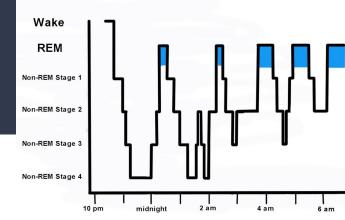
# Learning Sleep Stages from Radio Signals

A Conditional Adversarial Architecture

Mingmin Zhao, Shichao Yue, Dina Katabi, Tommi S. Jaakkola, Matt T. Bianchi

Presenters: Dennis Ai, Michael Chen, Hansohl Kim, Sam Kim, Jimmy Wu

## Stages of Sleep



NREM Stage 1	NREM Stage 2	NREM Stage 3	REM
Lightly asleep	Slowing heart rate, decreasing body temperature	Larger, slower brain waves	Deep and powerful dreams
Easy to wake up, sometimes sensations of falling	Harder to wake up, larger brain waves	Large, slow brain waves  Can sleep through most  disturbances	Eyes move rapidly, sleepwalking, increase in heart rate

## Sleep Lab

Requires sensors on scalp, chest, nose

## Sleep at Home

Box with RF sensor





## Current State-of-the-art Methods

#### Polysomnography (PSG)

- intrusive i.e. requires patient to wear sensors, electrodes, monitors, probes, etc
- expensive
- uncomfortable

#### Radio Technology

- capture physiological signals without body contact
- transmit low power radio signal
- less intrusive

## Key Challenge

Radio frequency reflections are highly dependent on the **measurement conditions** and the **individual** 



#### Multi-Source Domain Adaptation

#### domain = measurement condition + individual

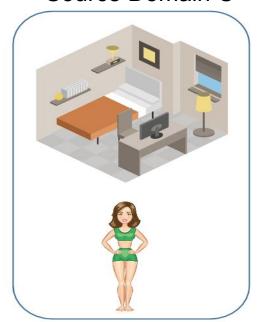
Source Domain A



Source Domain B

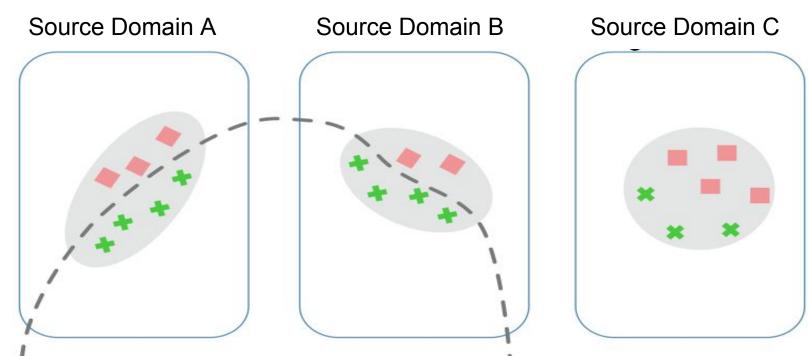


Source Domain C

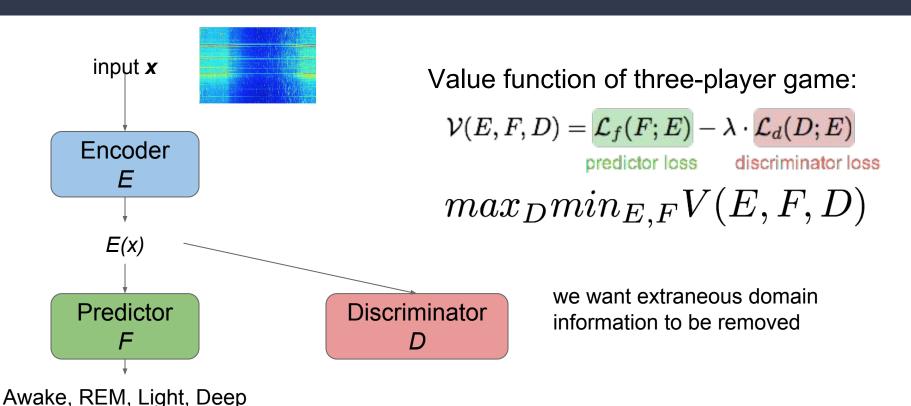


#### Multi-Source Domain Adaptation

domain = measurement condition + individual

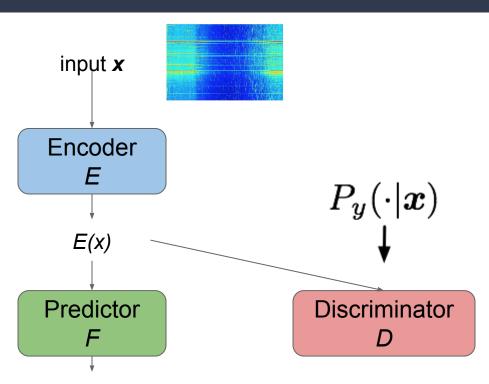


#### Conditional Adversary Architecture



## Predictor Encoder Architecture Discriminator LSTM LSTM LSTM

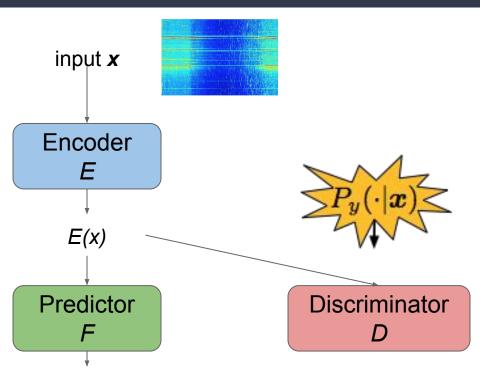
#### Conditional Adversary



Add a posterior of the label distribution (don't want this information removed)

