, {nzlat, maragos}@cs.ntua.gr, smyrnis@med.uoa.gr

Introduction – Problem Definition



What is a relapse?

- Deterioration in the condition of mental patients.
- Signs of relapses appear in various modalities, including **speech**.
 - Bipolar Disorder: longer pauses between utterances, increased formant frequencies.
 - Schizophrenia: lower speech rate, decreased formant frequencies.
- Goal: Being able to **detect** and **predict** the appearance of relapses from spontaneous speech in patients in the psychotic spectrum.
 - Validation of subjective clinician evaluations.
 - Ability to intervene by predicting the appearance of relapses.
- Majority of studies in the literature tackle the problem using supervised learning approaches, either feature-based [1] or using deep learning [2].

[1] J. Gideon, E. M. Provost, and M. McInnis, "Mood State Prediction from Speech of Varying Acoustic Quality for Individuals with Bipolar Disorder," in Proc. ICASSP 2016

[2] L. He and C. Cao, "Automated Depression Analysis using Convolutional Neural Networks from Speech," Journal of Biomedical Informatics, vol. 83, pp. 103–111, 2018.

Motivation - Contribution



Motivation:

- Appearance of relapses in patients is scarce -> opting for an **unsupervised approach** in an anomaly detection framework, since models can be trained only using data from stable time periods.
- Previous work used a deterministic Convolutional Autoencoder [3] for personalized relapse detection and prediction.
- How can we scale this approach at a **universal** (patient-independent) setting?

Contribution:

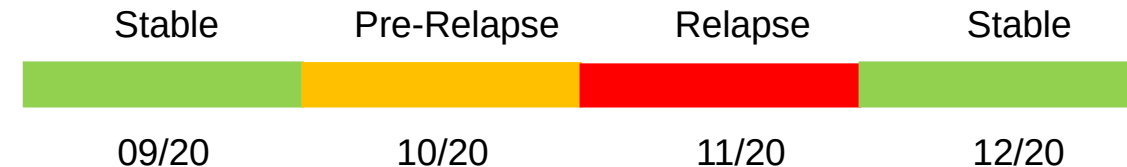
- Development of a **Convolutional Variational Autoencoder** (CVAE) on data collected from patient – clinician interviews.
- Training at spectrogram level, results aggregated in a per-session basis.
- Personalized models: **Comparable** performance between CVAEs and CAEs.
- Universal models: CVAEs **significantly outperform** CAEs, reach the performance of personalized models in conjunction with **personalized normalization** and norm pooling for temporal aggregation.

[3] C. Garoufis, A. Zlatintsi, P. Filintisi, N. Efthymiou, E. Kalisperakis, T. Karantinos, V. Garyfalli, L. Mantonakis, N. Smyrnis, and P. Maragos, “An Unsupervised Learning Approach for Detecting Relapses from Spontaneous Speech in Patients with Psychosis,” in Proc. BHI 2021

Data Collection



- Study participants: 24 patients in the bipolar or psychotic spectrum.
- Annotations of the condition of the patients as stable or relapsing by the expert clinicians, based on:
 - Monthly in-person clinical assessments between the patients and experienced clinicians, through which psychopathological scales are estimated.
 - Weekly unstructured (duration: 5-10 min) interviews conducted between patients and clinicians via a dedicated tablet app and then stored in a cloud server.
 - Communication between the clinicians and the patient's environment.
- In this work, we will use the **short unstructured interviews** between patients and clinicians.
- Further data categorization as:
 - **Clean**: Patient condition is annotated as stable.
 - **Relapse**: A relapse has been detected by the clinicians.
 - **Pre-relapse**: Interviews conducted up to 30 days prior to the appearance of a relapse.
- Both pre-relapse and relapse data are considered as **anomalous**.



Data Preprocessing



Collected Dataset Statistics:

- Total amount of data used: **375 interviews**, from **13 patients**.
- 8 of the patients experienced a relapse during the course of the study.
- The rest were selected on the basis of amount of available speech data.

