## **Supplementary Methods**

### Diagnostic criteria

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- 3 1. **Smoking history**: Smoking at least 1 cigarette every day for a consecutive year.
- 4 2. **Epilepsy**: Patients clinically present with seizures and exhibit seizure-like discharges on
- 5 EEG recordings during or between seizures. Diagnosis of epilepsy is made if seizures are
- 6 observed, or if there is a history of taking antiepileptic drugs in the past<sup>1</sup>.
- 7 3. **High blood pressure**: In the absence of antihypertensive medication, if the systolic blood
- 8 pressure is ≥ 140mmHg and/or diastolic blood pressure is ≥ 90mmHg in three separate
- 9 measurements on different days, the patient is diagnosed with hypertension. Even if the blood
- 10 pressure is below 140/90mmHg and the patient has a history of hypertension and is currently
- taking antihypertensive medication, they should still be diagnosed with hypertension<sup>2</sup>.
- 4. **Diabetes**: According to the diagnostic criteria proposed by the World Health Organization
- 13 Expert Committee on Diabetes in 1999: fasting blood glucose ≥7.0mmol/L (confirm on another
- day if no typical diabetes symptoms are present); or typical diabetes symptoms with random blood
- 15 glucose or venous plasma glucose ≥11.1mmol/L; or 2-hour blood glucose ≥11.1mmol/L in an oral
- 16 glucose tolerance test (confirm on another day if no typical diabetes symptoms are present); or
- 17 glycosylated hemoglobin  $\geq 6.5\%^3$ .
- 18 5. Coronary artery disease: Coronary angiography shows at least one coronary artery
- stenosis  $\geq 50\%$ , or a history of coronary artery bypass grafting<sup>4</sup>.
- 20 6. Pulmonary embolism: According to the "Diagnosis and Treatment Guidelines for
- 21 Pulmonary Thromboembolism," a CT pulmonary angiography showing filling defects in the
- 22 pulmonary artery is used to diagnose pulmonary embolism<sup>5</sup>.
- 23 7. **Pulmonary contusion**: With a clear history of chest trauma, the patient presents with
- 24 symptoms such as chest pain, shortness of breath, hemoptysis, and even respiratory distress.
- 25 Physical examination reveals crackles in the lungs and decreased breath sounds. Diagnosis of
- pulmonary contusion is confirmed through X-ray or CT imaging<sup>6</sup>.
- 8. Pneumonia: According to the "Guidelines for the Diagnosis and Treatment of
- 28 Community-Acquired Pneumonia in Chinese Adults," chest radiography showing newly
- 29 developed patchy infiltrates, lobar or segmental consolidation, ground-glass opacity, or interstitial

changes, along with any of the following features: (1) fever; (2) new onset cough, sputum production, or worsening of existing respiratory symptoms; (3) signs of lung consolidation and/or auscultation of crackles; (4) peripheral blood leukocytes  $>10\times10^9/L$  or  $<4\times10^9/L$ <sup>7</sup>

9. Consciousness disorder: The Glasgow Coma Scale was used in this study to assess the patients' level of consciousness. GCS consists of three aspects: eye opening response (1-4 points), verbal response (1-5 points), and motor response (1-6 points), with a total score of 15 points. A score of ≤8 points indicates severe consciousness disorder, 9-12 points indicates moderate consciousness disorder, 13-14 points indicates mild consciousness disorder, and 15 points indicates clear consciousness. A higher score indicates a milder consciousness disorder<sup>8</sup>.

10. **Swallowing disorder**: This study used the Wakita water swallowing test to evaluate patients' swallowing function. The Wakita water swallowing test is graded into 5 levels: Level I (Excellent): can swallow water in one attempt; Level II (Good): can swallow water without coughing at least 2 times; Level III (Fair): can swallow water in one attempt but coughs; Level IV (Acceptable): can swallow water at least 2 times but coughs; Level V (Poor): frequently coughs and cannot swallow all the water<sup>9</sup>.