Awake, REM, Light, Deep

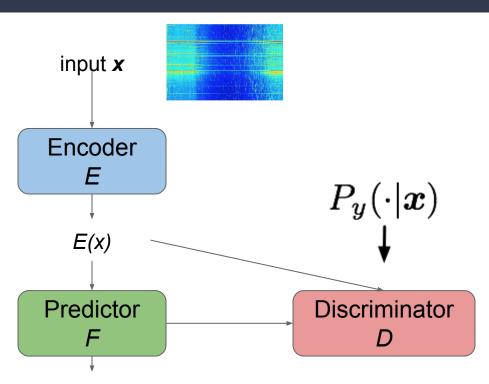
#### Conditional Adversary



Usually, posterior is not available during training!

Awake, REM, Light, Deep

#### Conditional Adversary



Solution: Condition the discriminator on predicted label distribution

Awake, REM, Light, Deep

#### Dataset

- 25 different bedrooms, 100 nights
- ~90K 30-second pairs of RF spectrograms and corresponding sleep stages
- Ground-truth labels: FDA-approved EEG-based sleep profiler of sleep stages

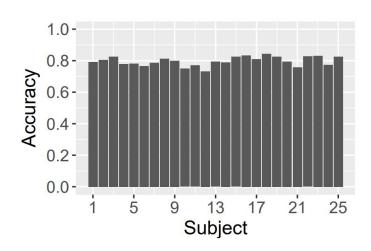






#### Testing

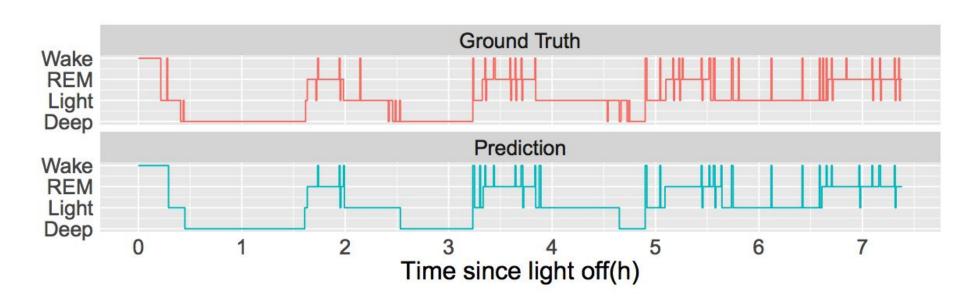
- Encoder: ResNet CNN + LSTM
- Predictor and Discriminator: 2-layer FC
- Each subject evaluated after training on all others
- Metrics: Accuracy + Cohen's Kappa
- Consistently high across subjects
- Standard Deviation: 2.9%



#### Results

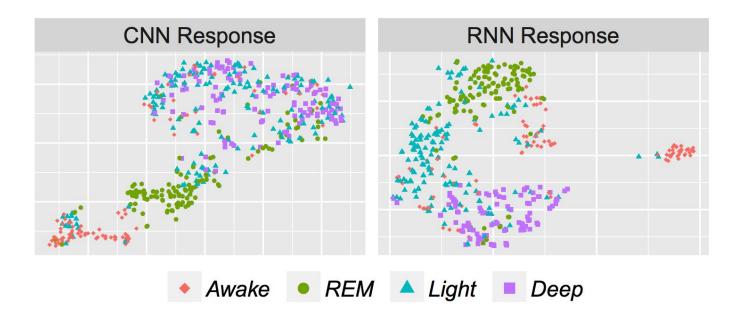
Signal Source		Accuracy (%)	Cohen's Kappa ( <i>k</i> )	Comfort
EEG		83	.76	Low
Cardiorespiratory		71	0.56	Medium
Actigraphy		Low	-	High
RF	State-of-the- art	64	0.49	High
	Conditional Adversary	79.8	0.70	High