Patient Speech Isolation:

- Audio extracted from interviews, and downsampled to 16 kHz.
- Speech excerpts corresponding to the patients isolated using kaldi.
- Final utterance statistics: **12107 utterances/30509 sec**.

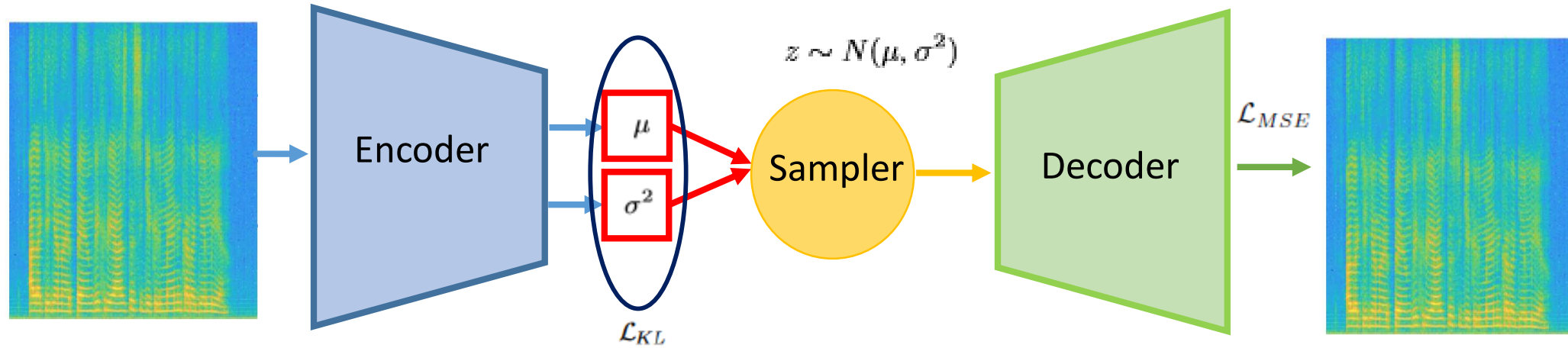
Feature Extraction:

- Computed **mel-spectrograms** for each speech utterance.
- Parameters: 512-sample window, 256-sample hop length, 128 mel bands.
- Spectrograms cut at slices of 64 frames (ca 1 sec.) -> 128x64 input representation, then standardized and log-scaled.

Patient demographics, as well as statistics on the amount of recorded and analyzed speech utterances.

Demographics	
Male/Female	8/5
Age (years)	27.5 ± 6.7
Education (years)	13.5 ± 1.9
Illness dur. (years)	7.9 ± 7.6
Recorded Data	
Num. of Interviews (total)	375
Num. of Interviews (mean $\pm$ std)	28.8 ± 8.7
Diarized speech duration (in sec)	30509
Diarized speech duration (in sec, mean $\pm$ std)	2347 ± 1550
Num. of Utterances (total)	12107
Num. of Utterances (mean $\pm$ std)	931 ± 527
Num. of Utterances (clean, mean $\pm$ std)	754 ± 425
Num. of Utterances (pre-relapse, mean $\pm$ std)	119 ± 126
Num. of Utterances (relapse, mean $\pm$ std)	169 ± 162

