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Wearable stethoscope for lung disease diagnosis

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Lung disease is one of the most widespread types of disease, especially in the era of COVID-19. Its diagnosis is of great importance, as different types have diverse treatments and prognoses. The most popular methods are computed tomography scanning, ultrasonogram, and bioimpedance sensors, but they are not suitable for wearable applications. Here, we developed a wearable stethoscope with an accompanying algorithm for lung disease diagnosis. It was demonstrated on 18 patients in hospital with three types of lung disease. After collecting and pre-processing lung sound signals, several machine learning methods with optimized features were applied and achieved high classification metrics. The features of the low-frequency wavelets decomposed from the lung sound signals were found to be important, serving as potential biomarkers for different types of lung disease. Overall, it was proven that our wearable stethoscope could provide a more user-friendly method and find greater application scenarios for lung disease diagnosis.

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Introduction

Lung diseases are one of the most common types of disease in the world. As the lung is a complex system, lung disease can be divided into many types with totally different causes and symptoms, affecting their diagnosis, treatment, and prognosis. In general, lung diseases can be classified by the affected region, such as the airways, air sacs, interstitium and others.¹ For lung diseases affecting the airways, chronic obstructive pulmonary disease (COPD) is one of the most common. COPD causes obstructed air flow from the lungs, resulting in symptoms such as difficulty breathing,² and is often triggered by long-term exposure to cigarette smoke.³ However, it is treatable by bronchodilators using inhalers after diagnosis. Among lung diseases affecting the air sacs, pneumonia is the most widespread, especially in the era of COVID-19. Pneumonia may cause the air sacs to fill with fluid or pus, and affects mostly children younger than 2 years old and people older than 65. After diagnosis, the symptoms may ease in a few days, while a feeling of tiredness can remain for a longer time. It is generally treated using antibiotics. Among lung diseases affecting the interstitium, interstitial lung disease (ILD) is prevalent, which causes progressive scarring of lung tissue due to long-term exposure to hazardous materials, such as asbestos.⁴ ILD is generally irreversible, so an early diagnosis is critical. Many ILD patients are initially

treated with a corticosteroid and other drugs to suppress the immune system.

Generally, as lung diseases occur in different parts of the respiratory system, early and accurate diagnosis is critical for different treatment strategies with various prognoses. Currently, diagnosis is performed using pulmonary function tests, which test the amount of the air inhaled and exhaled by the lungs, arterial blood gas analysis, pulse oximetry, and sputum testing. Furthermore, chest X-rays, computed tomography (CT) scans,⁵ and echocardiograms can be utilized to determine the severity and location of lung diseases.^{6,7} However, for diagnosis of relatively healthy people, portable or even wearable biomedical devices may find more application scenarios than conventional instruments in the hospital, and provide a key factor for the early diagnosis of lung diseases.^{8,9} Also, for people in hospital, a wearable device provides a more friendly method than disturbing the patient by moving them from their ward to the department with the relevant instruments, especially for patients in an intensive care unit (ICU).

Current wearable technology is based on sensing the physical vital signs or biochemical signals of the subject. Optical methods such as photoplethysmography (PPG) use visible or IR light to sense the signal, but penetrate only a few millimetres into the skin and thus cannot reach the lungs; it is mostly used to detect arterial blood vessels. Electrophysiological methods such as electrocardiography (ECG), electroencephalograms (EEG) or electromyography (EMG) cannot be applied to the lungs,¹⁰ because they do not generate strong enough electrical signals. Alternatively, bioimpedance devices have been studied to monitor tidal

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volume and respiratory rate^{11,12} and used to classify respiration disorders^{13,14} such as apnea and hypopnea.¹⁵ However, bioimpedance measurements require the injection of a weak current into the body, which can be noticeable to the subject, affecting the user experience and making this method unsuitable for long-term monitoring. Respiration causes vibrations or micromotions on the chest, which can be directly sensed by strain or inertial sensors,¹⁶ but the resulting information is not as rich as that of lung sounds.^{17–19} As a result, wearable lung sensing is focused on the generated sounds using ultrasonogram (USG) methods. However, the ultrasonic technique is harmful to the tissue and not recommended for usage longer than 30 min, which may result in abnormal lung sounds being missed. Additionally, its complicated piezoelectric sensor array makes it not widely affordable. Nevertheless, the stethoscope, which also detects sound signals, is the tool mostly widely used by doctors to detect lung diseases. It passively receives lung sounds, and thus it could be worn for hours or days, and experienced doctors could make an initial diagnosis of the type of lung disease directly using the stethoscope alone. In addition to the sensing method, current research is focused on different types of lung sounds, such as crackling, rhonchi, wheezing and stridor.^{20–27} However, the lung sound type is not directly related to the diagnosis of the type of lung disease, and one type of disease is sometimes related to several types of abnormal lung sounds.