## Categorical variable label encoding

During the correlation analysis, categorical variables were converted into dummy variables for use in machine learning. For binary classification variables, 1 represented "yes" and 0 represented "no". For multiclass variables, different categories were represented by Arabic numerals from 1 to the total number of categories.

Variables	Category	<b>Dummy variables</b>	
Maal gandar	yes	1	
Meal gender	no	0	
W. D. I	yes	1	
Wet Rales	no	0	
Smoking	yes	1	
	no	0	
C : D: 1	yes	1	
Consciousness Disorder	no	0	
Dryanhaaia	yes	1	
Dysphagia	no	0	
Eniloness	yes	1	
Epilepsy	no	0	

	yes	1		
Viscous sputum	no	0		
		1		
Upper Respiratory Tract	yes			
Structural Abnormalities		0		
(URSA)				
	no			
II .	yes	1		
Hypertension	no	0		
		1		
Caranary Haart Disaasa	yes			
Coronary Heart Disease		0		
	no			
Diabetes Mellitus	yes	1		
Diabetes Memtus	no	0		
Pulmonary Infection	yes	1		
runnonary infection	no	0		
Dulmanany Embalian	yes	1		
Pulmonary Embolism	no	0		
Dalara Garata da	yes	1		
Pulmonary Contusion	no	0		
M (10) 1	yes	1		
Mental Disorders	no	0		
	DD ≤ 30 days	1		
D. D. (.	$30 \text{ days} < \text{DD} \le 60 \text{ days}$	2		
Disease Duration:	60 days < DD ≤ 90 days	3		
	DD > 90 days	4		
	Cerebral Trauma	1		
	Cerebral Hemorrhage	2		
	Cerebral Infarction	3		
	Subarachnoid Hemorrhage	4		
Primary Disease:	Ischemic Hypoxic Brain Disease	5		
	Central Nervous System	6		
	Infectious Diseases			
	Spinal Cord Injury	7		
	Other Diseases	8		
	Quadriplegia	1		
Limb Paralysis:	Hemiplegia	2		
	Paraplegia	3		
	Pseudomonas aeruginosa	1		
Sputum Culture Pathogens:	Klebsiella pneumoniae	2		
_	pneumoniae subtype			

Acinetobacter baumannii	3
Escherichia coli	4
Serratia marcescens	5
Other Pathogens	6
Two or more pathogens	7
None	8

50 Table S1. Categorical variables and dummy variables.

## Hyperparameter optimization and parameter tuning

In this study, a total of eight models were constructed, including K-Nearest Neighbors (K-NN), Support Vector Machine (SVM) <sup>10</sup>, Gaussian Naive Bayes (GNB), Decision Tree (DT), Random Forest (RF) <sup>11</sup>, Extra Trees Classifier (ET), Gradient Boosting (GB), and Logistic Regression (LR), to compare the predictive performance of each model. Modeling parameters were selected using cross-validation and hyperparameter optimization. The screened features collected in this study were used as feature variables, with extubation outcomes as the target variable. In the dataset of 501 patients, 80% of the data was used as the training set, while 20% was used as the test set. Accuracy was calculated using the relevant formulas <sup>10</sup>. The raw data processing was performed using the Numpy package in Python 2020, while dataset splitting and model construction were carried out using the Sklearn package<sup>12</sup>. The details of hyperparameter optimization were provided in Supplementary Material, Table S2.

Feature	Range	Optimal value	
KNN			
k_values	1-40	24	
SVM			
Kernel	linear', 'poly', 'rbf', 'sigmoid'	linear'	
max_iter	10-1000	500	
Regularization parameter (C)	0.01-100	0.1	
probability	Ture-False	Ture	
Gaussian Naive Bayes			
var_smoothing	10 <sup>-1</sup> -10 <sup>-9</sup>	0.01	
<b>Decision Tree</b>			
max_depth	3-20	3	
min_samples_leaf	1-50	3	
criterion	gini', 'entropy'	entropy'	
min_samples_split	2-50	5	
splitter	best', 'random'	best'	
Random Forest			

