

参赛学生姓名： 王 傲

中学： 上海外国语大学附属外国语学校

省份： 上海

国家/地区： 中华人民共和国

指导老师姓名： 潘丽坤（英才计划导师）

指导老师单位： 华东师范大学

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Development of Semiconductor Photothermal
Materials by Tuning Bandgap for Enhancing
Solar Water Evaporation

Machine Learning-Driven Development of Semiconductor Photothermal Materials by Tuning Bandgap for Enhancing Solar Water Evaporation

Ao Wang

Shanghai Foreign Language School Affiliated to SISU, Shanghai, 200083, China

Abstract

Solar water evaporation outstands as a promising method to solve the global freshwater shortage due to its lower cost, green and sustainable. Semiconductors photothermal materials (SPMs), with modifiable bandgap, and high performance, exhibit excellent prospects for practical solar evaporation. However, the precise influence of SPMs properties on evaporation performance remains unclear, necessitating long time and large costs to develop by trial-and-error tests.

Machine learning (ML) using data-driven approach to achieve feature classification and performance prediction. This work employed four ML models were employed to analyze the database collected from the latest research articles of SPMs, using data-driven approach to achieve feature classification and performance prediction. Shapley Additive Explanation analysis revealed the impact features for evaporation performance of SPMs. Moreover, experimental validation for the ML model's reliability was achieved doping sodium borohydride (NaBH_4) to titanium dioxide (TiO_2) for narrowing bandgap by calcining under 300 °C, 500 °C and 700 °C for one hour, respectively. Various characterizations were analyzed through SEM, XPS and UV-Vis confirmed Ti^{4+} converted into Ti^{3+} , resulting in forming oxygen vacancy to narrow bandgap of four samples from 3.36 eV (P25), 3.33 eV (300 °C), 3.12 eV (500 °C 5) to 2.09 eV (700 °C). The experimental results indicate that the evaporation rate of TiO_2 -700 °C improved 17.5% than primary literature, with evaporation rates of 1.12, 1.62, 1.89 and 2.28 $\text{kg m}^{-2} \text{h}^{-1}$ under 1sun, 2sun, 3sun and 4sun, respectively. This study highlights that ML strategies can enhance the efficiency of exploring high-performance SPMs for solar-driven water evaporation.

Keywords: Solar evaporation performance, Machine learning, Semiconductor photothermal materials, Bandgap, Absorbance

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1. Introduction

Currently the problem of global freshwater shortage is becoming more and more serious due to rapid population growth and accelerating industrialization. Seawater desalination has emerged as an effective approach to address this issue [1,2]. Solar-driven photothermal evaporation technology is an eco-friendly and sustainable solution to the freshwater crisis [3,4]. It primarily operates via three modes: bottom heating, bulk heating, and interfacial heating [5-7]. Interfacial solar evaporation (ISE) captures solar energy and convert into heat at the air-photothermal material-water interface. This improves thermal conversion efficiency, minimizes heat loss, and boosts freshwater production [8,9]. The interfacial heating process involves two distinct stages: evaporation and diffusion. Evaporation occurs when disruption breaks interfacial hydrogen bonds, and evaporation enthalpy is lower than bonded free water, resulting in thermal diffusion across the interface [10]. Photothermal materials float on the water surface and directly convert absorbed solar energy into heat, driving water evaporation at the water-air interface. The performance of ISE is governed by material microstructures, insulation layer performance, and solar-thermal conversion efficiency [11,12].

Recently, a variety of solar photothermal materials have been reported, including carbon, metal, semiconductor materials [13]. Particularly, semiconductor photothermal materials (SPMs) have gained extensive attention owing to excellent light-to-heat conversion capability, good chemical stability and broad applicability across multiple industries [6,14-16]. For ISE, the performance of SPMs is mainly influenced by their narrow bandgap and solar absorbance. When photon energy exceeds the semiconductor bandgap, absorbed photons generate electron-hole pairs under sunlight [17]. Photo-generated carriers relax toward band extrema, converting surplus energy into heat via thermalization. A narrower bandgap facilitates light absorption over a wider wavelength range. High light absorbance boosts heat generation by forming charge carriers, which then recombine to release thermal energy. This process also involves the absorption of electromagnetic waves, such as infrared radiation [13,15]. Moreover, solar irradiation, environmental temperature and humidity exert significant impacts on evaporation efficiency (EE) in the desalination process [18]. However, the interplay between these factors and solar water evaporation remains complex and unclear. Current SPMs perform well in desalination, but novel SPMs

are still needed to achieve high evaporation rate (ER) in practice. Conventional design approaches involve complex, multi-step processes including material synthesis, property analysis, and evaporation evaluation, develop by trial-and-error tests. Therefore, optimal and resource-conserving methods for designing high-performance SPMs are imperative.

Machine learning (ML) has become a transformative tool in materials science, offering a powerful means to automatically identify patterns and learning from complex datasets, and generate data-driven predictions [19-25]. ML technology integrating with theory and experimental enables predicting material properties, optimize parameters and structures, designing novel materials, and uncovering new mechanisms beyond intuition [26,27]. The operational mechanism of photothermal material can be accurately predicted by training various ML models with large datasets curated from relevant literatures. According to Gradient Boosting Decision Tree (GBDT) ensure each new tree correct the residuals of the previous trees in iterative process,, Wang et al. applied GBDT regressor to to predict the desalination capacity of faradic materials [28]. Ali et al. explored the extreme gradient boosting (XGBT) improves speed and prediction accuracy by using a new tree learning algorithm, leveraging parallel and distributed computing [29]. Elsheikh et al. applied support vector machines (SVM) for improving the performance of solar stills and water yield prediction [30]. In our previous study, voting regressor (VR) was employed to improve regression and explore carbon-based materials for solar evaporation. These works predominantly investigated solar desalination capabilities of carbon-derived photothermal absorbers, so ML-assisted exploration of SPMs remains largely underexplored.

In this study, four ML models (GBDT, XGBT, SVM and VR) were employed to predict the ER and EE of SPMs, and to establish the evaporation performance relationships with structure properties and experiment conditions. Particularly, two key material features of bandgap (BG) and absorbance (A); a property parameter of evaporation system of thermal conductivity (TC), and three operational conditions illumination conditions (sun), room temperature (T), and relative humidity (H); which were considered as inputs. The schematic process of predicting the evaporation performance of SPMs through ML methods is demonstrated in Fig. 1.

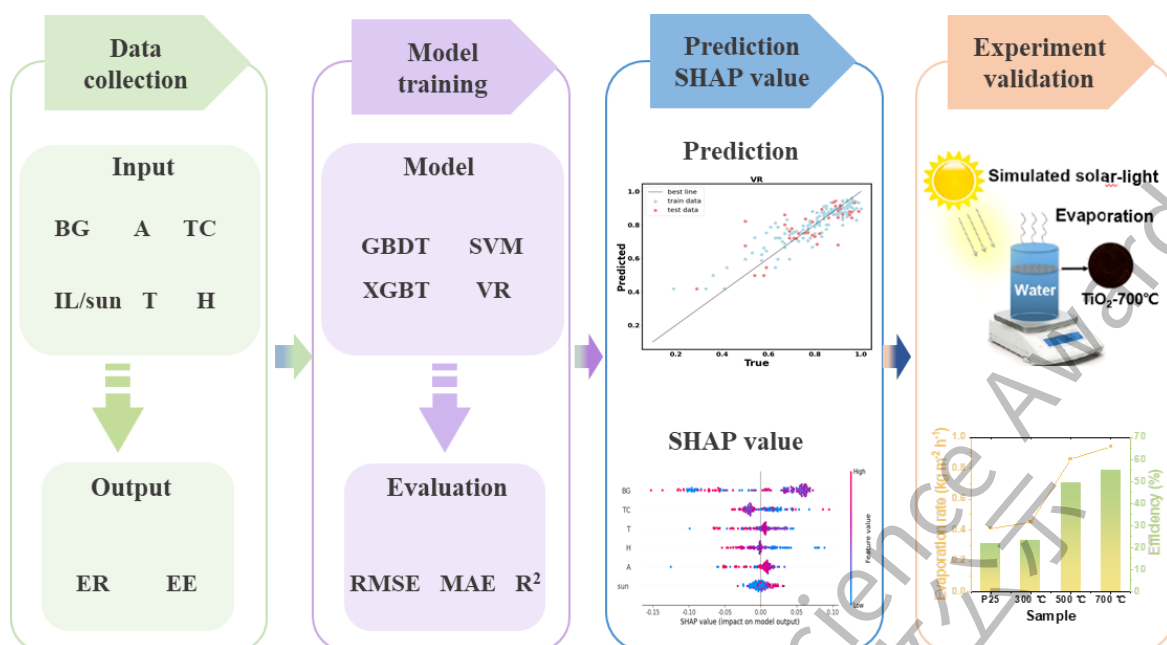


Fig. 1. Diagram for predicting ER and EE of SPMs using ML.

The key research objectives of this study are:

- Collect dataset (six input and two output features) from published literatures researched on semiconductor photothermal materials for solar water evaporation.
- Train machine learning models and evaluate performance by root mean square error (RMSE), mean absolute error (MAE) and R^2 values.
- Predictive ER and EE by four ML models, and analyze feature importance of six input features in the model was evaluated using Shapley Additive Explanation (SHAP).
- Prepare black TiO_2 samples by calcining mix of P25 and NaBH_4 under temperature 300°C, 500°C, 700°C, respectively.
- Evaporation performance will test under different solar irradiation (1 sun, 2 sun, 3 sun, 4 sun) to confirm the reliability and applicability of ML method.

By achieving these objectives, this study highlights that ML strategies can accelerate development of high-performance SPMs for solar water evaporation, improving the efficiency of seawater desalination.

2. Methods

2.1. Dataset construction and feature screening

The data collected for this research comprises 196 samples, all sourced from the

published literature. Comprehensive details of the dataset are presented in [Table S1](#), as shown in [Supplementary Information](#) document. Considering that data of features are missing exceeding 15% in [Table S1](#), Multiple Imputation was applied to handle missing data by synthesizing all available data to derive optimal imputed values [19]. In this dataset, ER and EE were used as the prediction target to evaluate solar water evaporation performance. The key properties of semiconductor materials BG, A are regarded to effectively influence the evaporation performance. Additionally, operational condition sun determines the light energy providing heat for evaporation process. Ambient environmental factors T and H also affect evaporation performance. To evaluate and identify the most accurate ML model for predicting the solar water evaporation performance, six commonly used features (BG, A, TC, sun, T, and H) served as input variables for the ML algorithms for predicting ER and EE.

2.2. Training ML models and accuracy assessment

In this work, VR used for ensemble method to combining the predictions from GBDT, XGBT, and SVM models, which leads to more robust predictions compared to individual model. The performance of the four ML models was evaluated through random allocation of the dataset with 80% for training sets and 20% for testing sets. Cross validation with five-fold was performed to train each ML model, each algorithm underwent five training and evaluation cycles. In addition, Bayesian optimization were applied to optimize the hyperparameters of each model, and [Table S2](#) and [Table S3](#) summarize the optimized hyperparameters for each model. Using the optimal hyperparameters, each model's performance was evaluated on the training sets and testing sets using RMSE, MAE, and coefficient of determination (R^2). These evaluation indicators are defined as:

$$RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^n (y_i - \hat{y}_i)^2} \quad (1)$$

$$MAE = \frac{1}{n} \sum_{i=1}^n |y_i - \hat{y}_i| \quad (2)$$

$$R^2 = 1 - \frac{\sum_{i=1}^n (y_i - \hat{y}_i)^2}{\sum_{i=1}^n (y_i - \bar{y})^2} \quad (3)$$

where, n denotes the sample size, y_i and $\hat{y}_i$ are actual value and predicted value, respectively, and $\bar{y}_i$ is the mean of y_i . Notably, for both values of RMSE and MAE are small signify higher predictive accuracy, smaller RMSE, and MAE indicate prediction, while the larger R^2 means better performance of the model ($0 \leq R^2 \leq 1$).

In the study, the weights for each base model in the VR were determined based on their performance on the validation set (specifically, the R^2).

3. Experimental

3.1. Materials and instruments

A commercial fine titanium dioxide (TiO_2) powder (P25) was used as starting material. The sodium borohydride (NaBH_4) was used as reduction agent. The P25 and NaBH_4 were acquired from Sinopharm Chemical Reagent Co. Ltd., and all chemicals were used without any further treatment. All experiments used deionized water.

The main instruments used in the sample preparation of this project are as follows:

- (1) High-precision electronic analytical balance, model FA1204B, Jucheng Group.
- (2) Tubular furnace, model OTF-1200X, Hefei Kexing Materials Technology Company.
- (3) Ultrasonic cleaning machine, model UC-7180L, Shenzhen Langjie Ultrasonic Electrical Appliance Company.
- (4) High-speed centrifuge, model TGL-15B, Shanghai Anting Scientific Instrument Factory.
- (5) Vacuum drying oven, model DHG-9055A, Shanghai Yiheng Scientific Instrument Company.

3.2. Samples preparation

Samples preparation process are demonstrated in Fig. 2, for the preparation of each sample, 0.4g TiO_2 and 0.15g NaBH_4 weighed and placed in a ceramic crucible. The two materials should be ground to ensure complete mixing. Then, they should be separately placed in ceramic crucible boxes and placed in a tubular heating furnace for baking. Four types of samples were prepared according to this mass ratio: the original sample, heated at 300°C for 1 hour, heated at 500°C for 1 hour, and heated at 700°C for 1 hour.

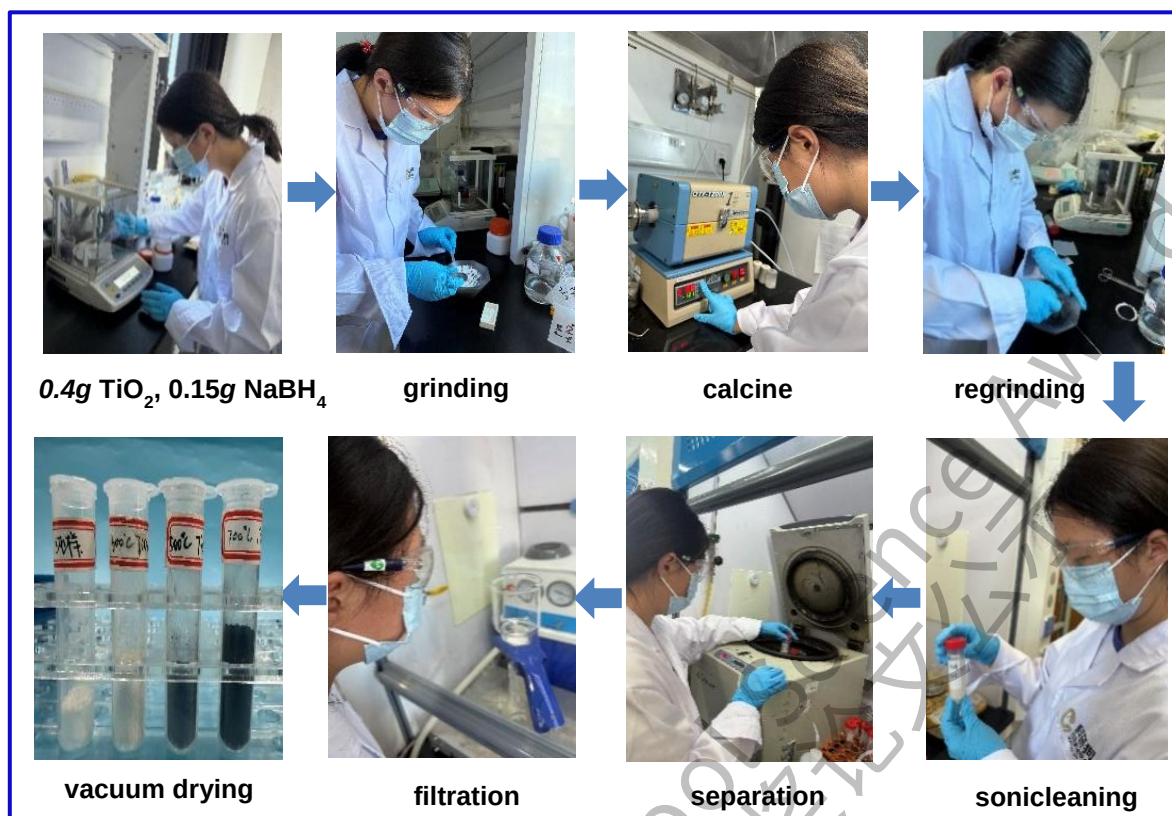


Fig. 2. Samples preparation process.

