

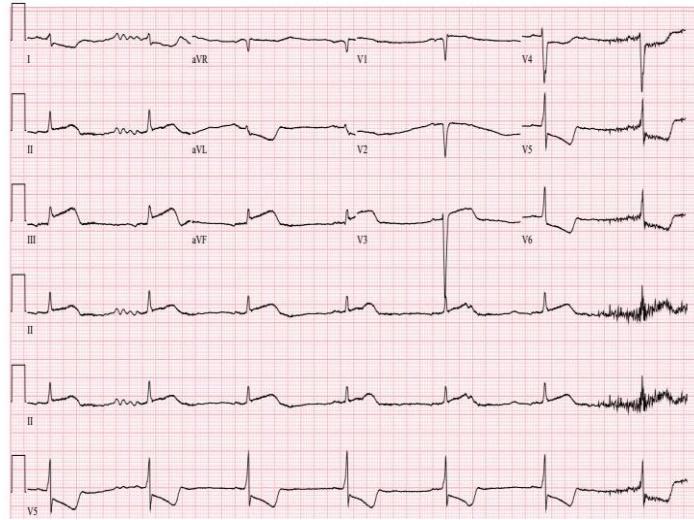


UNLOCKING PATTERNS IN EXPOSURE DATA WITH SHAPELETS

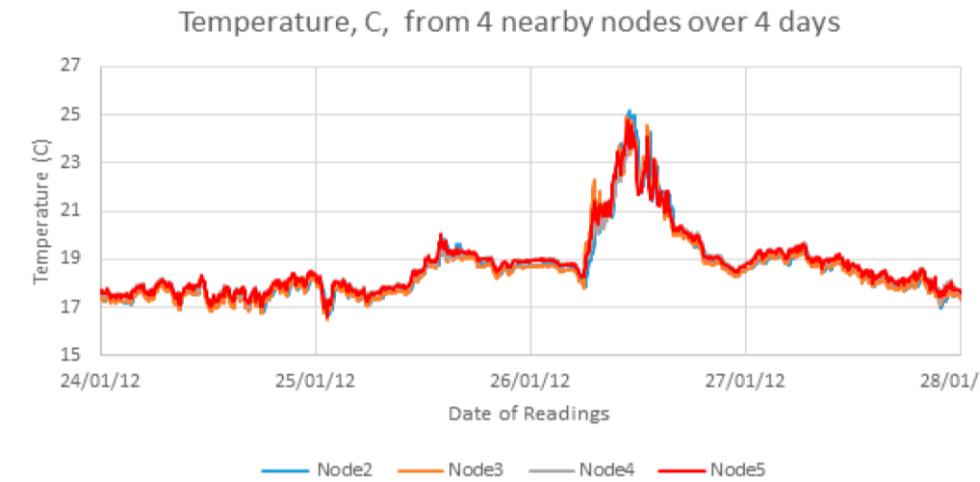
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BACKGROUND