max_depth	3-20	5	
max_features	sqrt', 'log2',	sqrt'	
min_samples_leaf	1-20	2	
n_estimators	100-5000	700	
random_state	1-20	3	
max_leaf_nodes	10-100	30	
min_samples_split	2-20	10	
Extra Tree Classifier			
max_depth	3-20	10	
max_features	sqrt', 'log2',	sqrt'	
min_samples_leaf	1-20	2	
n_estimators	100-5000	1000	
random_state	1-20	3	
max_leaf_nodes	10-100	30	
min_samples_split	2-20	10	
Gradient Boosting			
learning_rate	0.001-0.1	0.01	
loss	deviance', 'exponential'	exponential'	
max_depth	3-20	9	
max_leaf_nodes	5-50	5	
min_samples_leaf	1-50	25	
min_samples_split	2-20	5	
n_estimators	100-5000	500	
subsample	0.1-1	1	
AdaBoosting			
n_estimators	100-5000	200	
learning_rate	0.001-0.1	0.01	
Logistic Regression			
max_iter	0.001-1000	100	
nverse of regularization strength (C)	0.001-200	100	
solver	liblinear', 'newton-cg', 'lbfgs', 'sag', 'saga'	liblinear'	
penalty	L1', 'L2'	L2'	

63 Table S2. Range of hyperparameter optimization values.

## Model building methods and code

The raw data processing was performed via the *Numpy* package, while dataset splitting and model construction were carried out via the *Sklearn* package in *Python 3.12.4*<sup>12</sup>. The study constructed a total of 8 independent models, and the specific methods and construction process of these models are as follows:

### 1. Logistic Regression

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Logistic regression is a regression method widely used for modeling binary outcomes. It applies the logit function to the linear combination of independent variables to map the predicted results to the [0, 1] interval, estimating the probability of event occurrence<sup>13,14</sup>. Compared to traditional linear regression, logistic regression has significant advantages in dealing with non-continuous dependent variables (such as presence or absence of disease, survival or not). The method's statistical inference is based on maximum likelihood estimation, with a key focus on ensuring independence, avoiding multicollinearity, and maintaining a linear relationship for continuous variables on a log-odds scale<sup>15</sup>. In numerous clinical and epidemiological studies, logistic regression is widely used to investigate disease risk factors, predict clinical outcomes, and assess intervention effects.

#### Code:

lr = LogisticRegression(max iter=100,C = 100,solver='liblinear',penalty='12');

#### 2. K-Nearest Neighbors

The K-nearest neighbors (K-NN) algorithm has been widely recognized as a non-parametric classification and regression method based on instance learning since the 1990s<sup>16</sup>. The core idea is to calculate the distance between the sample to be classified and all samples in the training set, selecting the K nearest neighbors, and making predictions based on the neighbors' categories. The standard K-NN algorithm was first applied in academia, its main advantage being its simplicity in implementation, allowing for direct prediction using distance calculations without the need for a training phase<sup>16</sup>.

#### Code:

91 knn = KNeighborsClassifier(n neighbors=24);

## 3. Support Vector Machine

Support Vector Machine (SVM) is based on the principle of "structural risk minimization" in statistical learning theory, aiming to find a separating hyperplane with the maximum margin to effectively enhance the model's generalization ability<sup>17</sup>. In the basic linear SVM model, the algorithm transforms the training data into a high-dimensional space and determines the optimal hyperplane by solving a convex quadratic programming problem. Kernel function technique allows SVM to create non-linear decision boundaries for non-linearly separable problems<sup>18</sup>.Commonly used kernel functions include radial basis function (RBF), polynomial

kernel, and sigmoid kernel, among others. The choice of different kernel functions significantly impacts the performance of the model. In addition, SVM introduces slack variables during model training to handle noise and outliers in the data, allowing it to maintain high accuracy while preventing overfitting<sup>19</sup>.