Place the crucible in a tubular heating furnace, protected by nitrogen gas (flow rate of 0.2 L/min), and raise the temperature at a rate of 5°C per minute to the predetermined temperature, then hold it for 1 hour. Reach the predetermined temperatures (300°C, 500°C, and 700°C) respectively. After cooling to room temperature, remove the obtained titanium dioxide and grind it. Mix it for 1 hour by ultrasonic centrifugation, then wash and filter with water. Put the filtered sample into a vacuum box at 70°C to dry for 2 hours to obtain the prepared sample, as shown in Fig. 2. To facilitate distinction and comparison, we uniformly name the prepared samples. The original sample is named TiO₂-P25, while the samples after 1-hour heating at 300°C, 500°C and 700°C are respectively named TiO₂-300°C, TiO₂-500°C, and TiO₂-700°C.

3.3. Membranes preparation

Membranes preparation process as demonstrated in Fig. 3, we weighed 10 mg of different pure TiO₂ samples, respectively, at 300°C for 1 hour, 500°C for 1 hour, and 700°C for 1 hour. Then, we took 10 ml of deionized water for each sample and mixed them together. We placed the mixture in an ultrasonic machine for 1 hour to dissolve. After that, we

removed the mixture and used a standard cellulose paper with a diameter of 3.8 cm. We performed filtration to produce 4 different water-evaporated TiO_2 membranes.

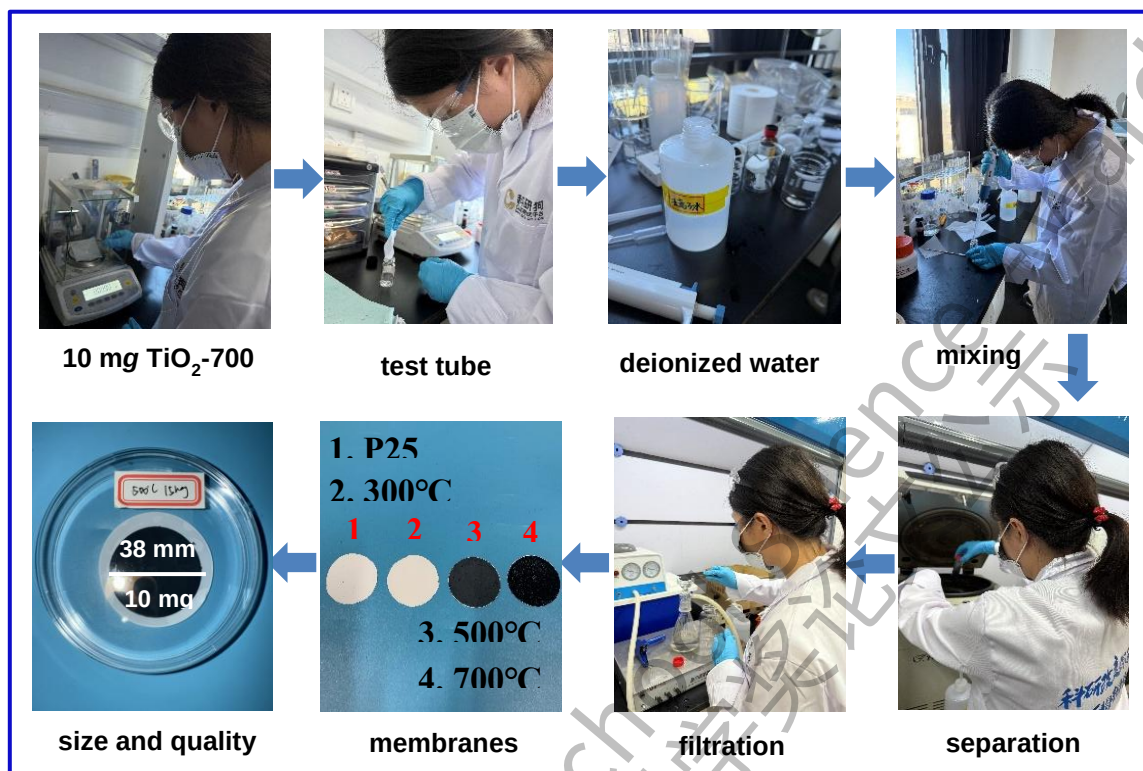


Fig. 3. Membrane preparation process.

3.4. Solar water evaporation experimental tests

The black TiO_2 powder were prepared by a NaBH_4 -assisted method. In brief, 2 g P25 powder were mixed with 0.75 g NaBH_4 powder, and then the mixed power was homogenized by grinding. Moreover, the powder mixture was transferred into tube furnace and purged with argon gas for 30 min. After that, the mixtures were calcined at 300 °C, 500 °C, and 700 °C under an argon gas for 1 h, respectively, and obtained black TiO_2 samples were named 300 °C, 500 °C and 700 °C. Finally, the black TiO_2 samples were washed with deionized water two times, respectively.

The efficiency of solar-driven water evaporation was quantified through the experimental setup presented in Fig. 4(a). Light source is a xenon lamp as a chamber optical system. Under the sun illumination, the mass reduction was continuously measured using a high precision analytical balance. As shown in Fig. 4(b), 10 mg TiO_2 samples (P25, 300 °C, 500 °C and 700 °C) were dissolved in 10 mL deionized water with ultrasound dispersion for 1 h at room temperature, and then deposited onto mixed cellulose ester (MCE) membranes

via vacuum filtration process to fabricate into evaporator, respectively. These membranes were cut into 38 mm diameter circle. Which were floated on water using a polyvinylidene foam insulating support and overlaid with an airlaid paper. Thermal conductivity of MCE membranes is $0.073 \text{ W m}^{-1} \text{ K}^{-1}$ as shown in Table S4. The membrane evaporator converts absorbed light into heat, leading to localized evaporation and steam generation. The variations in water surface temperature and the corresponding thermal distribution were captured using an infrared camera, as presented in Fig. 4(c). Absorbed light energy is rapidly converted into heat by the membrane evaporator, thereby inducing a rapid increase in its surface temperature and stabilize at approximately 51.3°C within just 10 min under three sun irradiations.

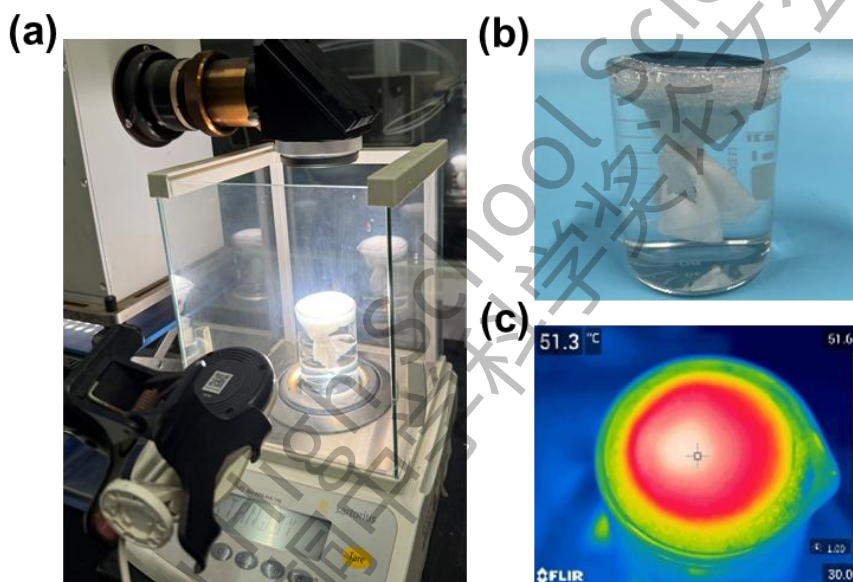


Fig. 4. Solar water evaporation experimental tests. (a) Testing system setup, (b) Modified TiO_2 membrane evaporators, (c) IR image of 700°C membrane surface.

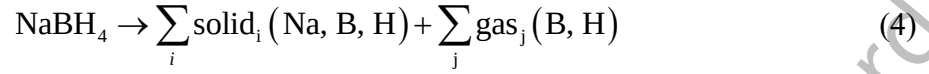
All tests were performed under controlled ambient conditions (26°C , 65% relative humidity). Following each test cycle, the samples were thoroughly dried at 50°C in a vacuum oven for subsequent reuse. Comprehensive details regarding the materials, characterization methods, and Supplementary Information provided evaporation testing procedure.

4. Experimental results and discussion

4.1. Doping mechanism of TiO_2

To reduce bandgap of original TiO_2 sample for increasing the visible light absorption,

NaBH₄ was employed as reduction agent, which can provide active hydrogen to convert Ti⁴⁺ into Ti³⁺ and form oxygen vacancies, as the primary origin of the Ti 3d bandgap states [31]. The thermal decomposition of NaBH₄ follows the below reaction:



where the solid residue and unreacted reactant can be easily removed by water.

4.2. Pearson correlation coefficient study

In this work, the interrelationships among features (BG, A, TC, sun, T, H) and targets (ER, EE) were evaluated using Pearson correlation coefficients, as presented in Fig. 5. Each matrix element represents the correlation coefficient between its respective row and column variables. This value falls in between -1 and 1, providing information on how the two variables correlate to each other. As shown in Fig. 5(a), the results show that a very high positive correlation (correlation coefficient of 0.92) between sun and ER, indicating that higher irradiance directly enhances solar energy capture. Mechanistically, increased photon flux elevates light-induced carrier generation efficiency in semiconductors, thus enhancing energy conversion. In contrast, no significant linear correlations were observed between the other features and the predicted variables of ER and EE, a key consideration in multivariate analysis.

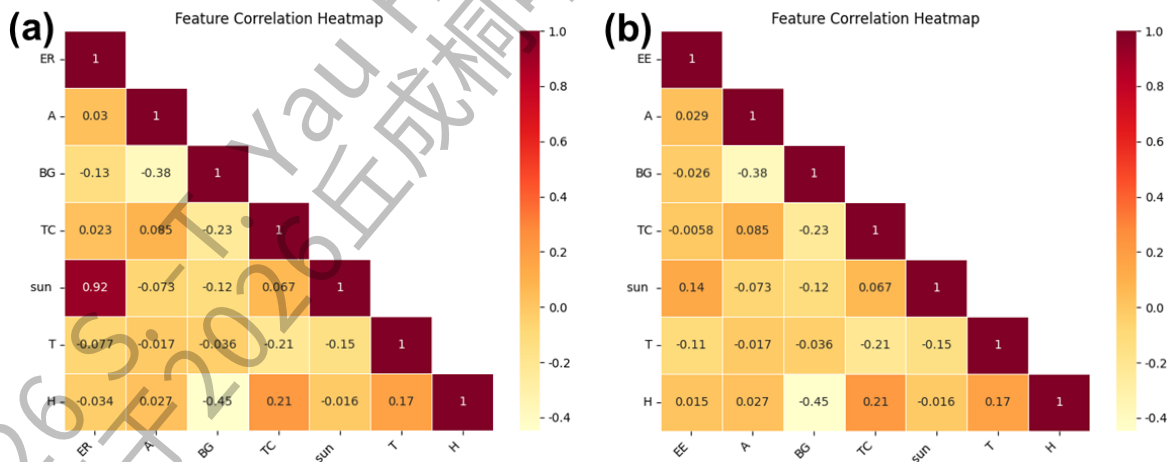


Fig. 5. Pearson correlation coefficient matrix between the six input features (A, BG, TC, sun, T, H) and (a) target variable (ER), (b) target variable (EE).

4.3. ER prediction

ER was employed as the primary performance metric to evaluate the evaporation

performance of SPMs. Model efficacy was quantified using RMSE and MAE on separately training sets and test sets, as shown in Table S5 and Table S6, with lower values indicating better performance. Comparative analysis indicates that the VR model as optimal, given its better error statistics.

Additionally, scatter plots illustrate the error by comparing the predicted versus actual ER for the four ML models, as shown in Fig. 6. Blue and red points denote the training and test data for ER, respectively. Obviously, these blue points and red points in Fig. 6(a) closely cluster around the best-fit line in the VR model. Furthermore, the prediction error (RMSE, MAE, R^2) of VR model are lower than other models as shown in Table S5, which indicates VR model has high accuracy. This advantage of VR model is due to integrate three distinct sets of ER prediction results of GBDT, SVM and XGBT models which are approximate and without overfitting, resulting in overperforms the individual models present as its estimators.

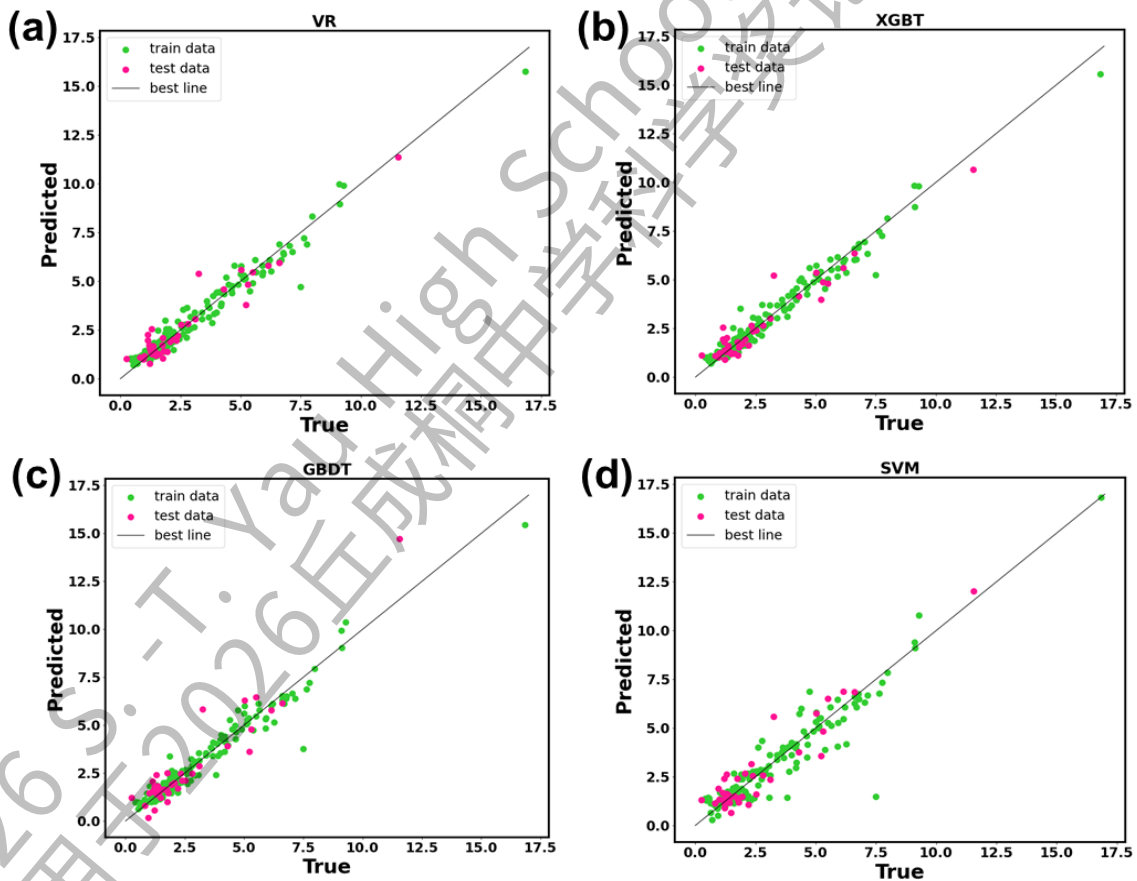


Fig. 6. ER prediction scatter plots for the four ML models: (a) VR, (b) XGBT, (c) GBDT, (d) SVM.

4.4. EE prediction

The EE is a critical factor in evaluating performance of SPMs on converting full-

spectrum solar energy into recoverable heat and depends on their specific properties, including bandgap, and light absorption capability. The EE prediction error of four ML models on the training sets and test sets are present in [Tables S7](#) and [Table S8](#). [Fig. 7](#). displays scatter plots comparing the predicted versus actual values for the four ML models. In these plots, training data for EE are shown in blue, while test data are denoted in red. Comparison of the predicted against actual values are illustrated via scatter plots in [Fig. 7](#), where blue and red points denote the training and test data for EE, respectively. [Fig.7\(a\)](#) indicates the VR model performs better due to it combining the predictions of multiple diverse estimators to smooth out errors and captures different aspects of the data. Therefore, VR model can achieve superior accuracy than single model on EE training data.