- Time series data appears in many domains



Health Monitoring



Sensor Network



Financial World

BACKGROUND

- Temporal Exposure: Duration & frequency
- Spatial Exposure: Substance & place



+



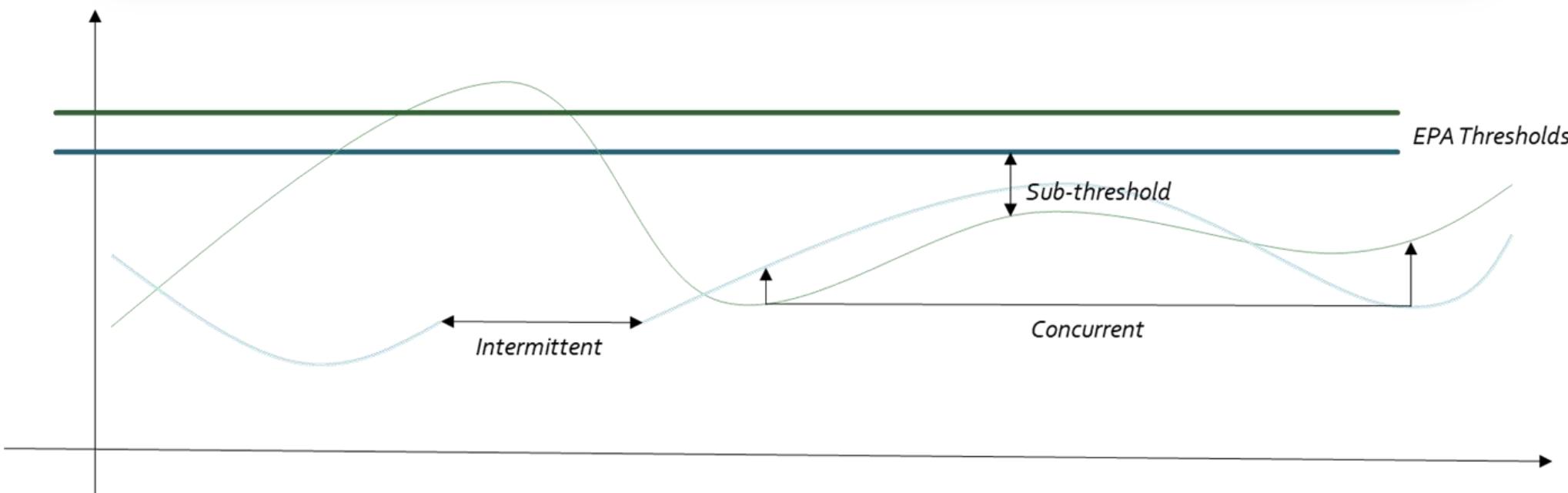
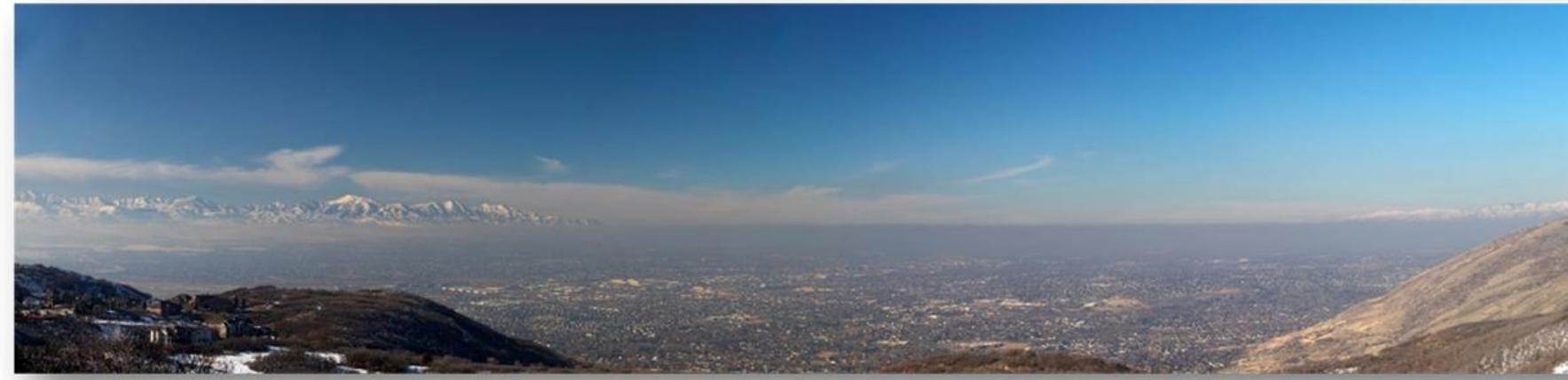
Temporal Exposure

