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A universal and accurate LPMI method for calculating mismatch in heterogeneous ice nucleation†

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The interfacial correlation factor $f(m, x)$, where m refers to the interaction among ice, water and the substrate and x refers to the ratio of the critical nucleation size to the surface topography characteristic size of the substrate, plays a crucial role in the classical theory of heterogeneous ice nucleation as it significantly impacts the energy of nucleation. Generally, a smaller value of $f(m, x)$ indicates a higher propensity for ice nucleation. The degree of structural compatibility between ice and the substrate greatly influences $f(m, x)$, particularly on specific substrates. Several approaches have been proposed to calculate the lattice matching based on this idea, which allows whether a surface is favorable for nucleation to be determined. However, none of these methods adequately correlates the mismatch index with ice growth phenomena. In this paper, we embarked on a new attempt to calculate the mismatch index by combining the lattice parameter and Miller index (LPMI). Droplet freezing experiments have been carried out on α -Al₂O₃ and silicon surfaces with different Miller indices to verify the rationality of the LPMI method. Furthermore, we validated the LPMI method extensively against other works and further demonstrated its readiness, accuracy and universality for freezing problems. The results consistently show that $\delta_d = 2|d_i - d_s|/(d_i + d_s)$ with interplanar spacing more accurately predicts heterogeneous ice nucleation rates across a wide range of substrates than $\delta_1 = (a_i - a_s)/a_i$ with the lattice parameter of ice and the substrate and is more generally applicable than $\delta_{2D} = (d_i - d_s)/d_i$ with the distances between two adjacent and congener atoms on the same plane. We believe that the proposed approach will aid in the selection of substrates for promoting or inhibiting heterogeneous nucleation on a specific substrate.

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1. Introduction

Water freezing is a ubiquitous natural phenomenon and a critical process in research and industry, ranging from meteorology¹ to cryopreservation.² Pure water can remain liquid at $-38\text{ }^{\circ}\text{C}$.³ Usually, ice crystallization is triggered at a closer theoretical freezing temperature when nucleating surfaces are present, so-called heterogeneous nucleation.⁴ It is a fact that water freezing on Earth is almost a heterogeneous nucleation process involving foreign particles. Besides the above-mentioned factors, a clear understanding of heterogeneous nucleation is essential for predicting the nucleation ability of the materials⁵ and designing rational materials to promote⁶ or restrain⁷

nucleation. However, a clear and unified picture has not been drawn from these investigations because heterogeneous nucleation is an intricate interplay of the water, ice embryo, and substrate. Fortunately, the free energy of heterogeneous ice nucleation can be expressed as $\Delta G_{\text{heter}} = f(m, x)\Delta G_{\text{homo}}$ based on the classical nucleation theory (CNT), and the value of ΔG_{heter} can indicate whether nucleation is likely to occur or not. When ΔG_{homo} is determined under given conditions, $f(m, x)$ is the key parameter for predicting nucleation. The interfacial correlation factor $f(m, x)$ ⁸ in classical nucleation theory is a comprehensive parameter, where x is associated with the surface morphology and corrugation of the substrate.⁹ The parameter $m = (\gamma_{\text{sw}} - \gamma_{\text{si}})/\gamma_{\text{iw}}$ is associated with the interfacial free energy among water, ice and the substrate (the subscripts s, i, and w are the substrate, ice and water, respectively).^{8,10} For the liquid phase, γ_{sw} depends on the binding affinity, which is expressed as the intermolecular force between water and the substrate from a microscopic perspective. Stronger binding affinity leads to better wettability at the macroscopic level.¹¹ For the ice phase, the interfacial free energy γ_{si} is associated with the binding affinity of the substrate and the structural match between the substrate and nucleating