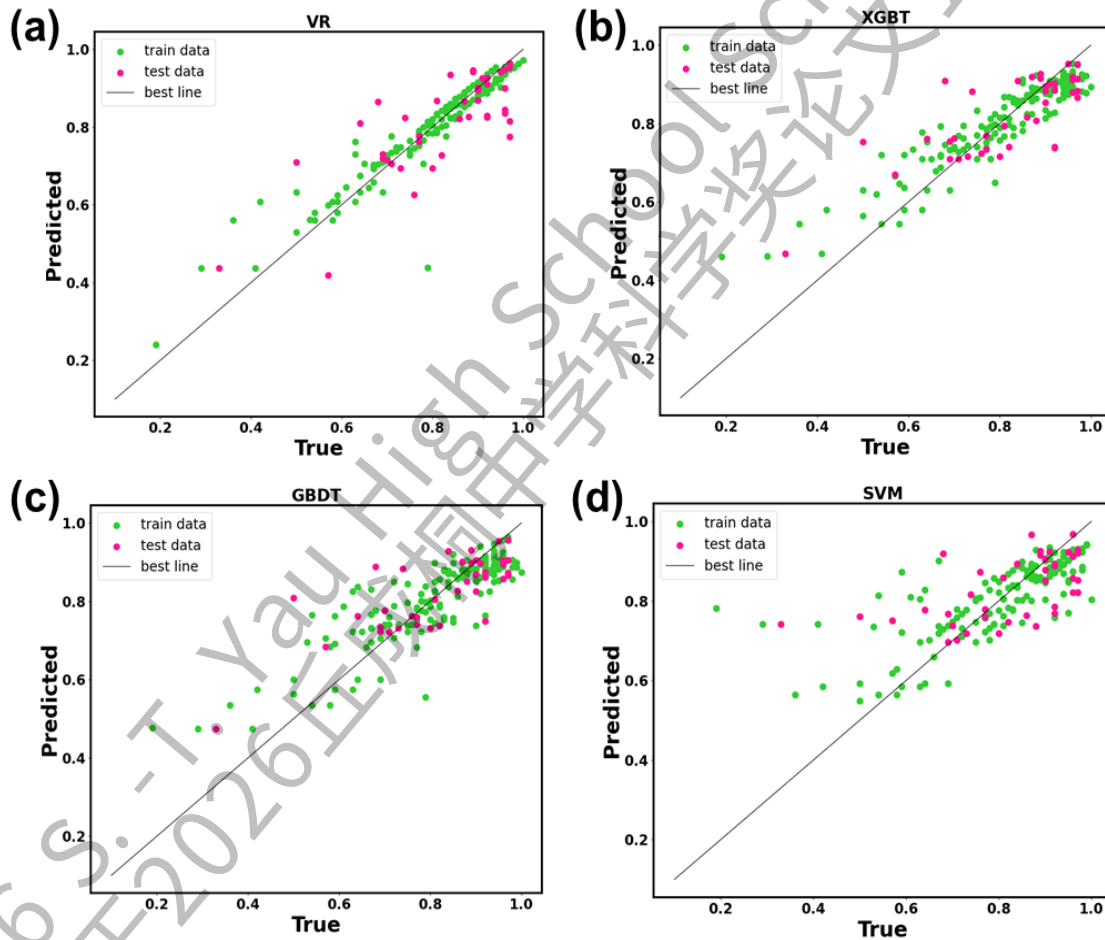


Fig. 7. EE prediction scatter plots for the four ML models: (a) VR, (b) XGBT, (c) GBDT, (d) SVM.

4.5. Feature importance and SHAP values analysis

Enhancing model interpretability is crucial for developing ML. SHAP values quantify the contribution of each feature to a specific prediction, and thus it can be used to analyze

which features are most important for the model and how they affect the outcome [32]. The SHAP plots shown in Fig. 8(a) highlight the impact of six features (BG, A, TC, sun, T, and H) on the model's output of ER. As shown in Fig. 8(a), among the six features, sun demonstrates a marked positive association with ER, indicating significant influence on ER prediction. When increasing sun light intensity, SPMs can achieve more solar energy absorption, resulting in improving water mass loss to increase ER. Similarly, TC is a system-level property dominated by the membrane, resulting in primary influencing ER. It has been reported that the interfacial heating is concentrated on the upper surface of the membrane evaporation with low thermal conductivity materials to prevent heat diffusion to the container, thereby improving ER [33]. Absorbance is essential for efficient photothermal conversion in SPMs, with higher absorbance leading to greater light harvesting and heat generation to improve ER [34]. BG reduction can decrease the intrinsic energy barrier for electron excitation from valence band to conduction band in SPMs, and prolong the activity under visible light irradiation. Ambient temperature and relative humidity which also directly impact ER because they affect energy available and vapor pressure, respectively.

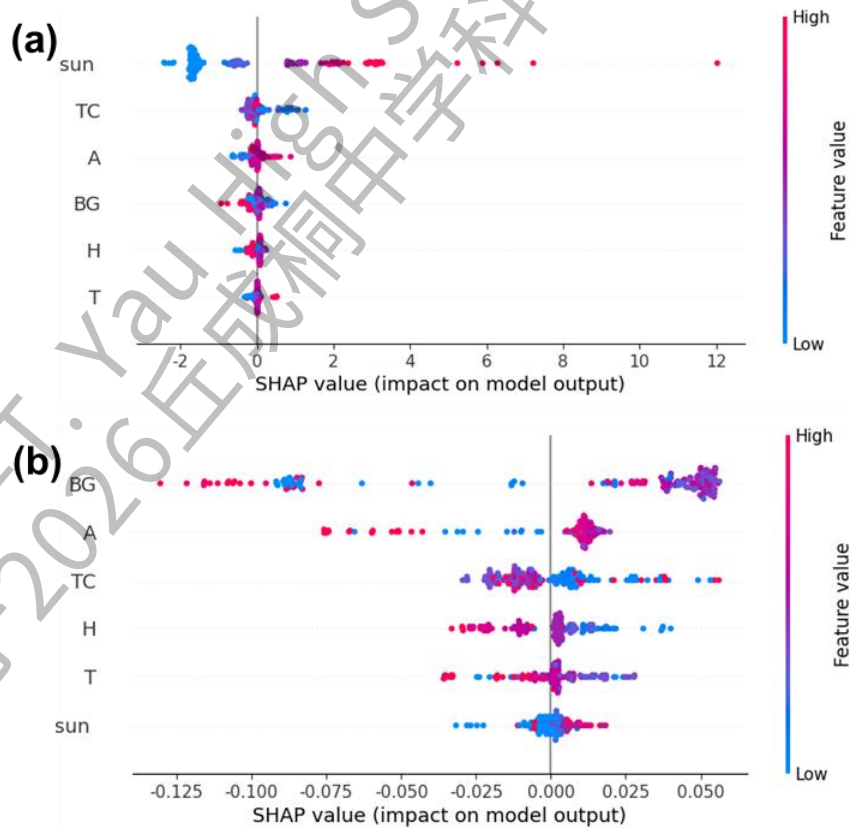


Fig. 8. Analyzing feature contribution with SHAP: (a) predicted ER using the VR model, (b) predicted EE using the VR model.

As shown in Fig. 8(b), the SHAP value for EE prediction indicates that no single feature exhibits a dominant influence, which is consistent with the model's difficulty in capturing the underlying physics of EE as shown in Fig. 4. Relatively speaking, BG and A have the most significant impacts on the SHAP value among all features, providing reliable physical interpretability. TC shows smaller influences on SHAP value of EE. This lack of clear, dominant feature contributions is consistent with the lower R^2 for EE prediction, suggesting that EE is governed by more complex, multi-factor interactions that are not yet fully captured by the current dataset. Larger and more diverse dataset may improve both the accuracy and the interpretability of EE prediction. To further validate the ML prediction, in the following section the solar water evaporation tests of SPMs was measured.

4.6. Experimental validation

To validate the predictive efficacy of the ML models, we employed black TiO_2 as a photothermal material for performance evaluation in interfacial evaporation. By using P25 as the starting material and NaBH_4 as the reducing agent, we thoroughly mixed them and then placed the mixture in a tube furnace for reaction. As depicted in Fig. S1, with the increase in reaction temperature, the color of the samples gradually changed from white to black.

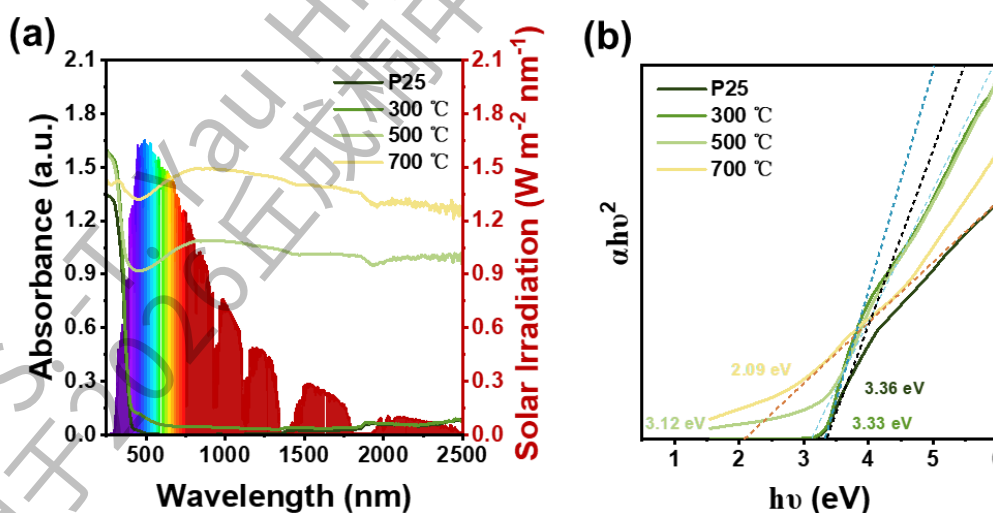


Fig. 9. The four different samples properties (a) absorbance, (b) bandgap.

As shown in Fig. 9(a), the UV-Vis-NIR absorbance spectra of the gray-black and black samples clearly demonstrate their excellent light absorption capabilities, whereas the white raw material and the sample calcined at 300°C exhibit extremely poor light absorption. As

the reaction temperature rises, the intensity of light absorption gradually increases, which is consistent with the color change of the samples. The bandgap energy of the samples is calculated based on their absorption spectra. As shown in Fig. 9(b), the bandgap of P25 is 3.36 eV. With the increase in reaction temperature, the bandgap gradually decreases, which is consistent with the phenomenon of gradually enhanced light absorption intensity in the absorption spectra. The gradual decrease in bandgap resulted from the creation of oxygen vacancies induced by reduction, which causes the transformation of Ti^{4+} to Ti^{3+} [35, 36].

To delve into the influence mechanism of valence states and oxygen vacancies on the bandgap, four samples were characterized by X-ray photoelectron spectroscopy (XPS) [37]. As shown in Fig. S2a, the XPS survey indicates that the sample is composed solely of Ti, O and C. As shown in Fig. 10(a), Ti 2p spectral analysis was carried out on P25 and TiO_2 calcined at different reaction temperatures. The analysis results reveal that within the energy range of 455 to 470 eV, XPS peaks for Ti 2p_{3/2} and Ti 2p_{1/2} can be observed at binding energies of 458.88 eV and 464.68 eV, respectively. These peaks are typical characteristic peaks of Ti^{4+} -O bonds. For the calcined TiO_2 , the XPS peak position of Ti 2p_{3/2} shifts from the initial 458.88 eV to 458.68 eV, 458.58 eV, and 458.58 eV successively. At the same time, the XPS peak position of Ti 2p_{1/2} also shifts from 464.68 eV to 464.48 eV, 464.18 eV, and 464.18 eV. In combination with Figs. S2(b)-(e), such shifts are observable in peak positions indicate the formation of Ti^{3+} [38]. Specifically, the peak at 458.88 eV can be decomposed into two Gaussian peaks, which are attributed to Ti^{3+} and Ti^{4+} , respectively. Through peak-fitting calculation, the $\text{Ti}^{4+}/\text{Ti}^{3+}$ peak area ratios for the samples reduced at different temperatures were determined to be 1.86, 1.56, 1.25, and 1.18, respectively. This ratio shows a gradual decreasing trend with increasing calcination temperature, indicating that the reduction degree of Ti^{4+} to Ti^{3+} is progressively enhanced.

Fig. 10(b) presents the O 1s XPS spectrum. From the spectrum, it can be observed that the XPS peak of Ti-O shows a certain degree of shift. In addition, there is a broad O 1s peak at 532.18 eV, which is attributed to the Ti-OH bond. This result confirms the presence of hydroxyl groups on the surface of black TiO_2 after the reduction reaction. To confirm the presence of hydroxyl groups, further analysis was conducted on the O 1s XPS spectra of each sample, with the results shown in Figs. 10(c)-(f). The analysis revealed the emergence

of two peaks at 530.18 eV and 531.78 eV. The peak at 530.18 eV corresponds to lattice oxygen ($\text{Ti}^{4+}\text{-O}$), while the other peak at 531.78 eV corresponds to oxygen vacancies ($\text{Ti}^{3+}\text{-O}$). Additionally, a new peak appeared at 532.68 eV in the calcined TiO_2 , which corresponds to adsorbed oxygen or surface hydroxyl groups (Ti-OH). These results of XPS analysis indicate abundant oxygen vacancies and Ti^{3+} into surface of TiO_2 sample with NaBH_4 reduction under high temperature.

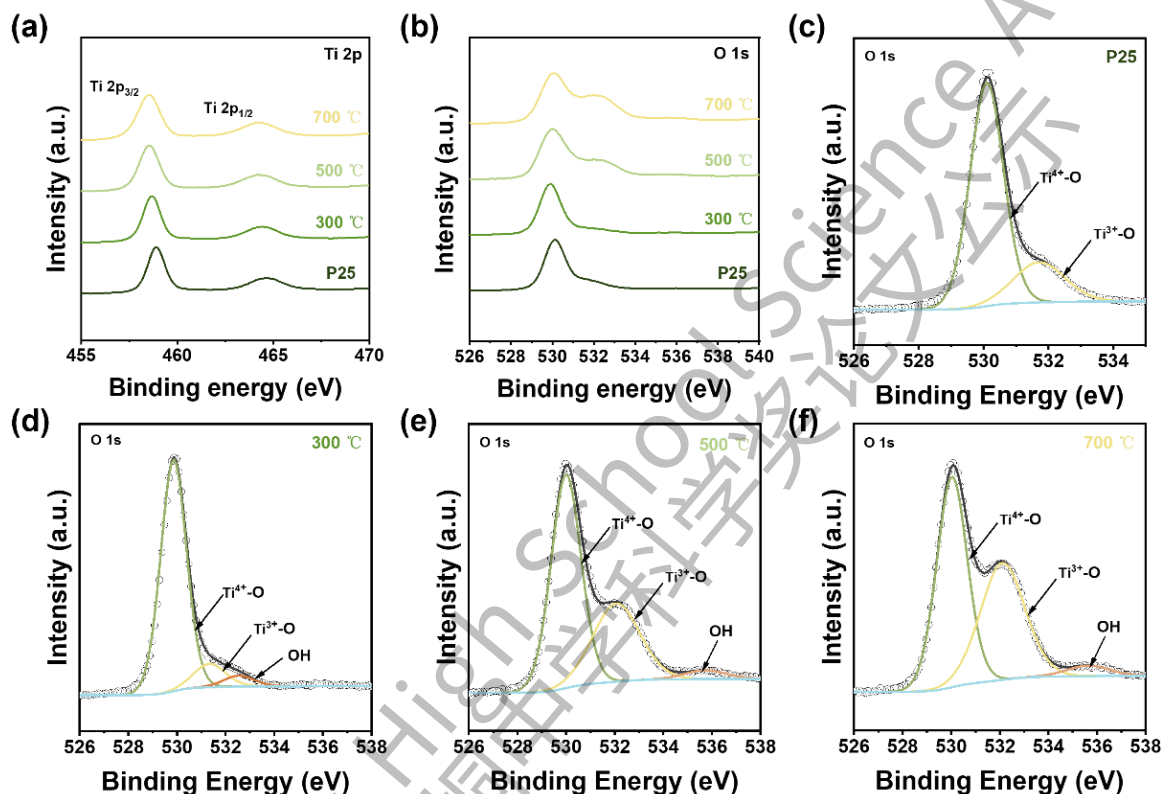


Fig. 10. (a) Ti 2p XPS spectra and (b) O 1s XPS spectra of the four TiO_2 samples. High-resolution O 1s spectra of (c) P25, (d) 300 °C, (e) 500 °C and (f) 700 °C.