#### Code:

svc = SVC(kernel='linear', max iter=500, C=0.1, probability=True);

#### 4. Gaussian Naive Bayes

The Gaussian Naive Bayes (GNB) classifier is based on Bayes' theorem, assuming all features are conditionally independent and continuous variables follow a Gaussian distribution. The essence of this approach is to compute the posterior probability and make classification decisions by maximizing the probability. In the field of disease prediction and classification, GNB is widely used for the early diagnosis of chronic diseases, cancer, and internal medicine conditions. In real medical data, it is often difficult to fully satisfy, which may affect the accuracy of the model's predictions. Therefore, scholars improve feature selection and data preprocessing techniques to minimize the impact of correlations, or use ensemble methods to integrate results from multiple classifiers<sup>20</sup>.

#### Code:

bays = GaussianNB(var smoothing=0.01);

#### 5. Decision Tree

Decision tree is a model with a tree-like structure, consisting of nodes that split data based on certain criteria until reaching a predetermined termination condition. The construction of a decision tree mainly involves two steps: attribute selection and tree pruning. During the attribute selection process, commonly used quantification metrics such as information gain, entropy, and Gini index are used to determine the optimal splitting attribute, thereby maximizing the inter-class differences at each branch<sup>21</sup>. To avoid overfitting during the construction process, pruning techniques are usually employed to simplify the complexity of the tree<sup>21</sup>.

#### Code:

tree = DecisionTreeClassifier( max\_depth=3, min\_samples\_leaf=3, criterion='entropy',

min\_samples\_split=5, splitter='best');

#### 6. Random Forest

Random forest (RF), as a non-parametric, ensemble machine learning algorithm, has shown many significant advantages in medical data mining and risk prediction. Firstly, RF's random sampling and majority voting mechanism make it robust in handling noisy and high-dimensional data. Additionally, its built-in variable importance assessment provides clinicians with an intuitive feature selection tool, effectively reducing the feature space and revealing underlying biological mechanisms<sup>22</sup>. Recent research has successfully applied random forests to various fields such as neuroimaging, cardiovascular diseases, diabetes prediction, and monitoring of drug adverse reactions, demonstrating high practical value of RF in all these areas<sup>23,24</sup>.

Code:

- rf = RandomForestClassifier(max depth=6, max features='sqrt', min samples leaf=2,
- n estimators=700, random state=7, max leaf nodes=30, min samples split=10);

#### 7. Extreme Random Tree

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- Extreme random tree classifier is a type of decision tree ensemble method, similar to RF, where the core idea is to vote on the predictions of multiple decision trees. However, ET achieves higher levels of randomness and generalization ability through the following key features:
- Random split points: During node splitting, ET does not select the data-driven optimal split point, but randomly selects from possible split values of features to further increase model diversity<sup>25</sup>.
- Ensemble decision: ET uses the predictions of multiple trees for majority voting or averaging to enhance model robustness and stability<sup>26</sup>.
  - Given these characteristics, ET has significant advantages in computational efficiency and model robustness compared to traditional decision trees and random forests. It also performs well in handling high-dimensional features and heterogeneous data, which is particularly important for the complexity of medical data <sup>27</sup>.

#### Code:

- extraTreeclassfier = ExtraTreesClassifier(max\_depth=10, max\_features='sqrt',
- min samples leaf=2, n estimators=1000, random state=3, max leaf nodes=30,
- min samples split=10);

#### 8. Gradient Boosting

The gradient boosting algorithm works by creating a sequence of weak learners (often decision trees) and updating each model by using the residuals from the previous model as the new target, moving closer to the true value with each iteration. This approach offers powerful nonlinear fitting capabilities and robustness, making it well-suited for high-dimensional data and complex pattern recognition tasks<sup>28</sup>. GB methods typically involve the following steps: initializing the model, usually with a simple mean prediction. In each iteration, calculate the current model's residual and fit it with a new weak learner. Update the overall model based on the predictions of weak learners, and introduce a learning rate to prevent overfitting. Repeat this process until reaching a predetermined number of iterations or the error decreases to a certain level. To improve model performance, researchers often use techniques such as cross-validation and Bayesian optimization for hyperparameter tuning, while incorporating regularization techniques to mitigate the risk of overfitting complex models<sup>29</sup>. In addition, there are efficient implementations like XGBoost, LightGBM, which have significantly optimized computing speed and memory usage, making them advantageous in large-scale medical data processing<sup>30</sup>.