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phase.^{8,12} According to Aleksandrov,⁸ the interfacial free energy γ_{si} can be estimated as a function of the binding affinity and the structural match, $\gamma_{\text{si}}(\varphi) \approx \gamma_{\text{si}}(\alpha_0) + \frac{\varepsilon b \delta}{4\pi(1-\nu)}$, where $\gamma_{\text{si}}(\alpha_0)$ is the minimum specific interfacial free energy at a given substrate, ε , b and ν are the elastic modulus, Burgers vector, and Poisson constant, respectively, and δ describes the structural match. For a given material, γ_{si} heavily depends on the structural match between ice and the substrate. A better structural match results in a smaller $f(m, x)$ and thus a lower barrier for heterogeneous ice nucleation.¹³ Therefore, the structural match plays a vital role in heterogeneous nucleation on the specific material substrate. The most widely studied approach in structural matching is to compare the mismatch of the lattice parameters between ice and the substrate. The lattice parameter difference between ice and the substrate is commonly employed to characterize the degree of conformity between them, which is typically defined as the level of mismatch.¹⁴ In heterogeneous ice nucleation, δ denotes the difference in lattice constants of ice and the substrate. According to the consensus of existing studies,^{8,15,16} a smaller mismatch between the substrate and ice corresponds to enhanced structural matching, thereby favoring the growth of ice nuclei on the substrate. In the atmospheric and environmental science context, various substances have been identified in experiments that work as very effective ice nuclei agents, such as silver iodide,¹⁷ kaolinite,¹⁸ and corundum ($\alpha\text{-Al}_2\text{O}_3$).¹⁹ The crystal structure of these substances matches well with ice, namely, the degree of mismatch is relatively small and is favorable for ice growth. Therefore, a general calculation of the ice–substrate mismatch helps to understand icing.

In earlier studies, a parameter characterizing δ was widely expressed as²⁰

$$\delta_1 = \frac{a_i - a_s}{a_i} \times 100\% \quad (1)$$

where a_i and a_s are the lattice parameters of ice and the substrate, respectively. To further characterize the quantitative relationship between ice and substrate lattice matching, Prupacher *et al.* proposed a new function.²¹

$$\delta_2 = \frac{ma_i - na_s}{ma_i} \times 100\% \quad (2)$$

where m and n are integers that were selected to minimize δ . When using the above two methods to calculate the mismatch between crystal planes, such as the primary prism plane (10 $\bar{1}$ 0) of ice and the substrate, the lattice parameters a and c need to be calculated separately. However, the atoms are not exactly in the same plane in the basal plane (0001) and the primary prism plane (10 $\bar{1}$ 0) of the ice hexagonal (I_h). Therefore, G. N. Patey *et al.* came up with a new parameter to describe the mismatch.¹⁵

$$\delta_{2D} = \frac{|d_i - d_s|}{d_i} \times 100\% \quad (3)$$

where d_i and d_s are the distances between two adjacent and congener atoms on the same plane for ice and the substrate, respectively, as shown in Table S1 (ESI†). They are analogous lattice distances, not necessarily lattice parameters. The distances d_i and d_s are selected depending on the location of the potential or known

plane “binding site,” usually interatomic distance.¹⁵ This method is ingenious and takes into account the atomistic morphological features of the plane, which allows us to see how the inner and outer atoms on the same crystal plane of the substrate interact with ice. However, δ_{2D} requires accurate knowledge of the crystal structure and the positions of the corresponding atoms. Meanwhile, δ_{2D} is not applicable (NA) in a complex crystal structure, such as the basal plane of boehmite. More details about the mismatch are listed in Table S1 (ESI†). In summary, the existing methods for characterizing the degree of mismatch have certain limitations. For δ_1 and δ_2 , the mismatch would be calculated separately if the lattice parameters a , b , and c are not equal. Moreover, both mismatches for basal and prism planes are relatively small in δ_1 and δ_2 , but ice nucleation is observed at the basal plane, not at the prism plane.¹⁵ For δ_{2D} , d_i and d_s require a detailed and accurate crystal structure. Furthermore, δ_{2D} is not applicable when the crystal structure is complicated, such as mica, boehmite, *etc.* To address these shortcomings comprehensively, we aim to propose a more effective approach to quantify the mismatch between ice and the substrate by utilizing interplanar spacing (which combines lattice parameters with Miller indices). Furthermore, experimental validation was conducted to support this expression of mismatch while comparing it with molecular dynamics simulations carried out by other researchers. The comparison demonstrates that our proposed δ_d aligns well with both experimental observations and numerical simulation results. We firmly believe that this methodology will greatly contribute to the analysis of heterogeneous ice nucleation.