To further investigate the effect of calcination temperature on the material structure, XRD analysis was performed on the samples reduced at different temperatures, and the results are shown in Fig. S3. For the pristine P25, the diffraction peaks at 25.3° , 38.0° , 48.1° , 53.9° , 54.8° , and 62.7° can be assigned to the (101), (004), (200), (105), (211), and (204) crystal planes of anatase-phase TiO_2 (JCPDS No. 21-1272), respectively. When the calcination temperature was 300 °C, the diffraction peak positions of the sample were essentially consistent with those of P25, indicating that the moderate-temperature reduction did not induce obvious phase changes. As the temperature increased to 500 °C, new diffraction peaks appeared at 27.3° , 36.2° , and 54.0° in addition to the anatase peaks. These

peaks correspond to the (110), (101), and (211) crystal planes of rutile-phase TiO_2 (JCPDS No. 21-1276), suggesting that the anatase-to-rutile transformation had begun at this temperature. When the calcination temperature was further raised to 700 °C, the XRD pattern changed dramatically: the characteristic peaks of anatase almost completely disappeared and were replaced by numerous diffraction peaks attributable to Ti_3O_5 (JCPDS No. 76-1066) along with a small amount of TiO (JCPDS No. 72-0020) [39]. This indicates that TiO_2 was deeply reduced to low-valence titanium oxides, and the anatase phase was completely converted to the rutile phase.

SEM observations were performed on the samples treated at different temperatures, and the results are shown in Fig. 11. As the calcination temperature increased, the pristine P25 nanoparticles gradually agglomerated and grew. When the temperature reached 700 °C, obvious rod-like crystals appeared in the sample. This morphological evolution further confirms that when the calcination temperature exceeds 500 °C, a phase transformation occurs within the material, with the crystal structure gradually converting from the anatase phase to the rutile phase. Moreover, under the high-temperature reduction conditions at 700 °C, low-valence titanium oxides, namely Ti_3O_5 and a small amount of TiO , were further generated.

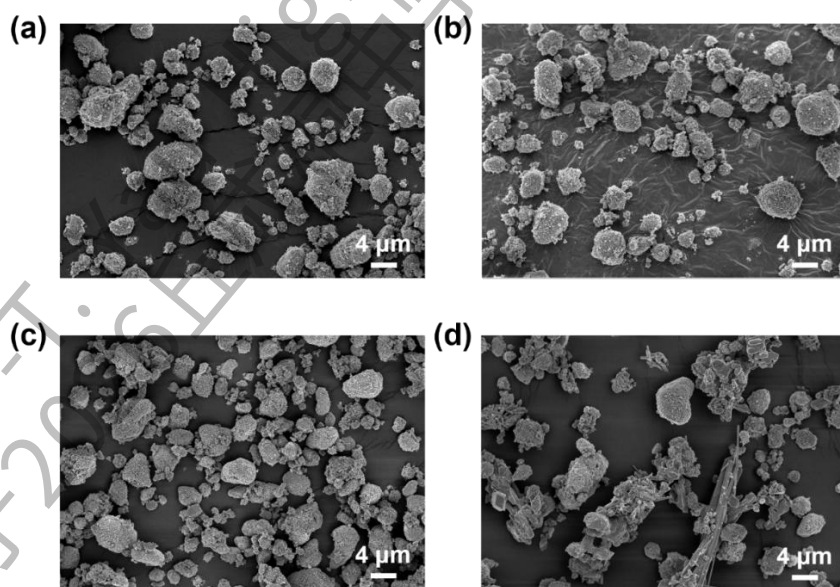


Fig. 11. SEM images of (a) P25, (b) 300 °C, (c) 500 °C and (d) 700 °C.

To further verify the phase transformation behavior occurring during the high-temperature reduction process, TEM characterization was performed on samples treated at different temperatures, with the results shown in Fig. 12. As the calcination temperature

increased, the pristine spherical particles gradually agglomerated. When the temperature reached 500 °C, amorphous flocculent edges appeared in some regions, and the particle boundaries became blurred, indicating the onset of lattice distortion, which is consistent with the initial stage of the anatase-to-rutile phase transition observed in the XRD results. At this stage, the lattice fringe spacing measured from High-resolution (HR-TEM) images is about 0.36 nm, which corresponds to the (101) plane of the anatase phase. When the temperature was further increased to 700 °C, the sample morphology changed significantly, with the spherical particles completely transforming into rod-like crystals. HR-TEM images revealed lattice fringes with a spacing of approximately 0.46 nm in the 700 °C sample, which can be indexed to the (110) crystal plane of the low-valence titanium oxide Ti_3O_5 , confirming that the phase had been completely reconstructed into a new crystalline phase with good crystallinity. In addition, lattice fringes corresponding to the rutile phase were also observed in the 700 °C sample, with a spacing of approximately 0.34 nm, assigned to the (110) crystal plane of rutile (Fig.S4). Notably, this spacing is larger than the theoretical value (0.3247 nm) for the standard rutile (110) plane, which can be attributed to lattice distortion and expansion caused by a high concentration of oxygen vacancies. The TEM observations are in good agreement with the XRD analysis, confirming that upon calcination at 700 °C, a phase transition indeed occurred and low-valence titanium oxides were generated.

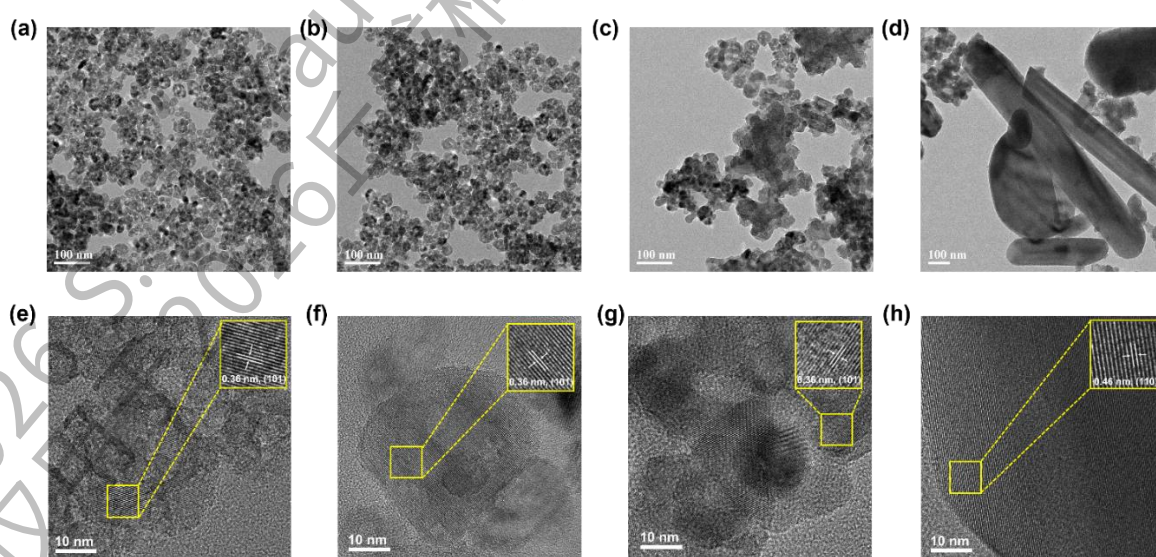


Fig. 12. TEM images of (a) P25, (b) 300 °C, (c) 500 °C and (d) 700 °C. HR-TEM images of (e) P25, (f) 300 °C, (g) 500 °C and (h) 700 °C.

Fig. 13(a) shows the Raman spectra of the four samples. The spectra of P25 and the sample reduced at 300 °C are consistent, exhibiting characteristic peaks of anatase TiO₂ at 147 cm⁻¹, 401 cm⁻¹, 516 cm⁻¹, and 641 cm⁻¹, indicating that low-temperature reduction did not induce phase transformation. The 500 °C sample still retains the anatase peak shape, but the strongest peak (144 cm⁻¹) blue-shifts to 154 cm⁻¹ with a slight increase in peak width, and the intensities of the other peaks decrease, suggesting that the sample is in the initial stage of the anatase-to-rutile phase transition, accompanied by lattice distortion caused by a small number of oxygen vacancies. For the 700 °C sample, the characteristic anatase peaks completely disappear and are replaced by a set of broadened new peak groups, indicating that at this temperature, TiO₂ is deeply reduced and the phase reconstructs into low-valence titanium oxides such as Ti₃O₅, while the high concentration of oxygen vacancies, Ti³⁺ and lattice disorder lead to significant broadening of the Raman peaks.

Fig. 13(b) shows the EPR spectra of pristine P25 and NaBH₄ reduced TiO₂ samples at different temperatures. No obvious paramagnetic signal is detected for pristine P25 or the sample reduced at 300 °C. As shown in the magnified view in Figure 11b, the weak sharp peaks observed in both samples are attributable to instrument baseline noise or paramagnetic impurities, rather than characteristic signals of oxygen vacancies or Ti³⁺. When the reduction temperature is increased to 500 °C, the EPR signal intensifies significantly, with the peak at $g \approx 2.002$ assigned to oxygen vacancies and the peak at $g \approx 1.968$ corresponding to Ti³⁺ species. Upon further increasing the temperature to 700 °C, the signal becomes substantially stronger, accompanied by pronounced broadening and an asymmetric line shape. These features are closely related to the deep reduction of the sample, the generation of abundant Ti³⁺ and oxygen vacancies, and the strong magnetic interactions arising from structural reconstruction into low-valence titanium oxides (Ti₃O₅), which is fully consistent with the Raman and XRD results. The significant increase in EPR linewidth further indicates a markedly higher local density of unpaired electrons and enhanced spin-spin coupling, confirming that treatment at 700 °C introduces high concentrations of oxygen vacancies and Ti³⁺ defects into TiO₂.

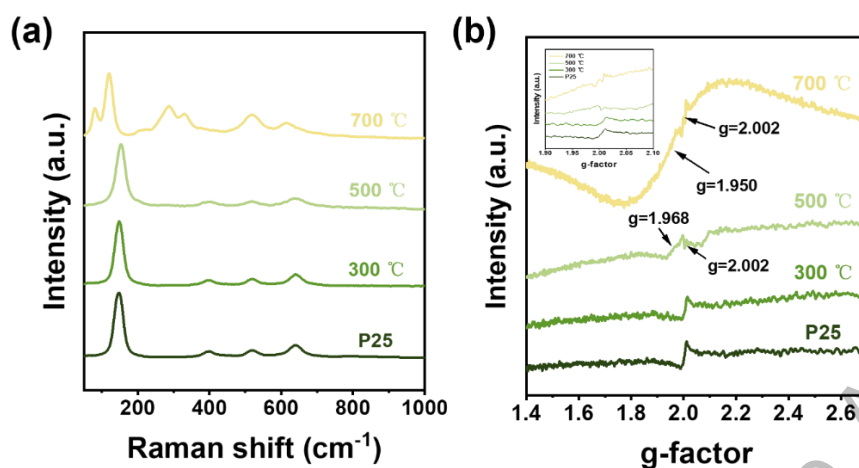


Fig. 13. (a) Raman spectroscopy of P25, 300 °C, 500 °C and 700 °C. (b) EPR spectra of P25, 300 °C, 500 °C and 700 °C, the inset shows a magnified view of the region within $g = 1.90$ to 2.10 .

The surface temperature of TiO_2 -700 membrane under 1 sun as shown in Fig. 14 under 1 sun for 0 min, 60 min, 70 min, 80 min and 90 min, respectively.

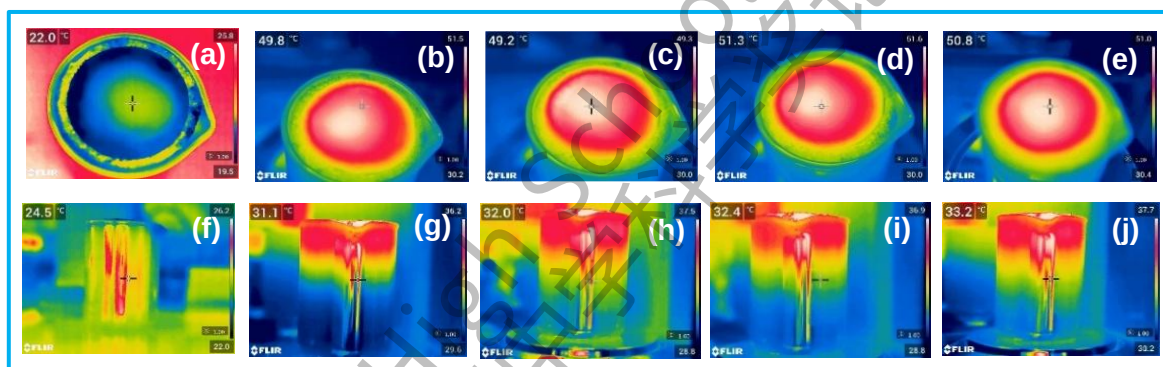


Fig. 14. Infrared radiation images of TiO_2 -700 membrane surface under 1 sun (a)-(f) 0 min, (b)-(g) 60 min, (c)-(h) 70 min, (d)-(i) 80 min, (e)-(j) 90 min..

The evaporation performance of four different samples was evaluated to validate the ML prediction. To quantify solar water evaporation of the four different samples, we recorded the mass change versus time under different light densities of one sun, two sun, three sun, and four sun, respectively. Fig. 15(a)-(b) show the variations in mass and surface temperature over time for pure water and four different TiO_2 membranes under one sun.

Fig. 15(c) compares the measured ER and EE data for four different samples under one sun. The ER values of pure water, P25, 300 °C, 500 °C and 700 °C are 0.22, 0.41, 0.45, 0.86 and $1.12 \text{ kg m}^{-2} \text{ h}^{-1}$, respectively, with corresponding EE values of 10.9%, 21.8%, 23.2%, 49.7% and 67.8%. Comparing the 500 °C and 700 °C with 300 °C, ER and EE achieves a

significantly improvement, primarily attributed to the bandgap of samples decreases with increasing of calcined temperature, and it can absorb light across the full spectrum, with a stronger light absorption capacity. Overall, for the four different samples, the consistency between the experimental correlation of these factors and the ML results confirm BG plays a pivotal role in determining the EE.

To further verify the influence of sun on evaporation performance, we tested ER and EE of the membrane with 700 °C under different irradiance levels. Fig. 15(d) displays water mass loss of the 700 °C sample exhibits a continuous increase with rising sun. Fig. 15(e) shows the surface temperature fast increases at initial time, eventually reaching saturation and keep stability with prolonging the irradiation time. Fig. 15(f) presents the ER values of 700 °C evaporator are 1.12, 1.62, 1.89 and 2.28 kg m⁻² h⁻¹ under one sun, two sun, three sun, and four sun, respectively, with corresponding EE values of 67.8%, 51.2%, 40.7% and 37.7%. Which manifests the four sun have significant enhancement of ER for 700 °C evaporator. Obviously, EE exhibits a gradual decline with increasing sun. This phenomenon is primarily ascribed to the inherently limited light absorption of the material calcined at 700 °C, which restricts the maximum surface temperature increment. Therefore, the evaporation rate fails to grow linearly with increasing light intensity, demonstrating a typical nonlinear behavior. In addition, elevated heat loss via thermal radiation and convection occurs under high illumination, accompanied by severe interfacial overheating. These combined effects deteriorate the photothermal conversion performance and gradually reduce the evaporation efficiency [40-42]. Additionally, as described in Table S9, the 700 °C demonstrated a higher water evaporation performance superior to those of previously reported in the literature. This indicates that narrower BG promotes the adsorption of sunlight. Collectively, research results verify the distinct yet significant influences of material feature A on the ER and BG on the EE of SPMs. It confirming the SHAP analysis and establishing the validity of material performance forecasts by the ML methodologies employed.