In order to take advantage of the stethoscope, make it digital, and solve the issue of lung disease diagnosis, we optimized a wearable small-scale electronic stethoscope (WSES) system previously developed by us¹⁷ and applied it to 18 patients with lung diseases in hospital. The system has a microphone IC chip with an integrated microelectromechanical system (MEMS). Its small size, low power consumption, and high signal-to-noise ratio (SNR) makes this wearable stethoscope applicable for patients in hospital. A user-friendly mobile application was developed to

collect signals from the device for further diagnosis, making our device cable-free. In order to diagnose the type of lung disease, several machine learning methods were compared to obtain the best performance. Our results showed that the system could reach a diagnosis accuracy higher than 90% and could be applied in hospital to assist doctors in lung disease diagnosis. Fig. 1 shows an overall flow chart of our study design. Lung sounds from patients with different lung diseases were collected and pre-processed to remove noise and artifacts. Then, features in the time domain, from wavelet decomposition and demography, were extracted. Next, different feature sets were compared, and nine machine learning algorithms were applied to find the optimized algorithms with the highest performance metrics.

Methods

Subjects and ethical considerations

This study was conducted on 18 patients with lung disease recruited from Nanjing Drum Tower Hospital. There were nine male and nine female patients with a mean age of 74.2 years (standard deviation \pm 10.4 years), with the specific demographic and pathological characteristics detailed in Table 1. All subjects had normal hearing, normal vision, and normal speech function. The subjects were conscious and able to communicate with doctors, and the gold standard for lung disease type was evaluation by the doctor before the experiment. Prior to the experiment, subjects were informed of the procedure and related precautions, and that the experiment was not harmful to humans. All participants signed an informed consent form and the study was approved by the Ethics Committee of Southeast University. Data were obtained in accordance with the guidelines of the University Ethics Committee and the ethical principles of the Declaration of Helsinki for medical research involving human beings.

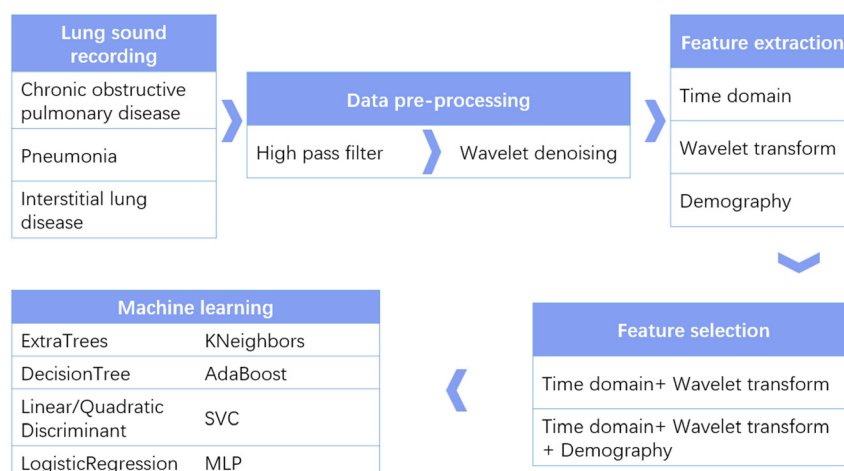


Fig. 1 Flow chart of diagnosis of the type of lung disease using our wearable stethoscope system.