Methodology: Variational Autoencoders



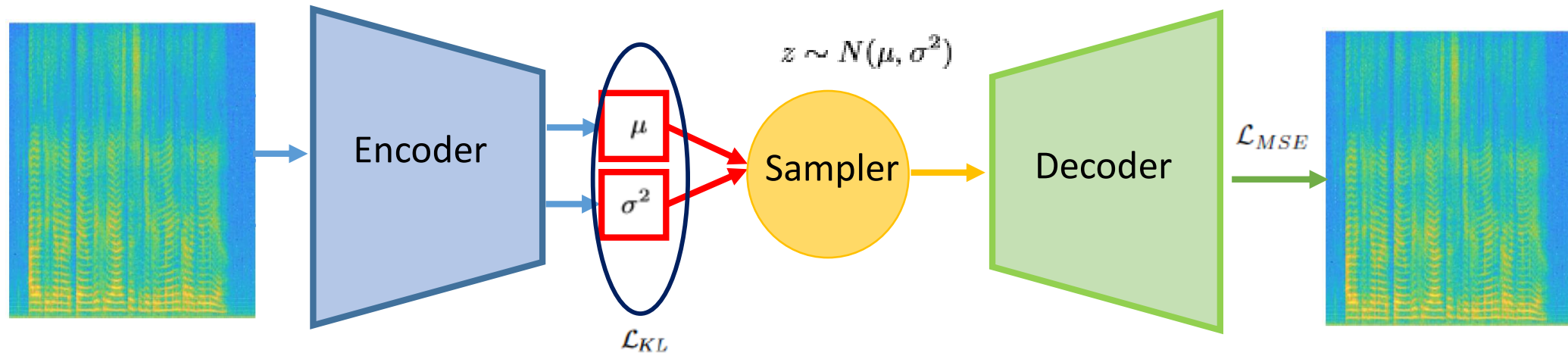
- Probabilistic variant of classical autoencoders, first developed in [4]
- Encoder (inference model): **Encodes** its input into a low-dimensional latent representation, assumed to follow an isotropic Gaussian distribution.
- Decoder (generative model): Attempts to **reconstruct** the input from a sample drawn from the learned distribution.
- Used in a variety of audio-related tasks, such as speech enhancement [5] and speech representation learning [6]

[4] D. P. Kingma and M. Welling, "Auto-Encoding Variational Bayes," arXiv preprint arXiv:1312.6114, 2013.

[5] H. Fang, G. Carbajal, S. Wermter, and T. Gerkmann, "Variational Autoencoder for Speech Enhancement with a Noise-Aware Encoder", in Proc. ICASSP 2021

[6] J. Chorowski, R. Weiss, S. Bengio, and A. van den Oord, "Unsupervised Speech Representation Learning using WaveNet Autoencoders," IEEE/ACM Trans. on Audio, Speech, and Language Process., vol. 27, no. 12, pp. 2041–2053, 2019.

Methodology: Variational Autoencoders



- **Encoder:** 3 convolutional blocks, alternating 2D-convolutional layers and 2D max pooling layers, **increasing** number of filters + pair of parallel layers to estimate μ and σ^2 .
- **Decoder:** 4 convolutional blocks, alternating 2D-upsampling layers and 2D convolutional layers, **decreasing** number of filters.
- Activations: LeakyReLU (no activation at the output layer)
- Loss functions:
 - MSE loss at the **output** of the network, between the true and estimated spectrograms.
 - KL loss at the **distribution** of the encoded embeddings, between the learned distribution and the spherical isotropic Gaussian, $N(0, I)$.

Net. Block	N_{filt}	(k_x, k_y)	(p_x, p_y)	(u_x, u_y)
Conv_DS1	N	(5,5)	(2,2)	-
Conv_DS2	2N	(5,5)	(4,2)	-
Conv_DS3	4N	(5,5)	(4,4)	-
Conv_DS4(μ)	8N	(4,4)	(4,4)	-
Conv_DS4(σ)	8N	(4,4)	(4,4)	-
Sample	-	-	-	-
Conv_US1	4N	(5,5)	-	(4,4)
Conv_US2	2N	(5,5)	-	(4,4)
Conv_US3	N	(5,5)	-	(4,2)
Conv_US4	1	(5,5)	-	(2,2)

Experimental Setup



- Experiments for both **personalized** (separate model for each patient) and **universal** (one model for all patients) cases.
- Baseline: The CAE model presented in [3].
- $N = 32$ filters at the outer convolutional layer, for both CAE and CVAE models.