## Ground Truth vs. Predicted Sleep Staging

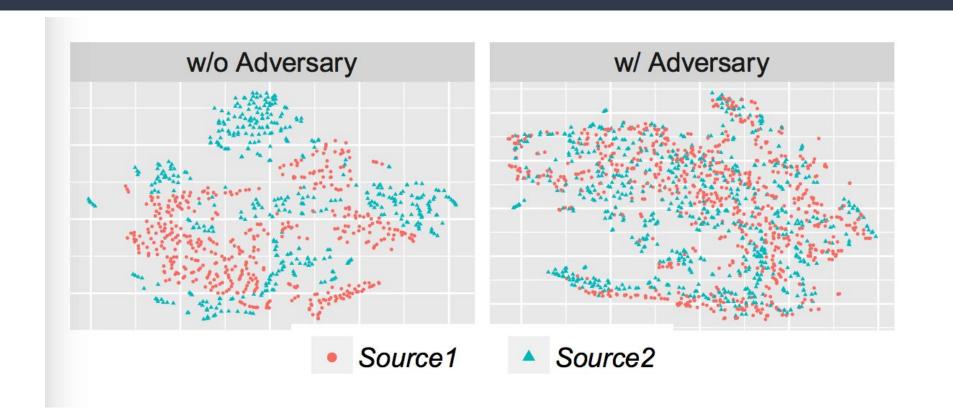


#### t-SNE Visualizations

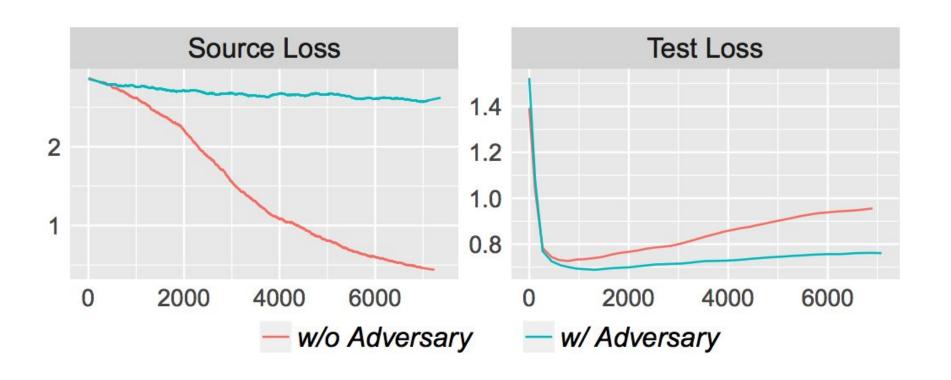
CNN Response - successful at separating Wake, REM from Light and Deep Sleep RNN Response - successful at separating Light and Deep Sleep



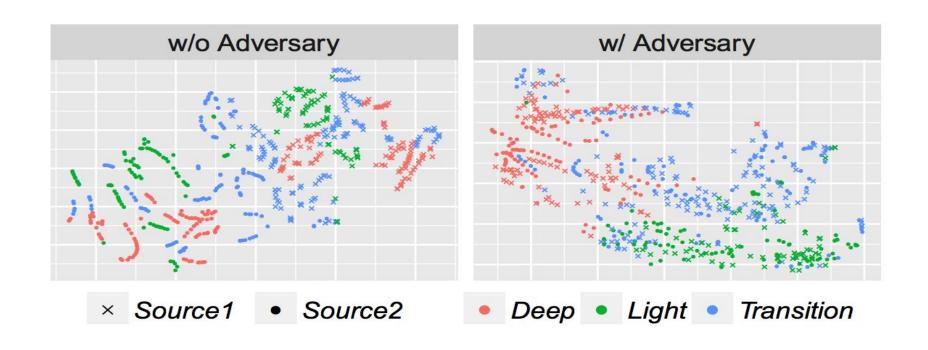
#### Role of Adversarial Discriminator



#### Role of Adversarial Discriminator



## Conditioning on the Posterior Distribution



#### Criticisms

RF signals are inherently noisy, which a network could learn to remove with lots of data, which the authors didn't have. Could have leveraged more signal processing techniques.

Encoder could just learn to output the posterior distribution.

Could include more information about analyzing encoding versus labels/environment.

The paper does not specify the details and hyperparameters of their architecture and thus does not allow its results to easily be replicated.

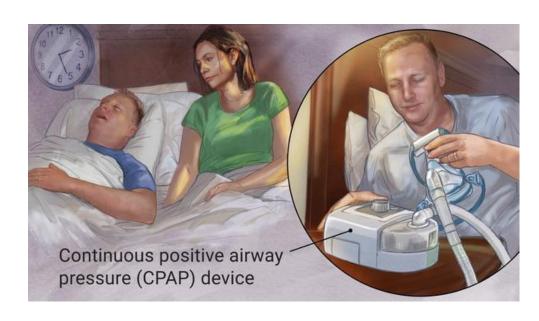
#### Future Work

In the confusion matrix, the true positive rate for the awake state is much lower than for other sleep stages (63% vs  $\sim$ 80%). Why is this the case?

What changes could be made to the encoder architecture to improve performance, better model transitions between sleep stages, or provide more fine-grained categorization of sleep stages?

Can this model truly be extended to non-healthy individuals, or individuals across a large variety of demographics (e.g. age)?

Can this solution be applied to diagnosis sleep apnea or other sleep disorders, which affects over 50 million people in the United States? This would mean measuring nasal or oral airflow, respiratory effort, and oximetry.



## Questions