Table 1 Demographic and pathological characteristics of 18 subjects. HR: heart rate/BPM, RR: respiration rate/RPM, SBP: systolic blood pressure/mmHg, DBP: diastolic blood pressure/mmHg

No	Sex	Age	HR	SpO ₂	RR	SBP	DBP
1	Male	91	60	99	30	137	55
2	Female	84	114	95	50	110	64
3	Male	87	75	98	17	130	77
4	Female	88	114	97	18	128	79
5	Female	75	96	98	36	105	75
6	Male	67	70	99	25	105	67
7	Female	70	89	100	13	143	78
8	Female	72	84	99	26	101	51
9	Female	82	76	99	18	138	64
10	Male	75	65	98	24	113	62
11	Female	55	53	95	22	110	66
12	Male	79	70	99	14	111	N/A
13	Female	80	96	98	21	159	71
14	Male	69	71	83	30	152	91
15	Male	63	106	88	15	120	72
16	Male	55	62	96	17	98	64
17	Female	74	72	99	24	126	73
18	Male	69	87	89	24	125	83

Hardware and experimental procedure

Fig. 2 presents the design of the device and a photo of a patient wearing it. The sensing chip is an ICS-43432 (InvenSense) with a size of 3.5 mm by 3.5 mm, and the microcontroller unit (MCU) is nRF52840 (Nordic Semiconductor) with a size of 3.5 mm by 3.6 mm. The ICS-43432 senses the sound from the body and transmits the digital signal to the nRF52840, which has a Bluetooth module to send the data to a mobile phone. The device operates at a sampling rate of 2.4 kHz, which is sufficient for lung sound collection. Blue medical tape (Huaxi Sanitary Materials) was placed between the device and the patient's skin for sanitary purposes, and the battery used was 250 mA h. The device cost is tens of US dollars, and it could be afforded by the general public. The firmware was developed with uVision 5.28 based on NordicSemi nRF SDK 15.3. The algorithm was demonstrated on MATLAB R2021a and Scikit-Learn 0.24.2 in Python 3.7.

In a quiet environment in the ward, the subjects wore our device lying flat, keeping their body posture unchanged as much as possible. We collected around three minutes of lung sounds for each patient. The signal quality was initially judged according to the frequency domain display in the app interface, which was used to decide whether to repeat the data acquisition process.

Data pre-processing and feature extraction

The app saved the lung sound data to the memory of the smartphone as a text file. The name of the file contains time information, such as "Tue Nov 08 10:09:47 GMT + 08:00 2023-282". The first six digits of the last string are timestamps and the seventh digit is the order of collected data. The text file is read into MATLAB, and the sound data of the corresponding period were searched and intercepted according to the timestamps recorded during the experiment. The data was processed using a 100 Hz high-pass filter and a wavelet denoising algorithm.

After the pre-processing, the features in the time domain, frequency domain, and nonlinear domain of the signal were extracted, as listed in Table 2. Other information listed in Table 1, such as HR, SpO₂, RR, SBP and DBP, was collected but not included as features because during the collection period they were not continuously monitored. They were only utilized to confirm that the patient status was stable. Here, nine features, namely, maximum value, minimum value, maximum minus minimum value, standard deviation, mean of absolute, median of absolute, mean, kurtosis, and mean of absolute of the derivative in the time domain were extracted. Next, the mean of absolute of coefficients, mean of power of coefficients, and standard deviation of coefficients of five wavelets were extracted, resulting in 15 features. Also, the ratio between the mean of absolute of coefficients was

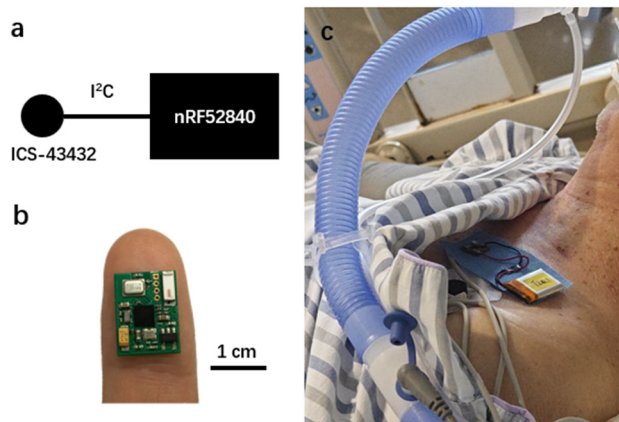


Fig. 2 a) Schematic and b) photo of the wearable stethoscope. c) Photo of a patient wearing our device.