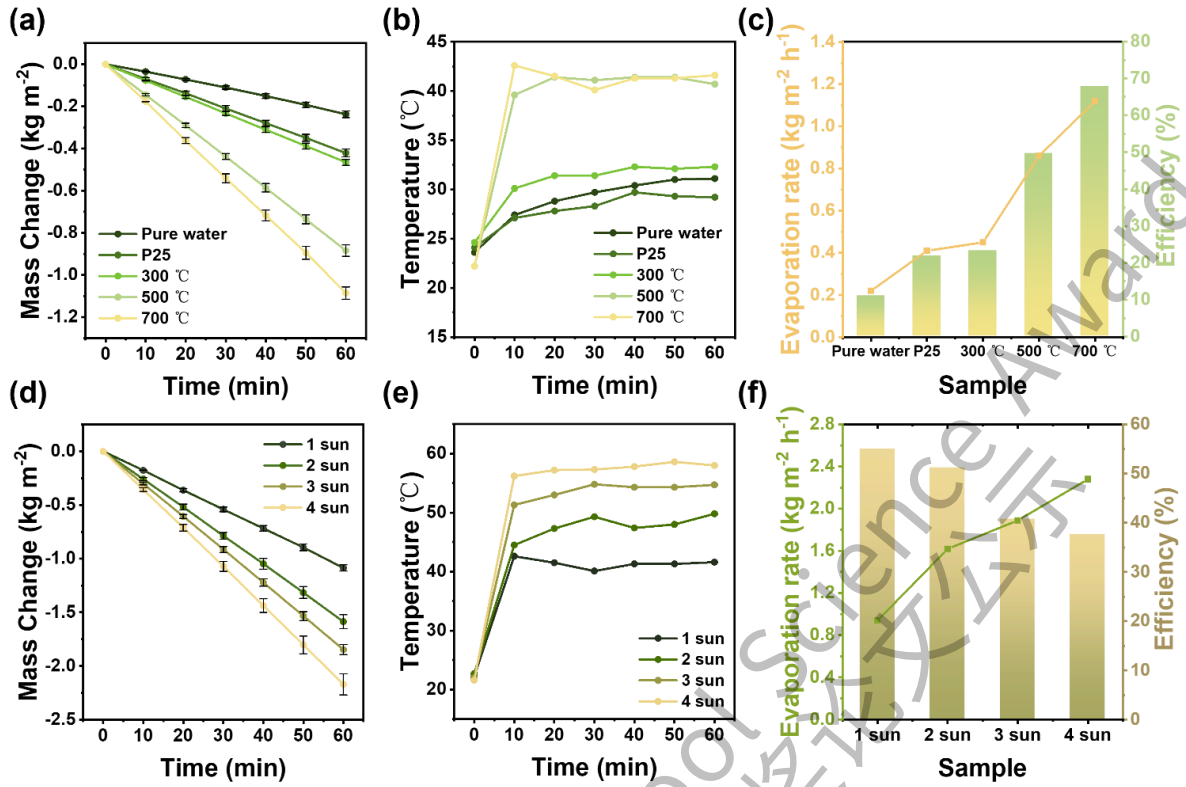


Fig. 15. (a) Water mass change for Pure water and four samples (1 sun). (b) Surface temperature profiles for Pure water and four membranes (1 sun). (c) Comparative analysis of ER and EE for Pure water and four membranes (1 sun). (d) Water mass change with 700 °C sample under various irradiance levels. (e) Changes in surface temperature of 700 °C sample under various irradiance levels. (f) The ER and EE values of 700 °C sample under different simulated intensities of sunlight.

5. Conclusions

In this study, ML models VR, XGBT, GBDT and SVM were employed to accurately forecast the evaporation rate and evaporation efficiency of SPMs. As the limited sample size in the current dataset ($n=196$), we applied Bayesian optimization and treated cross-validation as the objective function, minimizing the number of cross-validation evaluations, reducing overfitting to a single validation split. Moreover, the correlations between critical features and evaporation performance were investigated. The VR model achieves the highest R^2 for ER on test sets ($R^2=0.9337$). However, the R^2 of EE prediction is only moderate ($R^2=0.724$), suggests that the current dataset may not fully capture the complex factors governing evaporation, expanding the dataset samples to enhance model performance in the future work. SHAP value analysis identified sun and TC as the most significant factors for ER, while BG was the most influential for EE. The results highlight

BG as critical parameters deserving focused attention in the future study and design of novel SPMs. Four modified TiO_2 samples prepared by utilizing hydrogen-induced reduction reaction to reduce the band-gap of TiO_2 under different high temperature. To evaluate the prediction performance of the ML models, solar evaporation tests of four different samples were tested under different light density of one sun, two sun, three sun, and four sun, respectively. The characterization and evaporation experimental results show that when the calcination temperature reaches $700\text{ }^\circ\text{C}$, a phase transition occurs within the P25 sample, generating low-valence titanium oxides, with Ti^{4+} reduced to Ti^{3+} accompanied by the formation of oxygen vacancies. These changes lead to bandgap narrowing and enhanced light absorption, thereby effectively improving the water evaporation capability. This result further validates the reliability and applicability of the machine learning (ML) method. This study highlights the promise of ML approaches in innovating design strategies for maximizing energy conversion, and optimizing material features and operational conditions in photothermal evaporation technologies. Overall, this work presented a ML approach to support the development of SPMs, establishing a predictive framework between material characteristics and evaporation performance for material design.

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Appendix A: Supplementary Information (data collection and testing)

Table S1 Collected dataset from previous works.

EE %	ER kg/m ² h	BG eV	A %	TC W/mK	Sun kw/m ²	T °C	H %	Ref
81	1.4	1.3	0.959	0.02	1	23	90	[1]
87	2.8	1.3	0.959	0.02	2	23	90	
96	4.3	1.3	0.959	0.02	3	23	90	
96	3.68	1.3	0.838	0.02	3	23	90	
96	3.05	1.3	0.765	0.02	3	23	90	
80.5	1.3	1.2	0.93	0.06	1	26	75	[2]
83.5	2.8	1.2	0.93	0.06	2	26	75	
91	4.18	1.2	0.93	0.06	3	26	75	
91	5.65	1.2	0.93	0.06	4	26	75	
91	7.03	1.2	0.93	0.06	5	26	75	
67.7	1.15	0.45	0.99	0.525	1	23	55	[3]
70	2.2	0.45	0.99	0.525	2	23	55	
76.5	3.6	0.45	0.99	0.525	3	23	55	
82	5.1	0.45	0.99	0.525	4	23	55	
86.2	6.6	0.45	0.99	0.525	5	23	55	
84.5	1.46	/	0.9225	0.02	1	25	50	[4]
88.78	6.8	/	0.9225	0.02	5	25	50	
88	1.32	/	0.95	0.05	1	20	58	[5]
50.3	0.8	/	0.913	/	1	21	55	[6]
76.7	1.229	1.82	0.97	/	0.5	25	50	[7]
85.6	1.255	1.82	0.97	/	1	25	50	
86.5	2.52	1.82	0.97	/	2	25	50	
63	0.9	0.3	0.95	0.48	1	23	55	[8]
65	1.85	0.3	0.95	0.48	2	23	55	
67	3.1	0.3	0.95	0.48	3	23	55	
70	4.5	0.3	0.95	0.48	4	23	55	
71	5.6	0.3	0.95	0.48	5	23	55	
68.6	1.1	0.3	0.95	0.48	1	23	55	
71	2.25	0.3	0.95	0.48	2	23	55	
73	3.4	0.3	0.95	0.48	3	23	55	
75	4.9	0.3	0.95	0.48	4	23	55	
76.5	6.2	0.3	0.95	0.48	5	23	55	
75	1.35	0.3	0.95	0.48	1	23	55	
78	2.75	0.3	0.95	0.48	2	23	55	
80	4	0.3	0.95	0.48	3	23	55	
82	5.3	0.3	0.95	0.48	4	23	55	
83	6.7	0.3	0.95	0.48	5	23	55	
70.8	0.63	/	0.94	0.32	0.6	25	60	[9]
76.9	0.92	/	0.94	0.32	0.8	25	60	
81.4	1.22	/	0.94	0.32	1	25	60	

86.2	1.56	/	0.94	0.32	1.2	25	60	
53.7	0.45	/	0.868	/	1	25	50	[10]
85.8	1.44	/	0.94	/	1	25	50	
90	2.56	/	0.94	/	2	25	50	
90	3.85	/	0.94	/	3	25	50	
88	5	/	0.94	/	4	25	50	
92	1.48	3.2	0.944	/	1	21	50	[11]
92	1.48	3.2	0.944	/	1	21	50	[12]
90.4	1.43	2	0.9	/	1	25	50	[13]
91.7	2.91	2	0.9	/	2	25	50	
93.3	5.93	2	0.9	/	4	25	50	
68.6	1.09	2	0.85	/	1	25	50	[14]
75.1	2.38	2	0.85	/	2	25	50	
81.2	5.15	2	0.85	/	4	25	50	
63.9	1.01	2	0.93	0.067	1	26	/	[15]
94.9	1.63	/	0.95	0.04	1	24	25	[16]
88.4	4.04	/	0.95	0.04	3	24	25	
97.5	7.16	/	0.95	0.04	5	24	25	
86.6	1.657	/	0.97	0.06	1	/	/	[17]
75.7	0.81	1	0.96	/	0.76	25	/	[18]
59.3	0.64	1	0.96	/	0.76	25	/	
81.4	6.15	1	0.96	/	5.35	25	/	
62.6	4.73	1	0.96	/	5.35	25	/	
94	1.95	/	0.98	0.03	1	/	/	[19]
91.5	6.6	/	0.95	0.06	5	25	/	[20]
80	1.27	/	0.9615	0.037	1	20	60	[21]
61	1.95	/	0.9615	0.037	2	20	60	
67	5.9	/	0.9615	0.037	5.5	20	60	
92.1	1.32	0.1	0.925	0.02	1	20	55	[22]
95.3	5.03	0.1	0.925	0.02	5	20	55	
77.8	1.24	/	0.837	/	1	25	40	[23]
84.45	1.23	/	0.9627	/	1	25	55	[24]
94	1.45	/	0.9	0.243	1	26	50	[25]
67	1.07	/	0.903	/	1	25	45	[26]
86.9	1.28	2.5	0.906	0.02	1	/	/	[27]
92.4	1.81	/	/	0.035	1	20	50	[28]
/	4.63	/	/	0.035	3	20	50	
91.5	1.337	1.7	/	0.03	1	27	50	[29]
93.5	3.99	1.7	/	0.03	3	27	50	
94.1	6.78	1.7	/	0.03	5	27	50	
77.1	1.2	2.7	0.99	/	1	22	40	[30]
78.5	1.25	2.43	0.879	/	1	22	40	[31]
/	0.69	3.11	0.634	/	1	22	40	

93.8	1.22	1.7	/	0.03	1	/	/	[32]
89.5	1.86	2.76	0.948	0.145	1	28	30	[33]
72.5	4.61	2.76	0.948	0.145	3	28	30	
69.4	7.75	2.76	0.948	0.145	5	28	30	
87.5	3.25	2.4	0.657	/	5	/	/	[34]
93.8	4.4	1.1	0.834	/	5	/	/	
77.5	1.24	/	0.902	0.3	1	23	55	[35]
73.7	0.53	/	0.895	/	1	/	/	[36]
69.1	/	/	0.9	0.04	1	25	55	[37]
57.3	1	2.68	0.95	/	1	25	30	[38]
74.5	1.3	2.35	0.95	/	1	25	30	
83.1	1.45	2.35	0.99	/	1	25	30	
79.9	1.39	2.35	0.99	/	1	25	30	
81.6	1.86	0.7	/	0.08	1	24	55	[39]
97.7	7.5	0.7	/	0.08	1	24	55	
80.4	/	1.79	0.95	0.069	0.5	32	45	[40]
94.5	3.12	1.79	0.95	0.069	1	32	45	
89.2	4.2	1.79	0.95	0.069	1.5	32	45	
88.3	5.38	1.79	0.95	0.069	2	32	45	
95.9	2.2	/	0.98	0.77	1	25	60	[41]
95.6	2.09	/	/	/	1	25	25	[42]
87	2.2	/	0.93	0.39	1	/	/	[43]
91.2	1.57	/	0.97	0.55	1	24	25	[44]
94.5	1.5	/	0.98	/	1	25	60	[45]
91	1.09	2	0.95	/	1	25	40	[46]
91	2.38	2	0.95	/	2	25	40	
91	5.15	2	0.95	/	4	25	40	
98.5	1.79	/	0.983	0.71	1	25	60	[47]
85.1	1.355	/	0.94	0.06	1	25	31	[48]
91.4	2.911	/	0.94	0.06	2	25	31	
97.5	4.659	/	0.94	0.06	3	25	31	
96.6	6.152	/	0.94	0.06	4	25	31	
95.9	7.643	/	0.94	0.06	5	25	31	
67	0.95	1.27	/	/	1	26	23	[49]
79	4.31	1.27	/	/	4	26	23	
89.9	9.29	1.27	/	/	8	26	23	
97.1	1.546	0.85	/	/	1	/	/	[50]
90	/	0.85	/	/	7.6	/	/	
88	/	0.85	/	/	9.84	/	/	
90	/	0.85	/	/	14.5	/	/	
99	1.43	1.54	/	0.043	1	/	/	[51]
77.3	1.71	/	0.65	0.04	1.5	27	/	[52]
66	2.33	/	0.65	0.04	2.7	27	/	

58	2.61	/	0.65	0.04	3.4	27	/	
91.5	1.92	/	/	0.04	1	27	/	[53]
53	2.45	2.65	/	/	1	25	50	[54]
67	3.07	2.65	/	/	1	25	50	
81.4	3.81	2.65	/	/	1	25	50	
18.5	0.259	2.6	0.9	/	1	25	/	[55]
28.7	0.41	2.6	0.99	/	1	25	/	
33	0.48	2.6	0.99	/	1	25	/	
41.3	0.578	2.6	0.99	/	1	25	/	
78.7	1.101	2.6	0.99	/	1	25	/	
82.5	1.4	2.35	0.94	0.4	1	15	12	[56]
87	2.59	2.35	0.94	0.4	2	15	12	
94.5	5.51	2.35	0.94	0.4	4	15	12	
92	7.98	2.35	0.94	0.4	6	15	12	
89.5	9.12	2.35	0.94	0.4	7	15	12	
57.4	2.77	3.2	/	0.067	5	14	/	[57]
/	1.56	3.2	0.92	0.6	1	24	50	[58]
93.4	1.68	3.2	0.92	0.6	1.5	24	50	
64.1	0.94	3.2	/	0.3	1	20	/	[59]
89.4	1.77	/	0.968	0.1	1	/	/	[60]
91.3	1.96	/	0.99	0.1	1	/	/	
87	1.89	/	0.955	0.246	1	26	50	[61]
93	3.9	/	0.955	0.246	2	26	50	
98	5.9	/	0.955	0.246	3	26	50	
94.4	2.08	1.3	0.962	/	1	/	50	[62]
94.4	2.2	1.3	0.978	/	1	/	50	
100	1.61	/	0.94	0.432	1	25	50	[63]
85.3	1.36	1.7	0.97	0.35	1	27.3	50	[64]
84	2.7	1.7	0.97	0.35	2	27.3	50	
87	4.2	1.7	0.97	0.35	3	27.3	50	
83	5.2	1.7	0.97	0.35	4	27.3	50	
84	6.6	1.7	0.97	0.35	5	27.3	50	
95.5	2.53	/	/	/	1	23	40	[65]
88.7	2.07	1.89	0.92	/	1	25	50	[66]
93.5	1.01	1	0.969	0.02	0.5	25	40	[67]
95.3	2.06	1	0.969	0.02	1	25	40	
95.9	3.11	1	0.969	0.02	1.5	25	40	
96.5	4.17	1	0.969	0.02	2	25	40	
96.6	5.22	1	0.969	0.02	2.5	25	40	
96.7	6.27	1	0.969	0.02	3	25	40	
59	1.25	2.8	0.93	/	1	25	40	[68]
72.2	1.42	2.67	0.965	/	1	25	/	
67.5	1.31	1.19	0.96	/	1	25	/	
91	1.76	/	/	/	1	25	15	[69]

/	1.4	3	0.95	0.04	1	/	/	[70]
84.8	1.37	3	0.9	0.04	1	/	/	
91.5	3.7	3	0.9	0.04	3	/	/	
92	5.5	3	0.9	0.04	5	/	/	
72	1.14	/	/	/	1	30	72	[71]
94	1.525	/	0.95	/	1	/	/	[72]
75.6	3.63	2.4	/	/	3	25	50	[73]
93.5	1.16	2.1	/	/	1	/	/	[74]
95	2.3	0.47	/	/	1	25	55	[75]
96.6	1.54	1.7	/	/	1	/	/	[76]
85	1.21	1.7	0.86	0.17	1	25	50	[77]
92	1.917	/	/	/	1	/	/	[78]
93.8	1.62	/	0.97	/	1	22	50	[79]
90.4	1.64	/	0.95	0.4	1	28	Nan	[80]
78	3.4	/	0.72	/	3	/	/	[81]
82	2.6	/	0.967	/	1	/	30	[82]
36	1.2	0.3	0.99	0.2	2	28	60	[83]
54	1.74	0.3	0.99	0.2	2	28	60	
58	1.85	0.3	0.99	0.2	2	28	60	
42	1.3	0.3	0.99	0.08	2	28	60	
59	1.91	0.3	0.99	0.08	2	28	60	
63	2.02	0.3	0.99	0.08	2	28	60	
50	1.6	0.3	0.99	0.06	2	28	60	
64	2.06	0.3	0.99	0.06	2	28	60	
69	2.19	0.3	0.99	0.06	2	28	60	
81.3	2.26	2	/	/	2	22	50	[84]
91.3	2.59	2	/	/	2.5	22	50	
90	3.6	0.1	/	0.06	1	26	30	[85]
65	2.5	0.1	/	0.06	1	26	30	
78	3	0.1	/	0.06	1	26	30	
74	2.6	0.1	/	0.06	1	26	30	
67	2.3	0.1	/	0.06	1	26	30	
92	3	1.6	0.92	/	1	18	60	[86]
28	0.36	1.3	0.97	/	0.4	26	60	[87]
59	0.75	1.3	0.97	/	0.6	26	60	
79	1.19	1.3	0.97	/	0.8	26	60	
93	1.66	1.3	0.97	/	1	26	60	
81	1.18	/	0.87	0.05	1	22	60	[88]
85	2.48	/	0.87	0.05	2	22	60	
91	5.27	/	0.87	0.05	4	22	60	
62	4.95	2.07	/	/	1	22	60	[89]
0.64	1.53	2.88	/	0.05	1	22	60	[90]
80	1.17	1.5	0.91	0.05	1	22	60	[91]

