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"TimBre" Pilot Study Conducted Using Multi-country Training and Validation Data for Screening of Pulmonary Tuberculosis Using Cough (Acoustic Sounds), Clinical & Demographic Inputs

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ABSTRACT

TimBre from Docturnal offers multidirectional screening of Lung Ailments – Pulmonary Tuberculosis, Pneumonia, Covid19 & COPD. Detailed studies of TimBre in the past used third party Microphone Array that focused on a XY arrangement that provided high fidelity cough sounds with an average length of >5 seconds and real-time demographic data such as Height, Weight, BMI [1]. In the current study, cough sounds were harvested from 7 different countries (India, Vietnam, Philippines, Uganda, Tanzania, Madagascar, SA) using Mobile Phones from different manufacturers & recorded solicited coughs in a clinic for a duration of 0.5 seconds. A plethora of demographic and clinical variables were provided of which a subset was used by TimBre algorithm. Most importantly, the .WAV files were recorded in a single channel at a sampling rate of 44.1kHz & 16 bits. The study details two approaches wherein the first method was to concatenate all the 0.5 second Supers as-is in both training and scoring sets without any concatenation. The first approach on the TEST set yielded a sensitivity and specificity (table-1 with CI values at 0.05) of 75.41% and 68.30% respectively with an AUC of 0.78 as reported by UCSF R2D2 team. Both the approaches used a combination of Clinical, Demographic and Spectral Variables. Some additional variables included were derived (BMI) & excluded (Spectral) based on the feature importance scores.

The ML model performed better in the second approach and we anticipate it to improve further once an additional 714,922 .WAV files harvested as Longitudinal coughs shall be appended to the training set as a part of a subsequent pilot study.

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Abbreviations

UCSF R2D2	University of San Francisco Rapid Research in Diagnostics Development
COPD	Chronic Obstructive Pulmonary Disorder
AUC / ROC	Area Under Curve / Receiver Operating Characteristic Curve
.WAV	Wave Audio File Format
ТВ	Tuberculosis
EMR	Electronic Medical Record
XY	An arrangement of Microphone Arrays on a Microphone
BMI	Body Mass Index
HIV	Human Immunodeficiency Viruses
PDP	Partial Dependence Plot
MRMR	Maximum Relevance Minimum Redundancy
RUS	Random Under Sampling

FFT	Fast Fourier Transformation
ML / DL	Machine Learning / Deep Learning
CNN	Convolutional Neural Network
MFCC	Mel Frequency Cepstral Coefficient
Hz	Hertz
AWS	Amazon Web Services
XAI	Explainable Artificial Intelligence
AI	Artificial Intelligence
HR	Heart Rate
CI	Confidence Interval

Introduction

Tuberculosis (TB), a communicable disease caused by Mycobacterium tuberculosis, is a major cause of ill health and one of the leading causes of death worldwide. Until the COVID-19 pandemic, TB was the leading cause of death from a single infectious agent, ranking even above HIV/AIDS.

In 2020, an estimated 9.9 million people fell ill with TB and 1.3 million died of TB worldwide. However, approximately 40% of people with TB were not diagnosed or reported to public health authorities because of challenges in accessing health facilities or failure to be tested or treated when they do. The development of low-cost, non-invasive digital screening tools may improve some of the gaps in diagnosis.

As cough is a common symptom of TB, it has the potential to be used as a biomarker for diagnosis of disease. Several previous studies have demonstrated the potential for cough sounds to be used to screen for TB, though these were typically done in small samples or limited settings [1-3]. Further development and evaluation are critical to move the field forward.

Here we leverage data collected from people who presented to clinics across 7 countries with new or worsening cough for at least 2 weeks. Elicited coughs were recorded using the Hyfe Research app. Individuals were then comprehensively evaluated for TB with sputum-based molecular (Xpert MTB/RIF Ultra) and culture (MGIT or Lowenstein-Jensen) testing.