Table 2 Time domain, frequency domain and nonlinear domain features

Time domain features		
Max	Min	Max-min
Standard deviation	Mean of absolute	Median of absolute
Mean	Kurtosis	Mean of absolute of derivative
Wavelet decomposition features		
Mean of absolute of coefficients	Mean of power of coefficients	Standard deviation of coefficients
Demographic features		
Sex	Age	

derived as four features. Adding sex and age as features, in total there are 30 features involved.

Machine learning models

Supervised learning is the most applicable method for lung disease classification, in which input data with labels are propagated by an algorithm, which then learns the patterns associated with each label. A supervised machine learning model was trained with extracted features as the dataset and the predetermined lung disease types as labels. This study utilizes the base classification models ExtraTreesClassifier, DecisionTreeClassifier, LinearDiscriminantAnalysis, LogisticRegression, QuadraticDiscriminantAnalysis, KNeighborsClassifier, AdaBoostClassifier, SVC, and MLPClassifier from the Python machine learning library scikit-learn. Parameter optimization was performed using RandomizedSearchCV. The performance of the above machine learning models was judged by accuracy, precision, recall, and F1 score. Accuracy, recall, precision, and F1 scores were used to evaluate our algorithms. They are related to true positives (TP), true negatives (TN), false positives (FP), and false negatives (FN). Accuracy is calculated as $(TP + TN)/(TP + TN + FP + FN)$, recall is quantified as $(TP/(TP + FN))$ and precision is $(TP/(TP + FP))$. The F1 metric is the harmonic mean of precision and recall for overall performance evaluation. Also, a confusion matrix was studied.

The features were first standardized by removing the mean and scaling to unit variance. The dataset was divided into training and test subsets at a ratio of 9:1. The classifier was trained using the training subsets and then utilized for prediction on the test subsets. The algorithm's capability and performance to tell the difference between different lung diseases was studied. *K*-Fold was adopted for cross-validation. The dataset was split randomly into *K* parts, with one part as test subsets and the remaining as training subsets.

Results and discussion

Feature extraction and selection

The acquired lung sound signals were filtered using a 100 Hz high-pass filter, as lung sound frequency is typically higher than 100 Hz, and then decomposed using 'wavedec' at level 5 with the wavelet 'coif4' in MATLAB. Then, 'wthresh' and

'waverec' were utilized to denoise and reconstruct the lung sound signals. Fig. 3 shows the raw data and the signals denoised using wavelet thresholds.

After feature extraction, the ranking of feature importance was calculated, and is shown in Fig. 4a. Two sets of features (28 features vs. 30 features) were trialed (Table 3). Classification using 28 features in the time domain and wavelet decomposition results in an accuracy of 0.85 ± 0.04 , recall of 0.84 ± 0.04 , precision of 0.84 ± 0.03 , and F1 of 0.83 ± 0.03 , which is rather good already. Interestingly, Fig. 4b shows that after adding the demographic information, all performance metrics were boosted, with an accuracy of 0.99 ± 0.01 , recall of 0.98 ± 0.02 , precision of 0.99 ± 0.01 , and F1 of 0.99 ± 0.01 . Indeed, age and gender are the most important features, which indicates the relationships between lung disease type and demographic information. Thus, it was decided to include all 30 features for all machine learning methods. It should be noted that obviously, age and sex alone could not determine the lung disease type; all three lung disease groups included patients with a wide range of ages and of both sexes. In addition to age and sex, the features related to low-frequency wavelets ranked higher, such as coefficients from the decomposition at level 5 and level 4, which indicates that the lower-frequency lung sounds

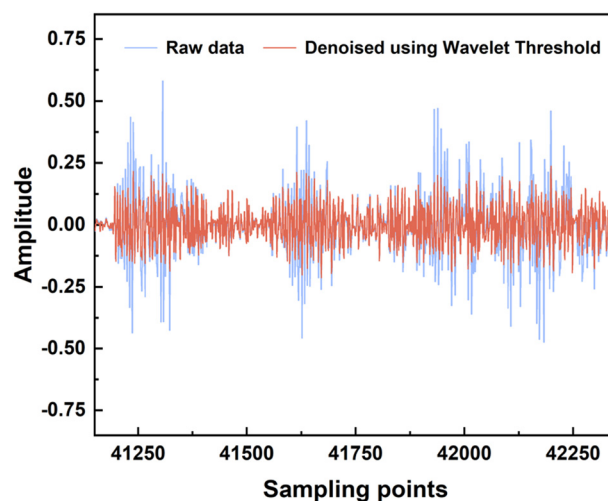


Fig. 3 Lung sound pre-processing using filters and then denoising using wavelet threshold.