87	2.53	1.5	0.91	0.05	2	22	60	
92	5.28	1.5	0.91	0.05	4	22	60	
80	1.12	1.5	0.94	0.56	1	22	50	[92]
67	0.46	/	0.9	0.03	0.5	26	60	[93]
76	1.62	/	0.9	0.03	1.5	26	60	
78	2.18	/	0.9	0.03	2	26	60	
90	1.52	0.9	0.95	/	1	26	60	[94]
31	0.5	1.6	0.27	/	1	25	60	[95]
36	0.58	1.6	0.56	/	1	25	60	
48	0.76	1.6	0.87	/	1	25	60	
55	0.88	1.6	0.91	/	1	25	60	
62	0.99	1.6	0.95	/	1	25	60	
92	1.32	1.12	0.86	/	1	28	40	[96]
90	3.84	1.12	0.87	/	3	28	40	
92	6.26	1.12	0.87	/	5	28	40	
97	1.55	1.2	0.84	/	1	26	40	[97]
83	1.44	1.2	0.96	/	1	26	40	
87	1.4	1.1	0.95	/	1	25	60	[98]
73	1.68	1.12	0.99	/	1	22	30	[99]
84	1.27	0.8	0.9	/	1	22	40	[100]
84	3.87	0.8	0.9	/	3	22	40	
84	6.61	0.8	0.9	/	5	22	40	
75	1.88	1.38	0.99	0.03	1	22	30	[101]
79	3.95	1.38	0.99	0.03	1	22	30	
79	5.65	1.38	0.99	0.03	1	22	30	
92	2.46	1.7	0.95	0.06	1	26	52	[102]
94	1.57	1.5	0.9	0.04	1	25	50	[103]
82	1.39	1.5	0.87	0.04	1	25	50	
84	1.42	1.5	0.85	0.04	1	25	50	
71	1.23	1.5	0.82	0.04	1	25	50	
68	1.79	1.22	0.95	/	1	23	30	[104]
95	2.35	1.22	0.97	/	1	23	30	
70	1.75	1.22	0.93	/	1	23	30	
64	1.58	1.22	0.89	/	1	23	30	
4	0.84	1.22	0.8	/	1	23	30	
85	1.78	1.8	0.97	0.47	1	25	40	[105]
81	1.23	1.8	0.91	0.47	1	25	40	
97	2.17	1.35	0.78	0.05	1	25	60	[106]
66	0.98	/	0.71	/	1	22	60	[107]
90	1.5	1.2	0.91	0.6	1	21	50	[108]
42	0.77	1.2	0.62	0.6	1	21	50	
73	1.25	1.2	0.73	0.6	1	21	50	
81	1.37	1.2	0.78	0.6	1	21	50	
85	1.43	1.2	0.87	0.6	1	21	50	

84	1.86	2.26	0.95	0.12	1	25	65	[109]
78	2.18	2.26	0.95	0.12	1	25	65	
93	2.25	2.26	0.95	0.12	1	25	65	
80	2.05	2.26	0.95	0.12	1	25	65	
76	1.13	2.26	0.95	0.12	0.5	25	65	
94	3.1	2.26	0.95	0.12	1.5	25	65	
96	4.19	2.26	0.95	0.12	2	25	65	
96	5.16	2.26	0.95	0.12	3	25	65	
97	6.71	2.26	0.95	0.12	4	25	65	
97	8.27	2.26	0.95	0.12	5	25	65	
93	1.78	1.92	0.95	0.22	1	22	49	[110]
95	2.5	1.92	0.95	0.22	1.5	22	49	
97	3.36	1.92	0.95	0.22	2	22	49	
89	1.36	1.3	0.89	0.05	1	22	60	[111]
88	4	1.3	0.89	0.05	3	22	60	
89	6.8	1.3	0.89	0.05	5	22	60	
76	0.84	1.7	0.98	/	0.5	23	65	[112]
96	1.58	1.7	0.98	/	1	23	65	
86	2.11	1.7	0.98	/	1.5	23	65	
89	2.7	1.7	0.98	/	2	23	65	
86	1.36	1.7	0.98	/	1	23	65	
60	0.96	1.7	0.98	/	1	23	65	
90	1.34	3.11	0.95	0.06	1	25	60	[113]
90	2.34	3.11	0.95	0.06	1	25	60	
99	2.82	2.1	0.95	0.05	1	35	6	[114]
97	1.67	0.45	0.99	/	2	25	50	[115]

Table S2 Optimized hyperparameters of four models for predicting ER.

GBDT	n_estimators = 200 learning_rate = 0.25, max_depth = 3, min_samples_split = 3, reg_alpha = 0.8, random_state=42
SVM	C = 2, gamma = 0.1, kernel = 'linear'

XGBT	n_estimators = 120, learning_rate = 0.2, max_depth = 5, random_state=35
VR	n_estimators = 100, weight_gb = 2, weight_svm = 1, weight_xgb = 4 random_state=35

Table S3 Optimized hyperparameters of four models for predicting EE.

GBDT	n_estimators = 180, learning_rate = 0.2, max_depth = 5, min_samples_split = 6, reg_alpha = 0.3, random_state=42
SVM	C = 6, gama = 1, kernel = 'rbf'
XGBT	n_estimators = 200, learning_rate = 0.2, max_depth = 5, random_state=35
VR	n_estimators = 120, weight_gb = 1, weight_svm = 6, weight_xgb = 2 random_state=35

Table S4 Thermal conductivity of MCE membrane

Sample	Ambient temperature °C	Thermal conductivity W/(m K)
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Table S5. RMSE, MAE, R^2 of predicted ER for each model on training sets.

ML Models	RMSE	MAE	R^2
VR	0.1908 ± 0.0343	0.2731 ± 0.012	0.9627 ± 0.0061
XGBT	0.0853 ± 0.0541	0.1928 ± 0.0936	0.9829 ± 0.01
GBDT	0.2732 ± 0.0085	0.3404 ± 0.0062	0.9504 ± 0.0078
SVM	0.7080 ± 0.0316	0.4539 ± 0.0358	0.8997 ± 0.0327

Table S6. RMSE, MAE, R^2 of predicted ER for each model on test sets.

ML Models	RMSE	MAE	R^2
VR	0.3101 ± 0.0153	0.3756 ± 0.0173	0.9337 ± 0.0033
XGBT	0.3260 ± 0.0376	0.3927 ± 0.0166	0.9225 ± 0.0191
GBDT	0.5530 ± 0.0318	0.4952 ± 0.0325	0.8960 ± 0.0107
SVM	0.6188 ± 0.0126	0.6155 ± 0.0072	0.8708 ± 0.0031

Table S7. RMSE, MAE, R^2 of predicted EE for each model on training sets.

ML Models	RMSE	MAE	R^2
VR	0.0021 ± 0.0001	0.0283 ± 0.001	0.9082 ± 0.006
XGBT	0.0039 ± 0.0003	0.043 ± 0.0052	0.8913 ± 0.0113
GBDT	0.0028 ± 0.0016	0.0302 ± 0.0015	0.8905 ± 0.0097
SVM	0.0012 ± 0.0008	0.0196 ± 0.0027	0.9087 ± 0.0011

Table S8. RMSE, MAE, R^2 of predicted EE for each model on test sets.

ML Models	RMSE	MAE	R^2
VR	0.0075 ± 0.0001	0.0628 ± 0.0003	0.7242 ± 0.0011
XGBT	0.0086 ± 0.0004	0.0681 ± 0.0031	0.697 ± 0.0221

GBDT	0.0092 ± 0.0002	0.0718 ± 0.0053	0.5921 ± 0.0161
SVM	0.0077 ± 0.0001	0.0649 ± 0.0003	0.6527 ± 0.0038

1.1 Programming languages of ML and the core software

All ML models were implemented in Python, installing Python 3 and packages (Numpy, Pandas, Scipy, Scikit-learn, Shap, Matplotlib, and Interpreter).

Open source software libraries lists:

(1) Voting Regression (VR) model

<https://scikit-learn.org/stable/modules/ensemble.html>

<https://github.com/scikit-learn/scikit-learn>

<https://scikit-learn.org/stable/modules/ensemble.html>

(2) Extreme gradient boosting (XGBT) model

<https://github.com/dmlc/xgboost>

<https://xgboost.readthedocs.io/>

(3) CatBoost model

<https://github.com/catboost/catboost>

<https://catboost.ai/en/docs/>

(4) Gaussian process regression (GPR) model

https://scikit-learn.org/stable/modules/gaussian_process.html

<https://github.com/scikit-learn/scikit-learn>

<https://scikit-learn.org/stable/modules/ensemble.html>

1.2 Characterizations

The crystal structure was measured by X-ray diffraction spectroscopy (XRD, Holland Panalytical PRO PW3040/60). The elemental composition and atomic ratios were characterized by X-ray photoelectron spectroscopy (XPS, Thermo Scientific K-Alpha). The surface temperature of membranes was taken by an infrared thermal imager (FLTR E95). The band gap was measured using an ultraviolet-visible-near-infrared spectrophotometer (Shimadzu UV-3600i Plus). Raman spectral data were measured using a Raman

spectrometer (Horiba LabRAM HR Evolution, Japan). Scanning electron microscopy (SEM) images were acquired using a Zeiss GeminiSEM 500 field-emission scanning electron microscope (FESEM). EPR measurements were performed on a Bruker ELEXSYS E580 X-band spectrometer with an ER4108 TMHS cavity. TEM images were acquired on a FEI Talos F200X transmission electron microscope.

1.3 Solar evaporation experiments

In solar steam generation experiment, a beaker containing 180 mL of distilled water was used as the container. The membrane evaporator was installed, supported by foam wrapped with dust-free paper. The entire setup was placed on an electronic balance (Sartorius BSA224S) to continuously monitor and record the mass change of the distilled water due to evaporation. Subsequently, the beaker was exposed to a 500 W xenon lamp equipped with a standard AM 1.5 G filter to conduct the evaporation experiment. The simulation of solar radiation intensity was measured using an optical power meter (CEL-NP2000-2). Every 10 minutes, surface temperature and mass changes of the membranes were recorded with an infrared camera (FLIR E95) and an electronic balance.

The solar evaporation efficiency (η , %) was calculated using the following equations:

$$\eta = \frac{\dot{m}\Delta H}{C_{opt}P} \quad (S1)$$

$$\Delta H = H_V + Q \quad (S2)$$

$$Q = c(T - T_1) \quad (S3)$$

$$H_V = 1.91846 \times 10^6 \times \left[\frac{T}{T - 33.91} \right]^2 \quad (S4)$$

where $\dot{m}$ is the evaporation rate ($\text{kg m}^{-2} \text{h}^{-1}$), $\dot{m} = \dot{m}_{light} - \dot{m}_{dark}$, $\dot{m}_{light}$ and $\dot{m}_{dark}$ are the solar water evaporation rate under the sunlight and dark conditions, respectively. C_{opt} is the optical concentration. P is the incident solar power (1 kW m^{-2} , one sun). ΔH is the liquid-vapor phase change enthalpy, H_V is the phase-change enthalpy (J kg^{-1}), and Q is the sensible heat of water. T is the surface temperature of solar evaporators, T_1 is the initial temperature of the water, and c is the specific heat of water ($4.2 \text{ J g}^{-1} \text{K}^{-1}$). In this work, only the phase-change enthalpy is considered when calculating the evaporation efficiency.



Fig. S1. Four different prepared samples.

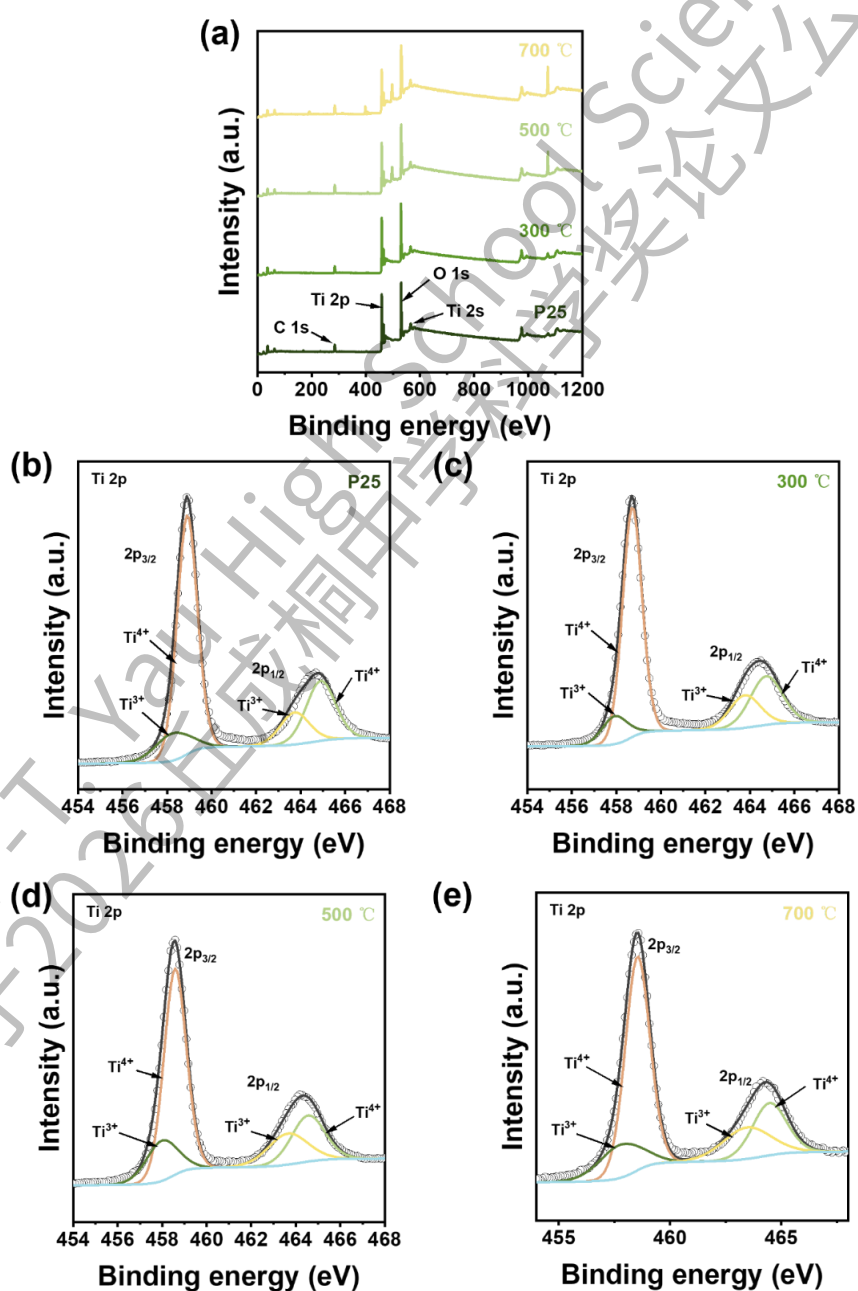


Fig. S2. (a) XPS survey. High-resolution Ti 2p spectra of (b) P25, (c) 300 °C, (d) 500 °C and (e) 700 °C.

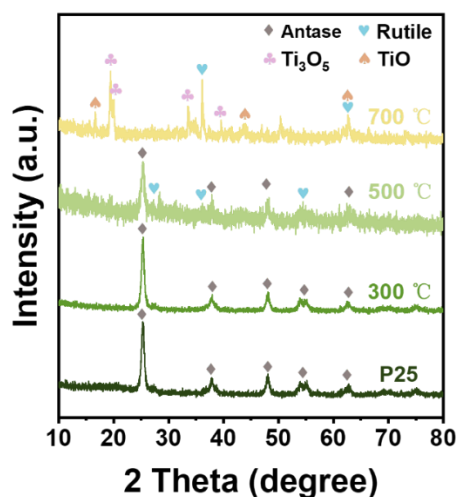


Fig. S3. XRD pattern.