#### Code:

gb = GradientBoostingClassifier(learning\_rate = 0.01, loss ='exponential', max\_depth = 9,
max\_leaf\_nodes = 5, min\_samples\_leaf = 25, min\_samples\_split = 5, n\_estimators = 500,
subsample= 1);

#### **9. WPV**

The basic principle of WPV integration method lies in the fact that different models may exhibit complementary performance advantages when facing complex tasks. By using the predicted probabilities of each model and assigning different weights based on their accuracy on the validation set, we can mitigate the negative impact of individual model errors on overall predictions. In addition, when designing the WPV model, it is necessary to consider not only the synergistic effects between different models, but also to effectively address the potential issues of data redundancy and high dimensionality that may exist between models. The voting ensemble learning method can produce results using either hard or soft voting. Hard voting is ideal for predicting distinct class labels, whereas soft voting is more suitable for predicting continuous values. This study focused on tweets categorized into different groups, suggesting that hard voting

is preferable over soft voting<sup>31</sup>.

189 **Code:** 

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WPV\_model = VotingClassifier(estimators=[('rf', rf), ('bays', bays), ('knn', knn)],

voting='soft', weights=[1, 1, 1], verbose=1)

## **Supplementary Results**

Ran	Model	Accuracy	AUC	Sensitivity	Specificity	Youden	PPV	NPV	Thre
k		(%)		(%)	(%)	index (%)	(%)	(%)	shold
1	WPV	85.15	0.87	81.08	87.50	68.58	78.9 5	88.89	NA
2	Extra Tree Classifie r	83.17	0.87	70.27	90.63	60.90	81.2	84.06	0.51
3	Random Forest	84.15	0.85	72.97	90.62	63.60	81.8	85.29	0.50
4	K-Neare st Neighbo ur	82.18	0.85	70.27	89.06	59.33	78.7 9	83.82	0.58
5	Gradient Boosting	81.19	0.85	72.97	85.93	58.91	75.0 0	84.61	0.53
6	Support Vector Machine	79.21	0.83	75.67	81.25	56.93	70.0	85.24	0.43
7	Decision Tree	79.21	0.82	75.68	81.25	56.92	70.0 0	85.24	0.27
8	Logistic Regressi on	77.23	0.84	75.67	78.13	53.80	66.6	84.75	0.60
9	Gaussian Naive Bayes	73.28	0.88	86.49	65.63	52.11	59.2 6	89.36	0.79

Table S3. Summary of the performance of all models.

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# Confusion matrix for all models

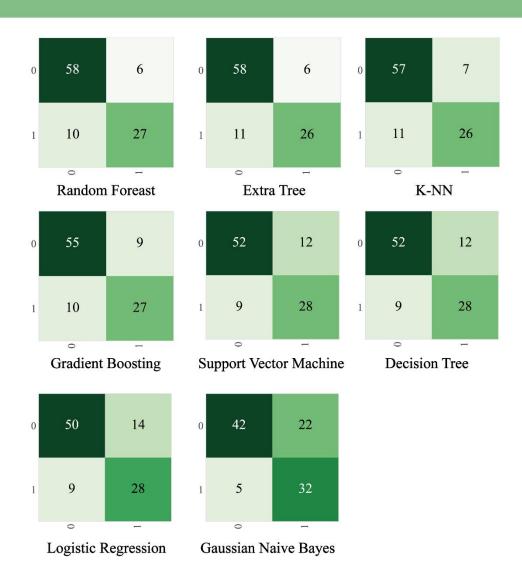
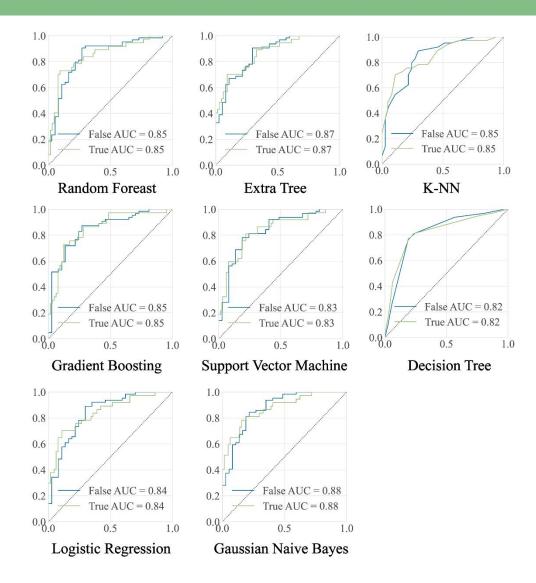