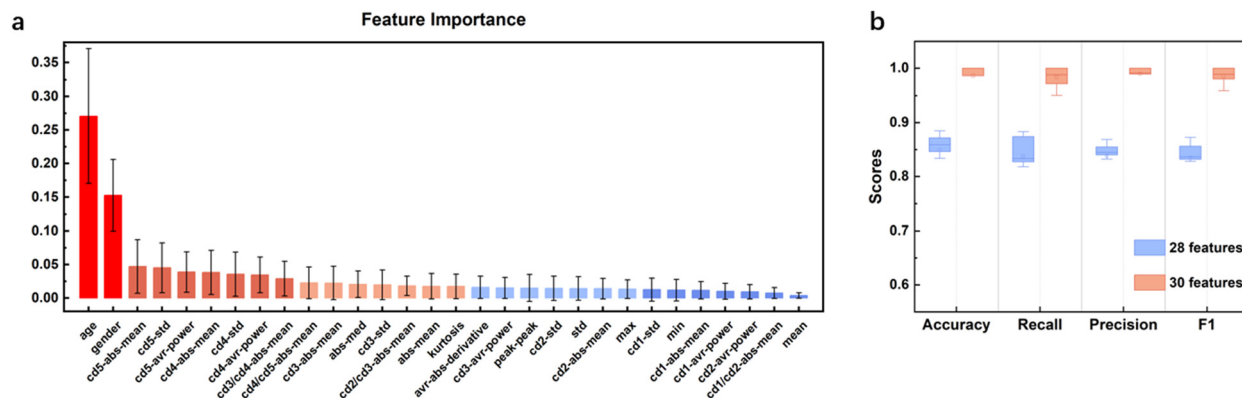


Fig. 4 a) Feature importance of ETC and b) classification performance of ETC for two sets of features.

Table 3 Comparison for 28 vs. 30 features

	Accuracy	Recall	Precision	F1
28	0.85 ± 0.04	0.84 ± 0.04	0.84 ± 0.03	0.83 ± 0.03
30	0.99 ± 0.01	0.98 ± 0.02	0.99 ± 0.01	0.99 ± 0.01

are biomarkers for diagnosis. It could also be observed that the higher the frequency of a wavelet feature, the lower its ranking in the frequency importance.

Classification results

After feature selection, nine machine learning methods were compared, and the model parameters were optimized. The classification results for the top four algorithms are shown in Fig. 5. K-Fold validation was utilized on accuracy, recall, precision and F1 to test the robustness of the model. The optimized hyperparameters for ExtraTreesClassifier were $n_estimators = 125$, $min_samples_split = 3$, $min_samples_leaf = 1$, $criterion = entropy$, with a mean cross-validation F1 score of 0.986 ± 0.016 . The optimized hyperparameters of MLPClassifier were $alpha = 0.0001$,

$hidden_layer_sizes = (200, 200, 200)$, $max_iter = 1000$, $learning_rate_init = 0.001$, $tol = 0.0001$, $beta_1 = 0.99$, $beta_2 = 0.99$, $epsilon = 1 \times 10^{-7}$ with a mean cross-validation F1 score of 0.966 ± 0.015 . The optimized hyperparameters of KNeighborsClassifier were $algorithm = ball_tree$, $leaf_size = 40$, $n_neighbors = 30$, $p = 1$, $weights = distance$ with a mean cross-validation F1 score of 0.942 ± 0.022 . The optimized hyperparameters of svm.SVC were $C = 2$, $kernel = rbf$, $gamma = scale$, $tol = 0.0001$ with a mean cross-validation F1 score of 0.954 ± 0.018 . ETC performed the best, followed by MLPC and KNN. Fig. 5b presents the confusion matrix of ETC, with especially high metrics for PI and IPF. The only missed label was an instance of “others” wrongly classified as “PI”. Thus, it is proven that using our wearable device with the corresponding optimized diagnosis algorithm, our system could accurately diagnose lung disease and could find further applications in clinics.