2. Definition and method

The mismatch denoted by interplanar spacing can be expressed as²²

$$\delta_d = 2|d_i - d_s|/(d_i + d_s) \quad (4)$$

where d_i and d_s are the interplanar spacing of the crystal panel in ice and the substrate, respectively. Interplanar spacing d_{hkl} between adjacent planes with Miller indices (hkl) is defined as the distance between the first plane and a parallel plane passing through the origin. Interplanar spacing can be visualized in the examples shown in Fig. 1, an example of a simple cubic lattice ($a = b = c$, $\alpha = \beta = \gamma = 90^\circ$) and a hexagonal lattice ($a = b$, $\alpha = \beta = 90^\circ$, $\gamma = 120^\circ$). The formulas of the interplanar spacing for different crystal systems are listed in Table S1 (ESI†) for reference.

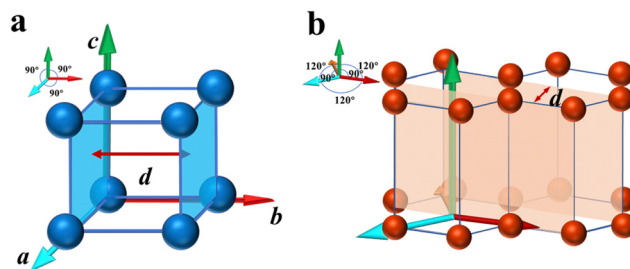


Fig. 1 Examples of some interplanar spacings: (a) (100) plane in a cubic lattice and (b) (10 $\bar{1}$ 0) plane in a hexagonal lattice.

Each crystal has a set of interplanar spacings of different sizes, which are a function of the lattice parameter and Miller indices. The interplanar spacing is calculated as

$$\frac{1}{d_{(hkl)}^2} = \left| \begin{array}{ccc} \frac{h}{a} & \cos \gamma & \cos \beta \\ \frac{h}{a} & \frac{k}{b} & 1 \\ \frac{l}{c} & \cos \alpha & 1 \end{array} \right|^2 + \left| \begin{array}{ccc} 1 & \frac{h}{a} & \cos \beta \\ \cos \gamma & \frac{k}{b} & \cos \alpha \\ \cos \beta & \frac{l}{c} & 1 \end{array} \right|^2 + \left| \begin{array}{ccc} 1 & \cos \gamma & \frac{h}{a} \\ \cos \gamma & 1 & \frac{k}{b} \\ \cos \beta & \cos \alpha & \frac{l}{c} \end{array} \right|^2 + \left| \begin{array}{ccc} 1 & \cos \gamma & \cos \beta \\ \cos \gamma & 1 & \cos \alpha \\ \cos \beta & \cos \alpha & 1 \end{array} \right|^2 \quad (5)$$

where a , b , c , α , β , and γ are the lattice parameters and (h, k, l) are the Miller indices of the crystal plane. The increase in Miller indices would result in a decrease in the interplanar spacing.²³ To verify the LPMI method, we performed droplet freezing experiments on the surfaces of monocrystalline silicon (cubic structure) and α -Al₂O₃ (hexagonal structure) with different Miller indices. The experimental details are elaborated in the ESI†. Moreover, we compared the different mismatch expressions with other results of heterogeneous ice nucleation in molecular dynamics. Unit cell parameters for crystal structures in this paper are shown in Table S2 (ESI†). The results are discussed in the following section.