Spatial Exposure

Health Outcome

OBJECTIVES

- Intermittent, sub-threshold effects



OBJECTIVES

ORIGINAL RESEARCH

Intermittent Exposure to Cigarette Smoke Increases Lung Tumors and the Severity of Emphysema More than Continuous Exposure

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Abstract

Lung cancer and chronic obstructive pulmonary disease are leading causes of morbidity and mortality worldwide, and cigarette smoking is a main risk factor for both. The presence of emphysema, an irreversible lung disease, further raises the risk of lung cancer in patients with chronic obstructive pulmonary disease. The mechanisms involved in smoke-induced tumorigenesis and emphysema are not fully understood, attributable to a lack of appropriate animal models. Here, we optimized a model of cigarette smoke (CS)-induced lung cancer and emphysema in A/J mice treated with 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone, a potent carcinogen. We investigated whether variations in CS exposure patterns with the same total amount and duration of exposure affect tumorigenesis and/or development of emphysema.

Continuous CS exposure for 3 months significantly suppressed 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone-induced development of adenomas and adenocarcinomas; however, emphysema independently developed during this period. Surprisingly, intermittent CS exposure increased the severity of emphysema and resulted in a higher incidence of adenocarcinomas. Furthermore, intermittent CS exposure elicited a marked increase in M2-polarized macrophages within and near the developed tumors. By employing a CS exposure protocol with repeated cycles of cessation and relapse, we provide evidence that intermittent CS exposure enhances tumorigenesis and emphysema progression more than that of continuous CS exposure.

Keywords: lung cancer; emphysema; cigarette smoke; M2 macrophage



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Predicting Lung Cancer Incidence from Air Pollution Exposures Using Shapelet-based Time Series Analysis

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Abstract

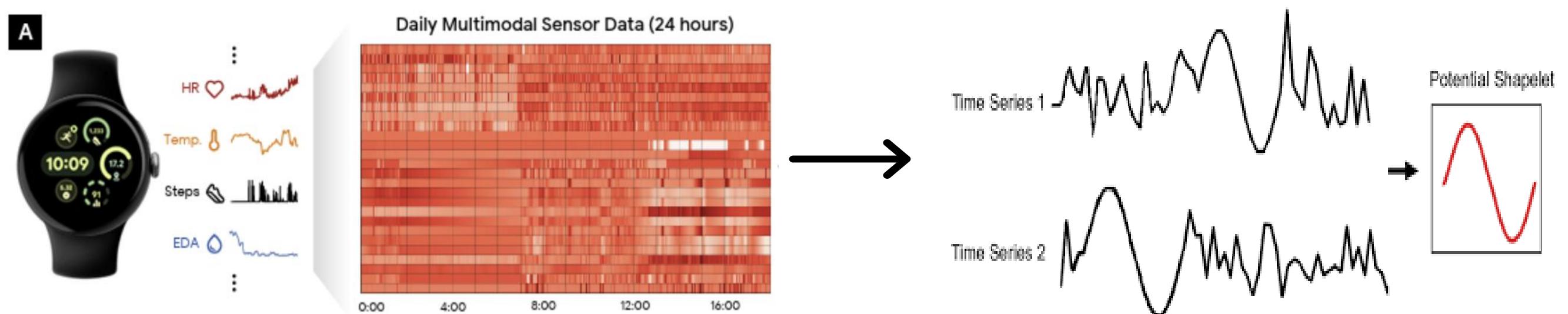
In this paper we investigated whether the geographical variation of lung cancer incidence can be predicted through examining the spatiotemporal trend of particulate matter air pollution levels. Regional trends of air pollution levels were analyzed by a novel shapelet-based time series analysis technique. First, we identified U.S. counties with reportedly high and low lung cancer incidence between 2008 and 2012 via the State Cancer Profiles provided by the National Cancer Institute. Then, we collected particulate matter exposure levels (PM_{2.5} and PM₁₀) of the counties for the previous decade (1998–2007) via the AirData dataset provided by the Environmental Protection Agency. Using shapelet-based time series pattern mining, regional environmental exposure profiles were examined to identify frequently occurring sequential exposure patterns. Finally, a binary classifier was designed to predict whether a U.S. region is expected to experience high lung cancer incidence based on the region's PM_{2.5} and PM₁₀ exposure the decade prior. The study confirmed the association between prolonged PM exposure and lung cancer risk. In addition, the study findings suggest that not only cumulative exposure levels but also the temporal variability of PM exposure influence lung cancer risk.