Nevertheless, several insights and limitations were identified. First, different biocompatible adhesives could be applied to attach our device to the patient's skin to minimize the environmental noise and match the sound impedance in order to obtain the highest SNR. Second, different types of sound could be categorized in detail to determine the

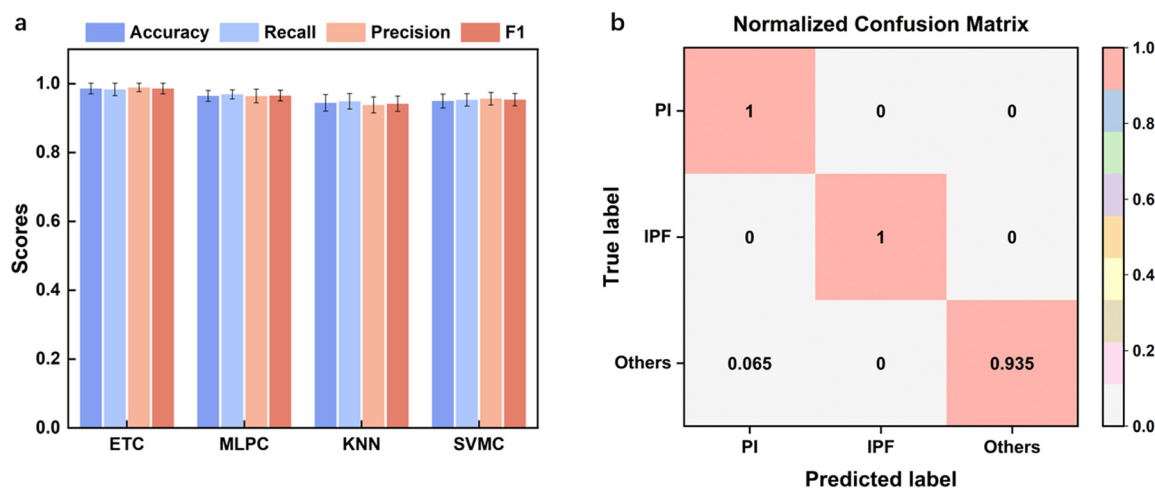


Fig. 5 Model performance characterization. a) Performance metrics of the best four models. b) Confusion matrix of the ETC model.



relationships between the sound type and lung disease type. Third, most of the patients were more than 60 years old, and the robustness of our system in lung disease diagnosis should be proven with other ranges of ages.

Conclusions

Lung disease type classification is critical for deciding the treatment strategy and prognosis monitoring. Currently, most lung disease studies with wearable technology are focused on sound type classification, which is not clinically significant. Here, we introduced a wearable stethoscope and applied it to 18 lung disease patients with the three most widespread disease types. Machine learning methods were used to achieve high performance metrics, proving its potential for application in clinics.