Figure S1. Confusion matrices of all models. K-Nearest Neighbors (K-NN).

## ROCAUC for all models



**Figure S2. AUC curves of all models.** Area under receiver operating characteristic curve (AUC), K-Nearest Neighbors (K-NN).

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# Learning Curve for all models

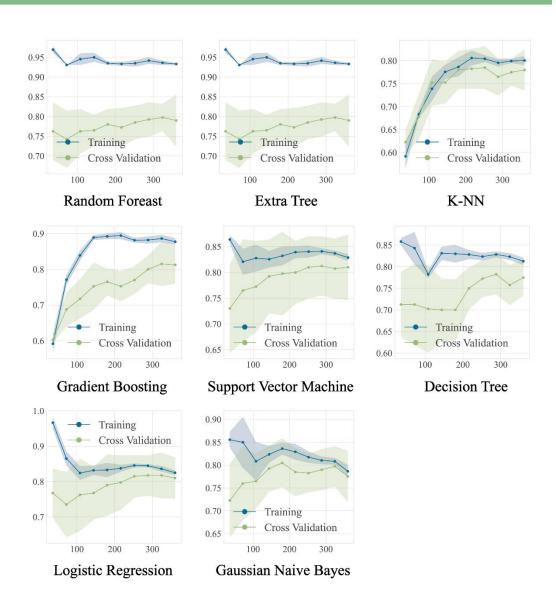


Figure S3. Learning curves of all models. K-Nearest Neighbors (K-NN).

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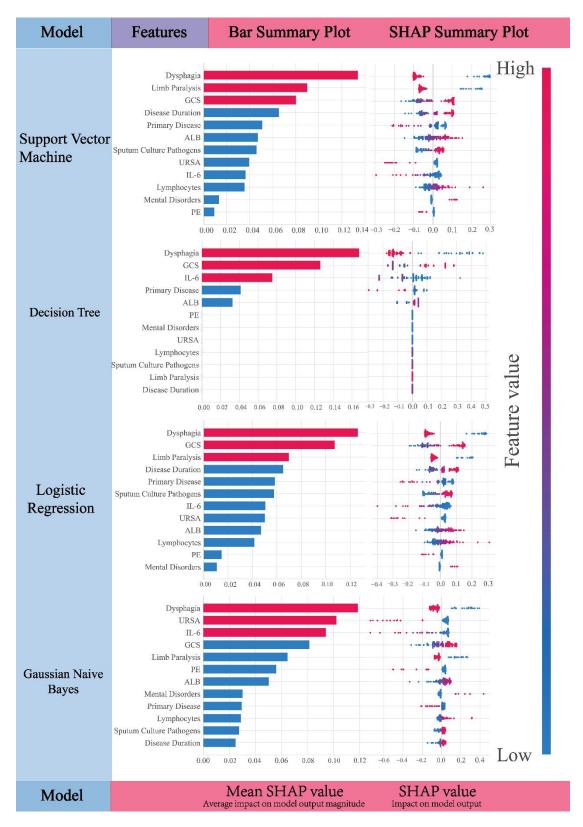


Figure S4. Bar summary plots and SHAP summary plots for the remaining four models. Shapley additive explanations (SHAP), Glasgow Coma Scale (GCS), Upper Respiratory Tract Structural Abnormalities (URSA), Interleukin 6 (IL-6), Albumin (ALB), Pulmonary Embolism (PE), K-Nearest Neighbors (K-NN).

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