Ref: Naofumi Kameyama, Shotaro Chubachi, Hegab AE, et al. Intermittent Exposure to Cigarette Smoke Increases Lung Tumors and the Severity of Emphysema More than Continuous Exposure. *American Journal of Respiratory Cell and Molecular Biology.* 2018;59(2):179-188. doi:https://doi.org/10.1165/rcmb.2017-0375oc

Ref: Yoon HJ, Xu S, Tourassi G. Predicting Lung Cancer Incidence from Air Pollution Exposures Using Shapelet-based Time Series Analysis. *IEEE EMBS Int Conf Biomed Health Inform.* 2016;2016:565-568. doi:10.1109/BHI.2016.7455960

SHAPELETS

- Discriminative sub-sequences of time series data that serve as a feature for classification or regression task



RESULTS

$600 \times 20 \times 2 \times 100 \approx 2,400,000$

(Counties) (pollutants) (Window size) (Number of shapelets / location)

$2,400,000 \times 20 \approx 48,000,000$

(Years)

Shapelet Library

Data Type: Daily

Pollutant: SO2

Period: From 01/01/2020 to 10/31/2022

Geographic Area: Salt Lake City

Shapelet Length: 7 Days

Apply Filters

Showing 1-20 of 1200 shapelets

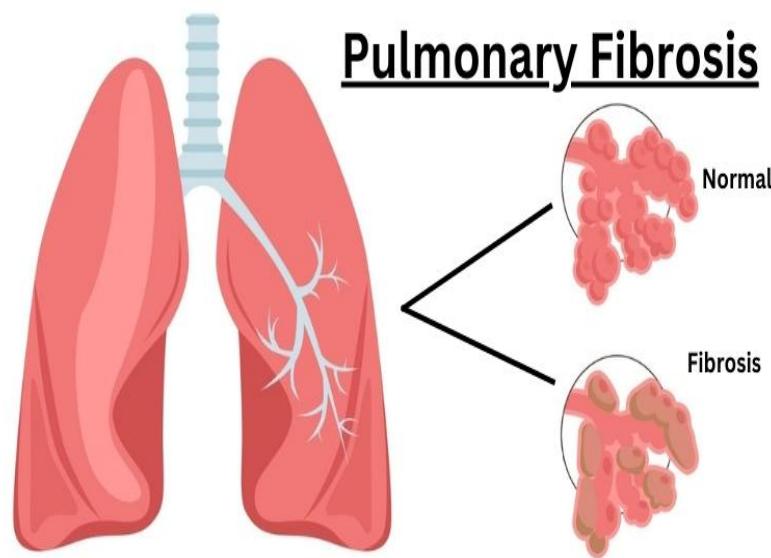
Shapelet ID	Location	Period	Year	Quality	Mean	Duration
Utah_Salt_Lake_2005_shapelets.pkl_2020_000	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.047429	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_001	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.049078	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_002	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.048256	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_003	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.064530	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_004	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.057244	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_005	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.034431	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_006	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.030494	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_007	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.033309	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_008	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.132224	0.0000	7d
Utah_Salt_Lake_2005_shapelets.pkl_2020_009	Utah_Salt_Lake_2005_shapelets.pkl	2020-01-01 to 2020-01-07	2020	0.168759	0.0000	7d



<https://ehie-shapelets.ctsi.utah.edu/>

NEXT STEPS

- Linking exposure motifs to health outcomes



ACKNOWLEDGEMENTS



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