Author contributions

C. Z. and H. L. conceived the research idea. C. Q. and W. Z. performed experiments. W. T. analyzed the data. C. Q. and C. Z. wrote the paper. All authors discussed the results and reviewed the manuscript.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 T. K. Burki, *Lancet Respir. Med.*, 2019, **7**, 1015–1016.
- 2 D. Blanco-Almazan, W. Groenendaal, M. Lozano-Garcia, L. Estrada-Petrocelli, L. Lijnen, C. Smeets, D. Ruttens, F. Catthoor and R. Jane, *IEEE Trans. Biomed. Eng.*, 2021, **68**, 298–307.
- 3 J. E. Bibault and L. Xing, *Lancet Digital Health*, 2020, **2**, e216–e217.
- 4 M. S. Wijsenbeek, C. C. Moor, K. A. Johansson, P. D. Jackson, Y. H. Khor, Y. Kondoh, S. K. Rajan, G. C. Tabaj, B. E. Varela, P. van der Wal, R. N. van Zyl-Smit, M. Kreuter and T. M. Maher, *Lancet Respir. Med.*, 2023, **11**, 97–110.
- 5 Q. Dou, T. Y. So, M. Jiang, Q. Liu, V. Vardhanabhuti, G. Kaissis, Z. Li, W. Si, H. H. C. Lee, K. Yu, Z. Feng, L. Dong, E. Burian, F. Jungmann, R. Braren, M. Makowski, B. Kainz, D. Rueckert, B. Glocker, S. C. H. Yu and P. A. Heng, *npj Digit. Med.*, 2021, **4**, 60.
- 6 R. Aggarwal, V. Sounderajah, G. Martin, D. S. W. Ting, A. Karthikesalingam, D. King, H. Ashrafian and A. Darzi, *npj Digit. Med.*, 2021, **4**, 65.
- 7 H. Barnes, S. M. Humphries, P. M. George, D. Assayag, I. Glaspole, J. A. Mackintosh, T. J. Corte, M. Glassberg, K. A. Johansson, L. Calandriello, F. Felder, A. Wells and S. Walsh, *Lancet Digital Health*, 2023, **5**, e41–e50.
- 8 S. H. Lee, Y. S. Kim, M. K. Yeo, M. Mahmood, N. Zavanelli, C. Chung, J. Y. Heo, Y. Kim, S. S. Jung and W. H. Yeo, *Sci. Adv.*, 2022, **8**, eabo5867.
- 9 S. H. Lee, Y. S. Kim and W. H. Yeo, *Adv. Healthcare Mater.*, 2021, **10**, e2101400.
- 10 Y. Wang, W. Tian, J. Xu, Y. Tian, C. Xu, B. Ma, Q. Hao, C. Zhao and H. Liu, *IEEE Sens. J.*, 2023, **23**, 21767–21775.
- 11 J. A. Berkebille, S. A. Mabrouk, V. G. Ganti, A. V. Srivatsa, J. A. Sanchez-Perez and O. T. Inan, *IEEE Trans. Biomed. Eng.*, 2022, **69**, 1909–1919.
- 12 A. R. Fekr, K. Radecka and Z. Zilic, *IEEE J. Biomed. Health Inform.*, 2015, **19**, 1532–1548.
- 13 A. R. Fekr, M. Janidarmian, K. Radecka and Z. Zilic, *IEEE J. Biomed. Health Inform.*, 2016, **20**, 733–747.
- 14 C. Qiu, F. Wu, W. Han and M. R. Yuce, *IEEE Trans. Biomed. Eng.*, 2022, **69**, 2970–2981.
- 15 T. Van Steenkiste, W. Groenendaal, P. Dreesen, S. Lee, S. Klerkx, R. de Francisco, D. Deschrijver and T. Dhaene, *IEEE J. Biomed. Health Inform.*, 2020, **24**, 2589–2598.
- 16 D. Jarchi, S. J. Rodgers, L. Tarassenko and D. A. Clifton, *IEEE Sens. J.*, 2018, **18**, 4981–4989.
- 17 J. Xu, W. Zeng, C. Zhao, J. Tong and H. Liu, *Sens. Diagn.*, 2023, **2**, 657–664.
- 18 J. Xu, C. Zhao, B. Ding, X. Gu, W. Zeng, L. Qiu, H. Yu, Y. Shen and H. Liu, *IEEE Sens. J.*, 2022, **22**, 11526–11534.
- 19 C. Zhao, W. Zeng, D. Hu and H. Liu, *IEEE Sens. J.*, 2021, **21**, 15962–15971.
- 20 J. A. Dar, K. K. Srivastava and A. Mishra, *Comput. Biol. Med.*, 2023, **164**, 107311.
- 21 N. S. Haider, *Biomed. Signal Process. Control.*, 2021, **64**, 102313.
- 22 D.-M. Huang, J. Huang, K. Qiao, N.-S. Zhong, H.-Z. Lu and W.-J. Wang, *Mil. Med. Res.*, 2023, **10**, 44.
- 23 A. Kandaswamy, C. S. Kumar, R. P. Ramanathan, S. Jayaraman and N. Malmurugan, *Comput. Biol. Med.*, 2004, **34**, 523–537.
- 24 A. Mondal, P. Banerjee and H. Tang, *Comput. Methods Programs Biomed.*, 2018, **159**, 199–209.
- 25 G. Shah, P. Koch and C. B. Papadias, *IEEE J. Biomed. Health Inform.*, 2015, **19**, 151–157.
- 26 K. H. Tsai, W. C. Wang, C. H. Cheng, C. Y. Tsai, J. K. Wang, T. H. Lin, S. H. Fang, L. C. Chen and Y. Tsao, *IEEE J. Biomed. Health Inform.*, 2020, **24**, 3203–3214.
- 27 W. Wang, D. Qin, S. Wang, Y. Fang and Y. Zheng, *Comput. Biol. Med.*, 2023, **164**, 107282.