3. Results and discussion

3.1. Validation of the mismatch by experiments

The freezing process of droplets on monocrystalline silicon wafers and α -Al₂O₃ flakes is depicted in Fig. 2(a). Upon freezing, the ice–water interface progresses from the bottom to the top of the droplet. The impact of substrate mismatch on droplet freezing was analyzed in terms of the freezing delay time, which is a parameter conventionally considered as the reciprocal of the nucleation rate.⁸ The freezing delay time refers to the duration required for the droplet to freeze after reaching the target temperature on the substrate. The freezing delay time τ_c for each substrate is the mean of the measured value from 15 experiments. Fig. 2(b) illustrates that the freezing delay time is shorter on the monocrystalline silicon substrate (110) compared to (100) and (111) surfaces. Concerning α -Al₂O₃, the freezing delay time is longer on the C -plane (0001) in comparison to the M -plane (10 $\bar{1}0$) and A -plane (11 $\bar{2}0$). Table 1 presents the mismatch between the substrate and ice. The crystalline structure of ice was assumed to be hexagonal since the minimum temperature in the experiments was -15°C , and cubic ice remains stable under extreme conditions such as low temperatures ($<190\text{ K}$)²⁴ and under the influence of a strong external electric field.²⁵

The minimum δ_d of 1.9% observed between silicon (110) and the primary prism plane of I_h corresponds to the minimum freezing delay time observed on silicon (110). On α -alumina, the freezing delay time on the (0001) plane is longer compared to the (1 $\bar{1}0$) and (11 $\bar{2}0$) planes. The minimum δ_d between ice and α -Al₂O₃ was 5.2%. A smaller mismatch leads to a lower interfacial correlation factor $f(m, x)$, thereby reducing the barrier for heterogeneous ice nucleation. Consequently, the freezing delay time is shorter under consistent conditions. In other words, the freezing delay time on different Miller indices confirms the accuracy of the mismatch (determined by the interplanar spacing). Molecular dynamics (MD) studies on the first hydration layer^{13,26,27} reveal that when the lattice mismatch is small, the water dipole distribution closely resembles that of bulk ice, resulting in stronger water–water interactions. Conversely, larger deviations caused by a larger lattice mismatch accelerate the decay of dipole orientation. This leads to higher potential energy and a less stable hydrogen bond network.¹³ Therefore, a smaller δ_d value correlates with a shorter freezing delay time. Furthermore, the findings of other MD simulations provide additional evidence to support the significance of the mismatch δ_d .

3.2. Validation of the mismatch by molecular dynamics

In this section, we present the results of heterogeneous nucleation on β -AgI,^{15,16} kaolinite,^{15,28} mica,^{15,29} gibbsite,^{15,30} and hematite^{15,31} in existing studies. Complete information about the substrates can be found in Table S2 (ESI†). Furthermore, we compared the values of δ_d with δ_1 , δ_2 , and δ_{2D} as summarized in Table 2. The results further support the generality of δ_d , which considers Miller indices and lattice parameters. As shown in Table 2, nucleation occurred on β -AgI, kaolinite (prime plane), mica (basal plane), and PbI₂ (basal plane) substrates, but not on gibbsite, hematite, and boehmite substrates within the same simulated time scale. A smaller δ value indicates a higher similarity between the crystal structure of the substrate and that of ice, indicating a greater likelihood of ice formation. Ice nucleation was observed on both the basal and prism planes of β -AgI due to their good lattice matching with ice (I_h).^{15,16} By comparing the different δ values, we find that δ_d (by the interplanar spacing) is in good agreement with the MD simulation results. δ_1 (by lattice parameter) and δ_2 (by lattice parameter) partially align with the MD simulation results. However, there are discrepancies in the case of kaolinite nucleating the (10 $\bar{1}0$) prism plane of ice (I_h).^{15,32} The δ_1 values for both the basal (14.29%, 98.4%) and prism (14.29%, 2.23%) faces are relatively large. Additionally, for δ_2 , the match between gibbsite and ice in the prism plane (3.6%, 3.69%) is better than that of kaolinite and ice in the prism plane (5.16%, 0.8%), yet no nucleation of gibbsite is observed in simulations.¹⁹ Both δ_{2D} (by analogous lattice distance) and δ_d are consistent with the observations of kaolinite and gibbsite, but δ_{2D} is not applicable in situations involving crystals with complex structures such as mica, where defining the analogous lattice distance in the prism plane is challenging.