Materials and Methods Per StudyId Model

In this model, a total of 10787 WAV files in the training set were mapped against 1269 StudyIDs in the first approach resulting in 1231 concatenated .WAV files implying that 38 StudyId's were included in the training set using clinical and demographic variables only (without .WAV). The clinical variables added additional clinical information that were not used in our earlier studies for lack of accurate information (EMR typically) to avoid subjectivity [1]. However, the below model included/added PriorTB and Hemoptysis given its critical nature to contribute to the model in determining a drug resistant TB or hemoptysis that is an indicator of a bigger problem.

Clinical & Demographic variables used

'Gender', 'Age', 'Height', 'Weight', 'BMI', 'PriorTB', 'Hemoptysis'

Per	Study	ID	model	(Table	1)
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	Total StudyIds (Meta Data)	StudyIds with cough (.WAV)	.WAV files before concatenation	.WAV files post concatenation	Sensitivity	Specificity	AUC
Train	1269	1245	10787	1231	72%	67%	0.73
TEST (Prediction)	1276	1248	10008	1248	68.6% (65.48%, 71.61%)	71.7% (66.31%, 76.78%)	0.75 (0.7283, 0.7890)

ROC of the Model



Per Cough Model

In this model, 10,008 .WAV files were predicted using a model created out of 10787 .WAV files. The clinical variables added additional clinical information that were not used in our earlier studies for lack of accurate information (EMR typically) to avoid subjectivity [1]. However, the below model included/added PriorTB, Hemoptysis, HIVstatus, Nightsweats, Heartrate, Fever & Temperature given its critical nature to contribute to the model in determining the usage of classic variables that are used in current TB programs as a part of screening questionnaire despite the fact that many researchers from the TB Coda Dream Challenge indicated Temperature variable as an "awash" [4]. In any healthcare setting, adding these would improve the screening accuracy on top of differential diagnosis should there be a mechanism to do so via an EMR or any other such repository

Clinical & Demographic Variables used

'Gender', 'Age', 'Height', 'Weight', 'BMI', 'HIVstatus', 'PriorTB', 'Hemoptysis', 'Fever', 'Nightsweats', 'Heartrate', 'Temperature'

Per Cough model (Table 2)

	Total StudyIds (Meta Data)	StudyIds with cough (.WAV)	.WAV files before concatenation	.WAV files post concatenation	Sensitivity	Specificity	AUC
TRAINING	1269	1245	10383	NA	81%	81%	0.90
Score (Prediction)	1191	1248	10008	NA	75.41% (74.40%, 76.40%)	68.30% (66.48%, 70.07%)	0.78 (0.7707, 0.7918)

ROC of the Model



Interpretability & Explainability

- 1. Both the supervised learning models used RUSBoosted Ensemble technique given the fact that the TB (Positive) and TB (Negative) were to the tune of 30% and 70% respectively in the provisioned datasets representing an imbalanced class set. The feature selection used Kruskall-Wallis Algorithm given its distribution and outlier agnostic nature in comparison with ChiSquare, Annova & MRMR. The models were built out of the box using MATLAB R2022a Classification Learner from MathWorks [6].
- 2. Explainability & Interpretability of the per-cough model that used 10-fold Cross Validation revealed BMI, Heart Rate, Night Sweats and Spectral Centroid to be the differentiators in segregating classes as seen below (Box1 PDP) [10]. We presume that additional training data shall reveal more such patterns & we shall share the entire list of contributing spectral variables in Phase-2 of the study.











Results & Discussion

- The approach to use Random Under Sampling Boosted Models (Ensemble – RUS) on top of features extracted post FFT has provided a great deal of explainability and interpretability [7-8]. The fact that MFCC components were already used obviates the need for CNN models relying on Spectrograms there by retaining the explainability & interpretability [10]. Most importantly, the contrastive AI approach (repeatability) wherein one can clearly understand what variable values in the TESTING set shall alter the prediction result [9]. Example: altering the standard deviation for "spectral centroid" or a "BMI" value.
- 2. In the context of Pulmonary TB screening, the interpretability as depicted in the Box1-PDP (Partial Dependence Plot) above is a useful tool & can be interpreted as below:
- a. BMI: Any value greater than 22 starts reducing the probability score from 0.8 to 0.65 for TB Negative labels as seen from the PDP. The outliers are left intentionally & distinct values are depicted as bars on the X-Axis.
- b. Heart Rate: HR of 110 and above depicts an increase in the probability score for a TB Positive label.
- c. Night Sweats: Has a slight increase in the Positive labels as seen from the probability score and inversely, a dip is observed for the Negative labels moving from No to Yes value.
- d. Spectral Centroid: Depicts a slight dip in the probability score for Negative labels and for distinct values on the X-Axis, there is a slight increase for the Positive labels.