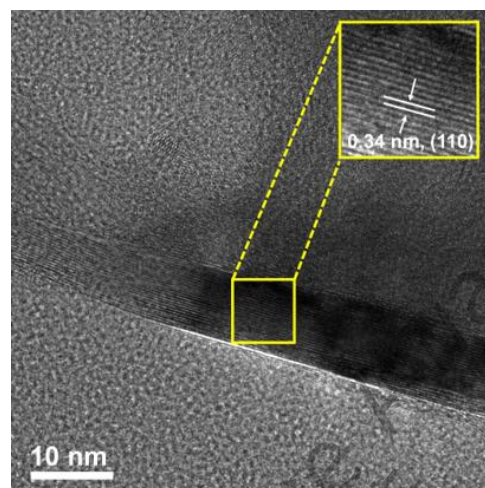


Fig. S4. HR-TEM image of 700 °C.

Table S9 Comparison of evaporation performance of different semiconductor materials

Semiconductor materials	Solar intensity (kw m ⁻²)	Evaporation Efficiency (%)	Evaporation rate (kg m ⁻² h ⁻¹)	Refs.
Black TiOx	1	50.3	0.8	[6]
Fe ₃ O ₄ MNPs	1	63	0.9	[8]
Ti ₃ C ₂	1	53.7	0.45	[10]
Fe ₃ O ₄ /C	1	67	1.07	[26]
WO ₃	1	57.3	1	[38]
Cu _{2-x} S NWs	1	67	0.95	[49]
TiN	1	64.1	0.94	[59]
TiO ₂ NWAs	1	59	1.25	[68]
TiO ₂ -700 °C	1	67.8	1.12	This work
Fe ₃ O ₄ MNPs	2	65	1.85	[8]
MoS ₂	2	61	1.95	[21]
Cu-Fe ₃ O ₄	2	36	1.2	[83]
Cu-Fe ₃ O ₄	2	42	1.3	[83]
TiO ₂ -700 °C	2	51.2	1.62	This work

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Appendix B: Experimental Process.



Fig. B1. Experimental tests of solar water evaporation.

Appendix C: 半导体光热材料性能研究的机器学习算法源代码（用 Python 语言编写，程序太多只附上了部分）

1、半导体多重插补处理缺失值 2.py

#处理缺失值程序

```
import numpy as np    #安装 numpy 库用于科学计算
import pandas as pd    #安装 pandas 库加载数据，并使用 numpy 进行数值操作
from sklearn.experimental import enable_iterative_imputer
from sklearn.impute import IterativeImputer
from sklearn.linear_model import BayesianRidge

pd.set_option('display.max_rows', None)
pd.set_option('display.max_columns', None)

filename = '半导体光热脱盐材料特征数据收集(终).csv'
data = pd.read_csv(filename,encoding='gbk')    #加载数据
print(data.head())    #查看数据
print(data.describe())
print(data.info())
print(data.shape)

# 将 DataFrame 转换为 NumPy 数组
#data1 = data.drop(['吸光材料','文献'],axis=1)    #假设['吸光材料','文献']是目标变量
data1 = data.replace({pd.NaT:np.nan})
data2 = data1.to_numpy()

# 初始化 IterativeImputer
imputer = IterativeImputer(BayesianRidge(), max_iter=10)

# 拟合数据并生成插补的数据
imputed_data = imputer.fit_transform(data2)

# 将插补的数据转换回 Pandas DataFrame
imputed_df = pd.DataFrame(imputed_data, columns=data1.columns)

imputed_df.to_csv('半导体插补数据.csv',index=False,encoding='gbk')
```

2、GBDT_model_training_best1.py

GBDT-ER 模型训练的程序—半导体光热材料的蒸发速率数据的机器学习

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split, cross_val_score
from sklearn.metrics import mean_squared_error, mean_absolute_error, r2_score
from sklearn.ensemble import GradientBoostingRegressor
from sklearn.model_selection import KFold
from sklearn.model_selection import GridSearchCV
from skopt import gp_minimize
from skopt.space import Real, Integer
from sklearn.preprocessing import StandardScaler
from matplotlib import pyplot as plt

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

# 导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename, encoding='gbk')
df2 = df.iloc[:, range(7)]
data = df.iloc[:, [2, 3, 4, 5, 6, 7]] # 所需要的材料特征
target = df.iloc[:, [0, 1]] # 前面两列
# target1 = df.iloc[:, 0]
# target2 = df.iloc[:, 1]

X = data.values
# Y_1 = target1.values
# Y_2 = target2.values
Y = target.values

# 标准化数据，归一化特征
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=10)
Y_train_1 = Y_train[:, 0]
Y_train_2 = Y_train[:, 1]
Y_test_1 = Y_test[:, 0]
Y_test_2 = Y_test[:, 1]

# 创建梯度提升回归模型
gb_reg = GradientBoostingRegressor()

# 定义超参数搜索空间
param_space = {
    'n_estimators': Integer(50, 300),
    'learning_rate': Real(0.01, 0.3),
    'max_depth': Integer(2, 5),

```

```

        'alpha': Real(0.5, 0.9),
        'min_samples_split': Integer(2, 10),
        'min_samples_leaf': Integer(2, 10),
    }

```

转换超参数空间格式

```

dimensions = [
    param_space['n_estimators'],
    param_space['learning_rate'],
    param_space['max_depth'],
    param_space['alpha'],
    param_space['min_samples_split'],
    param_space['min_samples_leaf'],
]

```

定义贝叶斯优化目标函数

```

def objective(params):
    n_estimators = params[0]
    learning_rate = params[1]
    max_depth = params[2]
    alpha = params[3]
    min_samples_split = params[4]
    min_samples_leaf = params[5]

    model = GradientBoostingRegressor(
        n_estimators=n_estimators,
        learning_rate=learning_rate,
        max_depth=max_depth,
        alpha=alpha,
        min_samples_split=min_samples_split,
        min_samples_leaf=min_samples_leaf,
        random_state=42
    )

```

#训练模型

使用交叉验证评估模型，对于回归问题使用负均方误差作为评分标准

```

scores = cross_val_score(model, X_train, Y_train_2, cv=5,
    scoring='neg_mean_squared_error')

```

返回负平均分数，因为 gp_minimize 尝试最小化目标函数

```

return -scores.mean()

```

执行贝叶斯优化

```

res = gp_minimize(
    objective,
    dimensions,

```

```

        n_calls=300,
        random_state=42,
        n_jobs=-1
    )

    best_params = {
        'n_estimators': res.x[0],
        'learning_rate': res.x[1],
        'max_depth': res.x[2],
        'alpha': res.x[3],
        'min_samples_split': res.x[4],
        'min_samples_leaf': res.x[5],
    }

    print("最佳超参数: ", best_params)

```

使用最佳超参数训练模型

```

best_model = GradientBoostingRegressor(
    n_estimators=best_params['n_estimators'],
    learning_rate=best_params['learning_rate'],
    max_depth=best_params['max_depth'],
    alpha=best_params['alpha'],
    min_samples_split=best_params['min_samples_split'],
    min_samples_leaf=best_params['min_samples_leaf'],
    random_state=42
)
best_model.fit(X_train, Y_train_2)

```

使用最佳模型进行预测

训练数据的预测

```
y_train_pred2 = best_model.predict(X_train)
```

测试数据的预测

```

predictions_2 = best_model.predict(X_test)
#result = pd.DataFrame([predictions_2, Y_test_2])
#result.to_csv('GBDT_PT_2.csv',index=False,encoding='gbk')

```

#计算训练数据的 MSE 和 MAE，评估模型性能

```

train_mse = mean_squared_error(Y_train_2,y_train_pred2)
train_mae = mean_absolute_error(Y_train_2,y_train_pred2)
train_r2 = r2_score(Y_train_2,y_train_pred2)

```

计算测试数据的 MSE 和 MAE

```
test_mse = mean_squared_error(Y_test_2, predictions_2)
```

```
test_mae = mean_absolute_error(Y_test_2, predictions_2)
test_r2 = r2_score(Y_test_2, predictions_2)
```

```
print("训练数据的 MSE:", train_mse)
print("训练数据的 MAE:", train_mae)
print("训练数据的 R2:", train_r2)
print("测试数据的 MSE:", test_mse)
print("测试数据的 MAE:", test_mae)
print("测试数据的 R2:", test_r2)
```

#绘制模型性能散点图

```
plt.figure(figsize=(12,10))
plt.plot([0,17], [0,17], color='black',linewidth=1,label='best line')
plt.scatter(Y_train_2, y_train_pred2,label='train data',color='lightblue',s=80)
plt.scatter(Y_test_2, predictions_2,label='test data',color='lightcoral',s=80)
plt.xlabel('True',fontsize=30,fontweight='bold')
plt.ylabel('Predicted',fontsize=30,fontweight='bold')
plt.title('GBDT',fontsize=24,fontweight='bold')
plt.legend(fontsize=20)
plt.xticks(fontsize=20,fontweight='bold')
plt.yticks(fontsize=20,fontweight='bold')
ax = plt.gca()
ax.spines['bottom'].set_linewidth(3)
ax.spines['left'].set_linewidth(3)
ax.spines['top'].set_linewidth(3)
ax.spines['right'].set_linewidth(3)
plt.show()
```

3、SVM_model_training_best1.py

SVM-ER 模型的训练程序—半导体光热材料的蒸发速率数据的机器学习

```
import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split,cross_val_score
from sklearn.metrics import mean_squared_error,mean_absolute_error,r2_score
from sklearn.svm import SVR
from sklearn.model_selection import KFold
from sklearn.model_selection import GridSearchCV
from hyperopt import hp, fmin, tpe, Trials
from sklearn.preprocessing import StandardScaler
from matplotlib import pyplot as plt
```

```

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=10)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建五折交叉验证
#kf = KFold(n_splits=5, shuffle=True, random_state=100)

# 创建 SVM 模型
svm = SVR()

# 定义超参数搜索空间
space = {
    'C': hp.loguniform('C', np.log(1e-3), np.log(1e3)), # 正则化参数
    'epsilon': hp.loguniform('epsilon', np.log(1e-3), np.log(1e3)), #  $\epsilon$  - 不敏感管的宽度
    'gamma': hp.loguniform('gamma', np.log(1e-3), np.log(1e3)), # 核函数的系数
    'kernel': hp.choice('kernel', ['linear', 'poly', 'rbf', 'sigmoid']) # 核函数类型
}

# 定义目标函数
def objective(params):
    params = {
        'C': params['C'],
        'epsilon': params['epsilon'],
        'gamma': params['gamma'],
        'kernel': params['kernel']
    }

```

```

# 使用交叉验证来评估模型性能
model = SVR(**params)
scores = cross_val_score(model, X_train, Y_train_2, cv=5, scoring='r2') # 使用 R 平方作为评估指标
loss = -np.mean(scores) # 因为 hyperopt 是用于最小化目标函数，所以取负值

return loss

# 运行贝叶斯优化
trials = Trials()
best = fmin(
    fn=objective,
    space=space,
    algo=tpe.suggest,
    max_evals=50, # 最大迭代次数
    trials=trials
)

# 输出最佳超参数
best_kernel = ['linear', 'poly', 'rbf', 'sigmoid'][best['kernel']]
best_params = {
    'C': best['C'],
    'epsilon': best['epsilon'],
    'gamma': best['gamma'],
    'kernel': best_kernel
}
print("最佳超参数: ", best_params)

# 使用最佳超参数训练最终模型
best_model = SVR(**best_params)

# 使用最佳模型进行预测

best_model.fit(X_train, Y_train_2)

# 训练数据的预测
y_train_pred2 = best_model.predict(X_train)

# 测试数据的预测
predictions_2 = best_model.predict(X_test)
#result = pd.DataFrame([predictions_2, Y_test_2])
#result.to_csv('GBDT_PT_2.csv',index=False,encoding='gbk')

#计算训练数据的 MSE 和 MAE

```

```

train_mse = mean_squared_error(Y_train_2,y_train_pred2)
train_mae = mean_absolute_error(Y_train_2,y_train_pred2)
train_r2 = r2_score(Y_train_2,y_train_pred2)

```

```

# 计算测试数据的 MSE 和 MAE

```

```

test_mse = mean_squared_error(Y_test_2, predictions_2)
test_mae = mean_absolute_error(Y_test_2, predictions_2)
test_r2 = r2_score(Y_test_2,predictions_2)

```

```

print("训练数据的 MSE:", train_mse)
print("训练数据的 MAE:", train_mae)
print("训练数据的 R2:", train_r2)
print("测试数据的 MSE:", test_mse)
print("测试数据的 MAE:", test_mae)
print("测试数据的 R2:", test_r2)

```

```

#绘制模型性能散点图

```

```

plt.figure(figsize=(12,10))
plt.plot([0,17], [0,17], color='black',linewidth=1,label='best line')
plt.scatter(Y_train_2, y_train_pred2,label='train data',color='lightblue',s=30)
plt.scatter(Y_test_2, predictions_2,label='test data',color='lightcoral',s=30)
plt.xlabel('True',fontsize=30,fontweight='bold')
plt.ylabel('Predicted',fontsize=30,fontweight='bold')
plt.title('SVM',fontsize=24,fontweight='bold')
plt.legend(fontsize=20)
plt.xticks(fontsize=20,fontweight='bold')
plt.yticks(fontsize=20,fontweight='bold')
ax = plt.gca()
ax.spines['bottom'].set_linewidth(3)
ax.spines['left'].set_linewidth(3)
ax.spines['top'].set_linewidth(3)
ax.spines['right'].set_linewidth(3)
plt.show()

```

4、XGboosting_model_training_best1.py

```

# XGBT-ER 模型的训练程序—半导体光热材料的蒸发速率数据的机器学习

```

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split
from sklearn.metrics import mean_squared_error,mean_absolute_error,r2_score

```

```

from sklearn.model_selection import KFold,cross_val_score
from sklearn.model_selection import GridSearchCV
from sklearn.preprocessing import StandardScaler
from skopt import gp_minimize
from skopt.space import Real, Integer
from xgboost import XGBRegressor
from matplotlib import pyplot as plt

```

```

pd.set_option('display.max_rows', None)
pd.set_option('display.max_columns', None)
plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

```

#导入数据

```

filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(2,7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

```

```

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

```

#标准化数据

```

X = StandardScaler().fit_transform(X)
X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=10)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

```

初始化 XGBoost 回归模型

```
xgb_model = XGBRegressor()
```

定义超参数搜索空间

```

param_space = {
    'learning_rate': Real(0.01, 0.8),
    'max_depth': Integer(3, 6),
    'n_estimators': Integer(50, 300),
    'gamma': Real(0, 0.5),
    'subsample': Real(0.5, 0.8),
    'reg_alpha': Real(0, 1)
}

```

```
}
```

```
dimensions = [  
    param_space['learning_rate'],  
    param_space['max_depth'],  
    param_space['n_estimators'],  
    param_space['gamma'],  
    param_space['subsample'],  
    param_space['reg_alpha']  
]
```

```
# 定义贝叶斯优化目标函数
```

```
def objective(params):
```

```
    learning_rate = params[0]
```

```
    max_depth = params[1]
```

```
    n_estimators = params[2]
```

```
    gamma = params[3]
```

```
    subsample = params[4]
```

```
    reg_alpha = params[5]
```

```
    model = XGBRegressor(  
        learning_rate=learning_rate,  
        max_depth=max_depth,  
        n_estimators=n_estimators,  
        gamma=gamma,  
        subsample=subsample,  
        reg_alpha=reg_alpha,  
        random_state=42  
    )
```

```
    scores = cross_val_score(model, X_train, Y_train_2, cv=5,  
scoring='neg_mean_squared_error')  
    return -scores.mean()
```

```
# 执行贝叶斯优化
```

```
res = gp_minimize(  
    objective,  
    dimensions,  
    n_calls=30,  
    random_state=42,  
    n_jobs=-1  
)
```

```
best_params = {  
    'learning_rate': res.x[0],  
    'max_depth': res.x[1],
```

```

        'n_estimators': res.x[2],
        'gamma': res.x[3],
        'subsample': res.x[4],
        'reg_alpha': res.x[5]
    }

```

```

print("最佳超参数: ", best_params)

```

```

# 使用最佳超参数训练模型

```

```

best_model = XGBRegressor(
    learning_rate=best_params['learning_rate'],
    max_depth=best_params['max_depth'],
    n_estimators=best_params['n_estimators'],
    gamma=best_params['gamma'],
    subsample=best_params['subsample'],
    reg_alpha=best_params['reg_alpha'],
    random_state=42
)

```

```

best_model.fit(X_train, Y_train_2)