We further compared the mismatch among artificial crystals based on the previous comparisons of actual crystal structures. Scaling α -Al₂O₃ eliminates the mismatch for the basal plane, resulting in ice nucleation on the surface. The four measures of

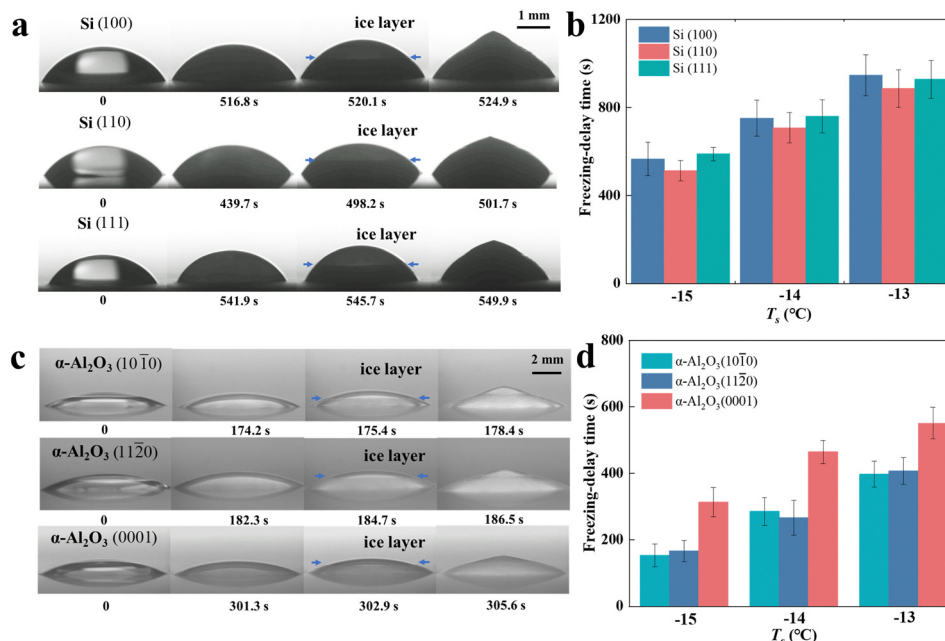


Fig. 2 (a) A droplet with a volume of 10 μ L freezes on a silicon substrate at $T_s = -15$ °C. (b) The freezing delay time of the droplet on silicon substrates. (c) A droplet with a volume of 10 μ L freezes on α -Al₂O₃ substrates at $T_s = -15$ °C. (d) The freezing delay time of the droplet on α -Al₂O₃ substrates. The smaller the δ_d , the shorter the freezing delay time. The δ_d values for Si (100), α -Al₂O₃ (1010) and α -Al₂O₃ (1120) are 1.9%, 5.2% and 5.2%, which are smaller than other crystal planes. Thus, freezing occurs easily on these substrates above. The error bar is the error of the mean.