Training Details:

- 5-fold cross-validation, data from the same session are assigned to the same fold.
 - Training: Only data from time periods where the patient condition was **stable**
 - Testing: Mixture of data from **stable** and **anomalous** (pre-relapsing or relapsing) time periods.
- Adam ($\text{lr}=0.0003$), batch size of 8.
- 200 epochs maximum, early stopping applied at 10 epochs.
- Loss weights: $W_{MSE} = 1$, and $W_{KL} = 0.01$
- Evaluation Metric: Mean **ROC-AUC score** over all sessions.

Experimental Setup – Ablation Studies



- Probe point for the anomaly scores:
 - CAE: Output reconstruction MSE
 - CVAE: Output reconstruction MSE and input embedding KL divergence.
- Temporal aggregation function of the per-session anomaly scores:
 - Average pooling (AP): $S = \frac{1}{N} \sum_{i=1}^N s_i$
 - Max pooling (MP): $S = \max(s_i)$
 - Norm pooling (NP): $S = \frac{1}{N} (\sum_{i=1}^N |s_i|^p)^{1/p}$, $p = 10$.
- Normalization scheme (universal models):
 - Global normalization, i.e. a **shared** normalization transform for all patients.
 - Per-patient normalization, i.e a **separate** normalization transform for each patient.

Results: Personalized Models



Average per-patient ROC-AUC score for the discrimination between sessions that correspond to stable, or anomalous, condition, for both CVAE and CAE personalized models.

Pooling Function	CAE [24]	CVAE	
		MSE	KL
AP	0.668 ± 0.035	0.673 ± 0.055	0.653 ± 0.052
MP	0.608 ± 0.060	0.617 ± 0.051	0.659 ± 0.045
NP	0.627 ± 0.058	0.640 ± 0.049	0.678 ± 0.049

- The performance of the proposed CVAE is **comparable** to that of the deterministic CAE in the personalized case.
- No statistically significant difference ($p > 0.05$) between models.
- Average pooling performs the best when using the reconstruction MSE as the anomaly score for both CAE and CVAE models.
- Norm pooling gives the best results when using the KL divergence as the anomaly score.

Per-patient ROC-AUC scores for the discrimination between sessions that correspond to stable, or anomalous, condition, for both CVAE and CAE personalized models.

Patient ID	CAE [24]	CVAE	
		MSE	KL
#1	0.546 ± 0.069	0.547 ± 0.081	0.523 ± 0.134
#2	0.448 ± 0.093	0.418 ± 0.182	0.388 ± 0.120
#3	0.711 ± 0.119	0.700 ± 0.159	0.656 ± 0.187
#4	0.676 ± 0.066	0.660 ± 0.034	0.768 ± 0.066
#5	0.781 ± 0.053	0.800 ± 0.095	0.774 ± 0.040
#6	0.512 ± 0.067	0.520 ± 0.173	0.696 ± 0.083
#7	0.877 ± 0.076	0.940 ± 0.074	0.720 ± 0.186
#8	0.800 ± 0.187	0.850 ± 0.292	0.900 ± 0.200
Average	0.668 ± 0.035	0.673 ± 0.055	0.678 ± 0.049

- Per patient results: ROC-AUC score **above 0.75** for 4/8 patients, **above 0.65** for 6/8

Results: Universal Models



- The CVAE model **outperforms** by a large margin the baseline CAE, especially when obtaining the anomaly score from the KL divergence.
- Application of **per-subject normalization** leads to performance equivalent to the one achieved by the personalized models.
- Statistically significant improvement ($p < 0.05$) over the baseline when using personalized normalization and the KL divergence as anomaly score.
- **Norm pooling** appears to perform the best as a temporal aggregation function.
- CVAEs perform better at a patient-independent setting -> speaker-invariant representations? [6]

Average ROC-AUC score for the discrimination between sessions that correspond to stable, or anomalous, condition, for both CVAE and CAE universal models.