```

```

# 使用最佳模型进行预测

```

```

# 训练数据的预测

```

```

y_train_pred2 = best_model.predict(X_train)

```

```

# 测试数据的预测

```

```

predictions_2 = best_model.predict(X_test)
#result = pd.DataFrame([predictions_2, Y_test_2])
#result.to_csv('GBDT_PT_2.csv',index=False,encoding='gbk')

```

```

#计算训练数据的 MSE 和 MAE

```

```

train_mse = mean_squared_error(Y_train_2,y_train_pred2)
train_mae = mean_absolute_error(Y_train_2,y_train_pred2)
train_r2 = r2_score(Y_train_2,y_train_pred2)

```

```

# 计算测试数据的 MSE 和 MAE

```

```

test_mse = mean_squared_error(Y_test_2, predictions_2)
test_mae = mean_absolute_error(Y_test_2, predictions_2)
test_r2 = r2_score(Y_test_2,predictions_2)

```

```

print("训练数据的 MSE:", train_mse)
print("训练数据的 MAE:", train_mae)
print("训练数据的 R2:", train_r2)
print("测试数据的 MSE:", test_mse)
print("测试数据的 MAE:", test_mae)

```

```
print("测试数据的 R2:", test_r2)
```

```
#绘制模型性能散点图
```

```
plt.figure(figsize=(12,10))
```

```
plt.plot([0,17], [0,17], color='black',linewidth=1,label='best line')
```

```
plt.scatter(Y_train_2, y_train_pred2,label='train data',color='lightblue',s=80)
```

```
plt.scatter(Y_test_2, predictions_2,label='test data',color='lightcoral',s=80)
```

```
plt.xlabel('True',fontsize=30,fontweight='bold')
```

```
plt.ylabel('Predicted',fontsize=30,fontweight='bold')
```

```
plt.title('XGBT',fontsize=24,fontweight='bold')
```

```
plt.legend(fontsize=20)
```

```
plt.xticks(fontsize=20,fontweight='bold')
```

```
plt.yticks(fontsize=20,fontweight='bold')
```

```
ax = plt.gca()
```

```
ax.spines['bottom'].set_linewidth(3)
```

```
ax.spines['left'].set_linewidth(3)
```

```
ax.spines['top'].set_linewidth(3)
```

```
ax.spines['right'].set_linewidth(3)
```

```
plt.show()
```

5、Voting_model_training_best1.py

VR-ER 模型的训练程序—半导体光热材料的蒸发速率数据的机器学习

```
from sklearn.model_selection import train_test_split
```

```
from sklearn.ensemble import VotingRegressor
```

```
from sklearn.metrics import mean_squared_error,mean_absolute_error,r2_score
```

```
from sklearn.preprocessing import StandardScaler
```

```
from sklearn.ensemble import GradientBoostingRegressor
```

```
from sklearn.svm import SVR
```

```
from xgboost import XGBRegressor
```

```
from matplotlib import pyplot as plt
```

```
import numpy as np
```

```
import pandas as pd
```

```
plt.rcParams['font.family'] = ['Microsoft YaHei']
```

```
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号
```

```
#导入数据
```

```
filename = '半导体插补数据.csv'
```

```
df = pd.read_csv(filename,encoding='gbk')
```

```
df2 = df.iloc[:,range(7)]
```

```
data = df.iloc[:,[2,3,4,5,6,7]]
```

```

target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=10)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建模型
svr = SVR(C=42.448789, kernel='rbf', epsilon=0.005106, gamma=0.004153)
xgb = XGBRegressor(gamma=0.491157, learning_rate=0.227627, n_estimators=289, max_depth=4,
                  reg_alpha=1.0, subsample=0.613525)
gb = GradientBoostingRegressor(alpha=0.9, learning_rate=0.3, n_estimators=50,
                               max_depth=2, min_samples_leaf=2, min_samples_split=2)

# 创建投票集成模型
voting_regressor = VotingRegressor([
    ('svr', svr),
    ('xgb', xgb),
    ('gb', gb)
], weights=[80, 800, 20])

voting_regressor.fit(X_train, Y_train_2)
# 训练数据的预测
y_train_pred2 = voting_regressor.predict(X_train)

# 测试数据的预测
predictions_2 = voting_regressor.predict(X_test)

#计算训练数据的 MSE 和 MAE

train_mse = mean_squared_error(Y_train_2, y_train_pred2)
train_mae = mean_absolute_error(Y_train_2, y_train_pred2)

```

```

train_r2 = r2_score(Y_train_2,y_train_pred2)

# 计算测试数据的 MSE 和 MAE
test_mse = mean_squared_error(Y_test_2, predictions_2)
test_mae = mean_absolute_error(Y_test_2, predictions_2)
test_r2 = r2_score(Y_test_2,predictions_2)

print("训练数据的 MSE:", train_mse)
print("训练数据的 MAE:", train_mae)
print("训练数据的 R2:", train_r2)
print("测试数据的 MSE:", test_mse)
print("测试数据的 MAE:", test_mae)
print("测试数据的 R2:", test_r2)

#绘制模型性能散点图
plt.figure(figsize=(12,10))
plt.plot([0,17], [0,17], color='black',linewidth=1,label='best line')
plt.scatter(Y_train_2, y_train_pred2,label='train data',color='lightblue',s=80)
plt.scatter(Y_test_2, predictions_2,label='test data',color='lightcoral',s=80)
plt.xlabel('True',fontsize=30,fontweight='bold')
plt.ylabel('Predicted',fontsize=30,fontweight='bold')
plt.title('VR',fontsize=24,fontweight='bold')
plt.legend(fontsize=20)
plt.xticks(fontsize=20,fontweight='bold')
plt.yticks(fontsize=20,fontweight='bold')
ax = plt.gca()
ax.spines['bottom'].set_linewidth(3)
ax.spines['left'].set_linewidth(3)
ax.spines['top'].set_linewidth(3)
ax.spines['right'].set_linewidth(3)
plt.show()

```

10、Shap_value_VOTE_vra.py

Shap-ER 模型的训练程序（用 VR 预测）-半导体光热材料的蒸发效率数据的机器学习

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split
from sklearn.preprocessing import StandardScaler
from sklearn.ensemble import GradientBoostingRegressor
from sklearn.svm import SVR
from xgboost import XGBRegressor
from sklearn.ensemble import VotingRegressor
from matplotlib import pyplot as plt
import shap

```

```

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=10)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建模型
svr = SVR(C=42.448789, kernel='rbf', epsilon=0.005106,gamma=0.004153)
xgb = XGBRegressor(gamma=0.491157,learning_rate=0.227627,n_estimators=289,max_depth=4,
                  reg_alpha=1.0,subsample=0.613525)
gb = GradientBoostingRegressor(alpha=0.9, learning_rate=0.3, n_estimators=50,
                                max_depth=2,
                                min_samples_leaf=2, min_samples_split=2)

# 创建投票集成模型
voting_regressor = VotingRegressor([
    ('svr', svr),
    ('xgb', xgb),
    ('gb', gb)
],weights=[80,800,20])

voting_regressor.fit(X_train,Y_train_2)

def vr_predict(X):

```

```

return voting_regressor.predict(X)

# 使用 Shap 库计算 Shap 值
explainer = shap.PermutationExplainer(vr_predict,X_train)
shap_values = explainer.shap_values(X_train)
#print(shap_values)
# 显示 Shap 值的密度散点图
shap.summary_plot(shap_values,X_train,feature_names=data.columns)

```

11、Shap_value_VOTE_cve.py

Shap-EE 模型的训练程序（用 **VR** 预测）--半导体光热材料的蒸发效率数据的机器学习

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split
from sklearn.preprocessing import StandardScaler
from sklearn.ensemble import GradientBoostingRegressor
from sklearn.svm import SVR
from xgboost import XGBRegressor
from sklearn.ensemble import VotingRegressor
from matplotlib import pyplot as plt
import shap

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

```

#导入数据

```

filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

```

```

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

```

#标准化数据

```

X = StandardScaler().fit_transform(X)

```

```

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=1)
Y_train_1 = Y_train[:,0]

```

```

Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建模型
svr = SVR(C=4.170161, kernel='rbf', epsilon=0.019841,gamma=0.044303)
xgb = XGBRegressor(gamma=0.265641,learning_rate=0.8,n_estimators=237,max_depth=6,
                    reg_alpha=0.0,subsample=0.657072)
gb = GradientBoostingRegressor(alpha=0.9, learning_rate=0.080013, n_estimators=76,
                                max_depth=2,
                                min_samples_leaf=4, min_samples_split=2)

# 创建投票集成模型
voting_regressor = VotingRegressor([
    ('svr', svr),
    ('xgb', xgb),
    ('gb', gb)
],weights=[30,5,5])

voting_regressor.fit(X_train,Y_train_1)

def vr_predict(X):
    return voting_regressor.predict(X)

# 使用 Shap 库计算 Shap 值
explainer = shap.PermutationExplainer(vr_predict,X_train)
shap_values = explainer.shap_values(X_train)
#print(shap_values)
# 显示 Shap 值的密度散点图
shap.summary_plot(shap_values,X_train,feature_names=data.columns)

```

12、数据相关性绘图 1.py

Feature Correlation Heatmap 的 ER 相关性热图程序

```

import numpy as np
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'

```

```

df = pd.read_csv(filename,encoding='gbk')
df1 = df.iloc[:,1,2,3,4,5,6,7]]
df2 = df.iloc[:,0,2,3,4,5,6,7]]
data = df.iloc[:,range(2,7)]
target = df.iloc[:,[0,1]]
target1 = df.iloc[:,0]
target2 = df.iloc[:,1]

X = data.values
Y_1 = target1.values
Y_2 = target2.values

#相关性分析
# 计算特征之间的相关性矩阵
correlation_matrix = df1.corr()
# 使用热力图可视化相关性矩阵
plt.figure(figsize=(8, 6))
#for (i, j), z in np.ndenumerate(correlation_matrix):
    #plt.text(j, i, f'{z:.2f}', ha='center', va='center', fontsize=12)
sns.heatmap(correlation_matrix, annot=True,cmap='YlOrRd',
linewidths=0.5,mask=np.triu(np.ones_like(correlation_matrix, dtype=bool),k=1))
plt.xticks(rotation=45, ha='right',fontsize=10) # 旋转 x 轴标签（特征名称）
plt.yticks(rotation=0,fontsize=10) # 确保 y 轴标签（特征名称）正着显示
plt.title('Feature Correlation Heatmap')
plt.show()

```

13、数据相关性绘图 2.py

Feature Correlation Heatmap 的 EE 相关性热图程序

```

import numpy as np
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df1 = df.iloc[:,0,2,3,4,5,6,7]]
df2 = df.iloc[:,1,2,3,4,5,6,7]]
data = df.iloc[:,range(2,7)]
target = df.iloc[:,[0,1]]
target1 = df.iloc[:,0]

```

```
target2 = df.iloc[:,1]
```

```
X = data.values
```

```
Y_1 = target1.values
```

```
Y_2 = target2.values
```

```
#相关性分析
```

```
# 计算特征之间的相关性矩阵
```

```
correlation_matrix = df1.corr()
```

```
# 使用热力图可视化相关性矩阵
```

```
plt.figure(figsize=(8, 6))
```

```
#for (i, j), z in np.ndenumerate(correlation_matrix):
```

```
    #plt.text(j, i, f'{z:.2f}', ha='center', va='center', fontsize=12)
```

```
sns.heatmap(correlation_matrix, annot=True,cmap='YlOrRd',
```

```
linewidths=0.5,mask=np.triu(np.ones_like(correlation_matrix, dtype=bool),k=1))
```

```
plt.xticks(rotation=45, ha='right',fontsize=10) # 旋转 x 轴标签（特征名称）
```

```
plt.yticks(rotation=0,fontsize=10) # 确保 y 轴标签（特征名称）正着显示
```

```
plt.title('Feature Correlation Heatmap')
```

```
plt.show()
```

14、# 根据 GBDT 模型结果做 Shap 法中 EE 图

```
Shap_value_GBDT_cve.py
```

```
Shap-GBDT-EE 根据 GBDT 做 Shap 表
```

```
import numpy as np
```

```
import pandas as pd
```

```
from sklearn.model_selection import train_test_split
```

```
from sklearn.metrics import mean_squared_error
```

```
from sklearn.model_selection import KFold,cross_val_score
```

```
from sklearn.model_selection import GridSearchCV
```

```
from sklearn.preprocessing import StandardScaler
```

```
from sklearn.ensemble import GradientBoostingRegressor
```

```
from matplotlib import pyplot as plt
```

```
import shap
```

```
plt.rcParams['font.family'] = ['Microsoft YaHei']
```

```
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号
```

```
#导入数据
```

```
filename = '半导体插补数据.csv'
```

```
df = pd.read_csv(filename,encoding='gbk')
```

```
df2 = df.iloc[:,range(7)]
```

```
data = df.iloc[:,[2,3,4,5,6,7]]
```

```
target = df.iloc[:,[0,1]]
```

```
#target1 = df.iloc[:,0]
```

```
#target2 = df.iloc[:,1]
```

```

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=100)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建梯度提升回归模型
gb_reg = GradientBoostingRegressor(alpha=0.7, learning_rate=0.1, n_estimators=250,
max_depth=3,
                                min_samples_leaf=2, min_samples_split=3)

gb_reg.fit(X_train, Y_train_1)

# 使用 Shap 库计算 Shap 值
explainer = shap.TreeExplainer(gb_reg, X_train)
shap_values = explainer.shap_values(X_train)
#print(shap_values)
# 显示 Shap 值的密度散点图
shap.summary_plot(shap_values,X_train,feature_names=data.columns)

```

15、# 根据 GBDT 模型结果做 Shap 法中 ER 图

Shap_value_GBDT_vra.py

Shap-GBDT-ER

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split
from sklearn.metrics import mean_squared_error
from sklearn.model_selection import KFold,cross_val_score
from sklearn.model_selection import GridSearchCV
from sklearn.preprocessing import StandardScaler
from sklearn.ensemble import GradientBoostingRegressor
from matplotlib import pyplot as plt
import shap

```

```
plt.rcParams['font.family'] = ['Microsoft YaHei']
```

```

plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values

#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=100)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建梯度提升回归模型
gb_reg = GradientBoostingRegressor(alpha=0.5, learning_rate=0.3, n_estimators=200,
max_depth=3,
min_samples_leaf=1, min_samples_split=3)

gb_reg.fit(X_train, Y_train_2)

# 使用 Shap 库计算 Shap 值
explainer = shap.TreeExplainer(gb_reg, X_train)
shap_values = explainer.shap_values(X_train)
#print(shap_values)
# 显示 Shap 值的密度散点图
shap.summary_plot(shap_values,X_train,feature_names=data.columns)

```

16、# 根据 SVM 模型结果做 Shap 法中 ER 图

Shap_value_SVM_vra.py

Shap-SVM-ER 用 SVM 生成 Shap 图像

```

import numpy as np
import pandas as pd
from sklearn.model_selection import train_test_split

```

```

from sklearn.metrics import mean_squared_error
from sklearn.svm import SVR
from sklearn.model_selection import KFold
from sklearn.model_selection import GridSearchCV
from sklearn.preprocessing import StandardScaler
from matplotlib import pyplot as plt
import shap

plt.rcParams['font.family'] = ['Microsoft YaHei']
plt.rcParams['axes.unicode_minus'] = False # 正确显示负号

#导入数据
filename = '半导体插补数据.csv'
df = pd.read_csv(filename,encoding='gbk')
df2 = df.iloc[:,range(7)]
data = df.iloc[:,[2,3,4,5,6,7]]
target = df.iloc[:,[0,1]]
#target1 = df.iloc[:,0]
#target2 = df.iloc[:,1]

X = data.values
#Y_1 = target1.values
#Y_2 = target2.values
Y = target.values#标准化数据
X = StandardScaler().fit_transform(X)

X_train, X_test, Y_train, Y_test = train_test_split(X, Y, test_size=0.2, random_state=100)
Y_train_1 = Y_train[:,0]
Y_train_2 = Y_train[:,1]
Y_test_1 = Y_test[:,0]
Y_test_2 = Y_test[:,1]

# 创建 SVM 模型
svm = SVR(C=10, kernel='rbf', epsilon=0.1,gamma=0.1)
svm.fit(X_train,Y_train_2)

# 定义一个函数来使用 SVR 模型进行预测
def svm_predict(X):
    return svm.predict(X)

# 使用 Shap 库计算 Shap 值
explainer = shap.KernelExplainer(svm_predict, X_train)
shap_values = explainer.shap_values(X_train)
#print(shap_values)
# 显示 Shap 值的密度散点图
shap.summary_plot(shap_values,X_train,feature_names=data.columns)

```