Note: PDPs are generated on the TRAINING set only & we are yet to receive labels for the TESTING set

- 3. The feature extraction used variables spread across different bands ranging from 0 to 5000Hz. Early data sets received during the last quarter of 2022 from UCSF team observed spurious explainability for Spectral Centroid in the 200Hz band. A constant effort has been made to analyze subsequent bands while not undermining the importance of low frequency components.
- Models used with and without demographic & clinical variables 4. on top of spectral features were evaluated and the inclusion of clinical & demographic variables always improved the accuracy. This has been the case from earlier pilots as well [1]. However, having this information under the guidance of a Physician or an EMR is of extreme importance instead of Subject providing this information that introduces some ambiguity as experienced in out earlier studies [1]. Since the study was blinded, we did not get enough information about how the data was collected & what to exclude to avoid subjectivity which otherwise shall perturb the algorithm. We presume that the clinical and demographic information was obtained from a clinical record as opposed to eliciting it from the Subjects. As an example, PriorTB has a good number of unknown/unsure values.
- 5. It is observed in the Coda TB dream challenge, participants had access to Longitudinal cough files & used a combination of CNN and other Ensemble models [4]. Once the Longitudinal cough data is made public, we would like to append the same to existing Ensemble models & also explore CNN models.

Data Acknowledgement

1. "The datasets used for the analyses described were contributed by Dr. Adithya Cattamanchi at UCSF and Dr. Simon Grandjean Lapierre at University of Montreal and were generated in collaboration with researchers at Stellenbosch University (PI Grant Theron), Walimu (PIs William Worodria and Alfred Andama); De La Salle Medical and Health Sciences Institute (PI Charles Yu), Vietnam National Tuberculosis Program (PI Nguyen Viet Nhung), Christian Medical College (PI DJ Christopher), Centre Infectiologie Charles Merieux Madagascar (PIs Mihaja Raberahona & Rivonirina Rakotoarivelo), and Ifakara Health Institute (PIs Issa Lyimo & Omar Lweno) with funding from the U.S. National Institutes of Health (U01 AI152087), The Patrick J. McGovern Foundation and Global Health Labs."

- 2. Solicited Training and Scoring data .WAV files used Hyfe.ai app https://www.hyfe.ai/ [5].
- 3. We sincerely thank Dr. Devan Jaganath from UCSF R2D2 team for analysis of the VALIDATION set results.
- 4. We sincerely thank Dr. Sophie Huddart for her patience in providing the calculated TEST results across multiple iterations based on posterior probability scores provided by TimBre algorithm.

Limitations

- 1. The study used Solicited coughs to build the training model. Additional 700,000 longitudinal cough files shall improve the accuracy & explainability.
- 2. A single channel mono WAV files could be replaced with a dual channel configuration for a more breadth of information
- 3. Mobile Phone make/models were unknown to determine uniformity across the results between training and testing files in that whether they were homogenous or mixed.

Next steps

- 1. Sage bionetworks shall open the Validation process that shall open up additional Longitudinal data for 714,922 wav files to be included in the ML/DL Model.
- 2. Explore the Maximum (Majority) rule as per UCSF recommendation for per cough model. Note that the current champion Ensemble model (RUS boosted) yielded a consistent result for each Study ID while other models had a mixed response (positive & negative) thus mandating the Majority rule.
- 3. Explore a combination of 1: Many (Concatenated-Study ID : individual-coughs) & Many:1 (individual-coughs : Concatenated-Study ID) approach for the validation & testing sets respectively.

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