Pers. Norm	Pool. Func.	CAE [24]	CVAE	
			MSE	KL
X	AP	0.504 ± 0.032	0.502 ± 0.016	0.532 ± 0.023
X	MP	0.542 ± 0.024	0.551 ± 0.023	0.592 ± 0.031
X	NP	0.531 ± 0.034	0.527 ± 0.024	0.581 ± 0.036
✓	AP	0.552 ± 0.036	0.585 ± 0.021	0.646 ± 0.035
✓	MP	0.541 ± 0.027	0.622 ± 0.034	0.685 ± 0.040
✓	NP	0.542 ± 0.028	0.618 ± 0.030	0.698 ± 0.042

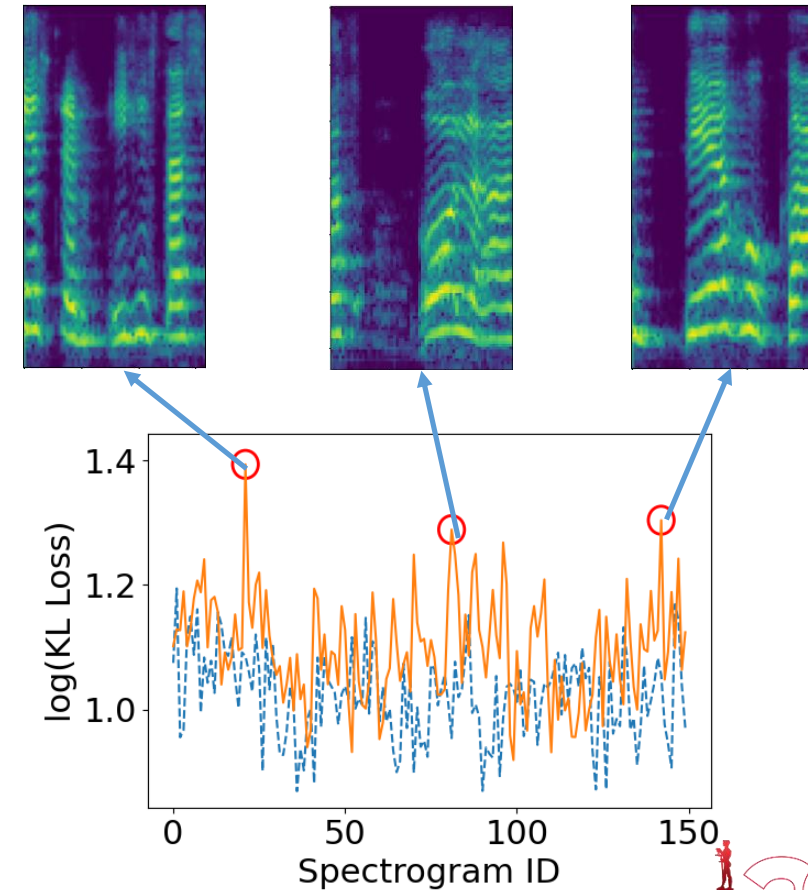
Results: Qualitative Analysis



- KL scores for two interview sessions over time:
 - Stable condition (dashed blue)
 - Relapsing condition (orange).

Observations:

- Not significantly higher KL scores during the relapsing session
- Appearance however of **a few peaks** (in red circles)
- Aural inspection of the respective segments -> **abrupt in-utterance disruptions** of the patient's speech flow.



Conclusions & Future Work



Explored the potential of Convolutional Variational Autoencoders (CVAEs) in speech-based relapse detection prediction in psychotic patients.

- Personalized case: **Comparable** performance to a CAE baseline.
- Universal case: **Significant improvement** over CAEs, in conjunction with a **personalized normalization** scheme.

What's next?

- Utilization of **multimodal information**, such as text transcripts, or data collected from smartwatches.
- Taking advantage of **longer-term dependencies** in interviews:
 - During the same utterance.
 - In successive utterances.



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