Table 1 Mismatch δ_d between the substrate and ice

Crystal	Silicon (100) (%)	Silicon (110) (%)	Silicon (111) (%)	α -Al ₂ O ₃ (1010) (%)	α -Al ₂ O ₃ (1120) (%)	α -Al ₂ O ₃ (0001) (%)
Ice (I _h) (1010)	32.5	1.9	26.1	5.2	48.7	30.3
Ice (I _h) (1120)	82.5	51.8	32.4	58.4	5.2	62.9
Ice (I _h) (0001)	30.1	62.8	80.4	56.3	102.2	51.7

δ align well with the observations. Conversely, scaling α -Al₂O₃ to hematite (referred to as AsH) increases the mismatch for the basal plane, leading to the absence of ice nucleation in both the basal and prism planes. In this case, δ_1 , δ_2 , and δ_d consistently correspond to the observations, except for a relatively small δ_{2D} value (3.3%, 0.58%) for the prism plane. Furthermore, scaling hematite to ice (referred to as HsI) eliminates the mismatch at the basal plane, resulting in ice nucleation under specific force fields. All measures of δ align well with the simulation results. By comparing different mismatches on real and fictional crystals, we find that δ_1 ,

δ_2 , δ_{2D} , and δ_d are to some extent consistent with the observations. However, δ_{2D} requires precise knowledge of the crystal structure and the positions of the corresponding atoms, making it inapplicable for certain complex crystal structures. Additionally, for different water-potentials (TIP4P/Ice and six-site) and the different water-substrate interactions (CLAFF and LJ), the result of ice nucleation in the same time interval is the same, which means that the potential and the water-substrate interaction have little influence on ice nucleation. Therefore, δ_d , which considers crystal parameters and Miller indices, provides a better

Table 2 δ_1 , δ_2 , δ_{2D} , and δ_d lattice mismatches of ice (I_h) for the surface considered in the molecular dynamics simulation

Surface label	$ \delta_1 $ (%)		$ \delta_2 $ (%)		$ \delta_{2D} $ (%)		$ \delta_d $ (%)		Nucleation
	Basal	Prism	Basal	Prism	Basal	Prism	Basal	Prism	
β -AgI ^{15,16}	2.08	(2.08, 2.23)	2.04	(2.04, 2.23)	2.04	(2.04, 2.23)	1.87	1.37	Y
Kaolinite ^{15,28}	(14.29, 98.4)	(14.29, 2.23)	(14.29, 0.8)	(5.16, 0.8)	14.6	(0.78, 1.85)	11.01	5.65	Y (prism)
Mica ^{15,29}	(15.2, 100.3)	(15.2, 173.7)	(15.2, 0.19)	(6.16, 0.19)	(0.2, 5.93)	NA	4.53	21.74	Y (basal)
PbI ₂ ¹⁵	1.13	(1.13, 6.95)	1.13	(1.13, 6.95)	1.13	NA	2.52	55.47	Y (basal)
Gibbsite ^{15,30}	(92.8, 12.7)	(92.8, 32.5)	(3.6, 12.7)	(3.6, 3.69)	14.1	(6.2, 0.59)	27.94	16.68	N
Hematite ^{15,31}	11.6	(11.6, 87.17)	11.6	(11.6, 2.71)	11.6	(3.3, 0.58)	124.67	11.01	N
Boehmite ¹⁵	(26.5, 17)	(26.5, 1.84)	(26.5, 17)	(26.5, 1.84)	NA	(17.09, 15.48)	116.6	33.78	N
AsI ¹⁵	0	51.7	0	(0, 8)	0	(13.4, 10.94)	0	51.7	Y (basal)
AsH ¹⁵	11.6	87.17	11.6	(11.6, 2.71)	11.6	(3.3, 0.58)	124.6	12.01	N
HsI ¹⁵	0	(0, 87.17)	0	(0, 8)	0	(13.4, 10.94)	0	51.7	Y (basal)

description of the mismatch between ice and crystals in heterogeneous nucleation.

3.3. Regression analyses of mismatch

We conducted a regression analysis on various methods for calculating mismatches. According to Classical Nucleation Theory, the freezing delay time can be calculated by eqn (6),³³

$$\tau_{\text{in}} = \frac{1}{V\Phi K(T)e^{-\frac{f(m,x)\Delta G_{\text{homo}}}{K_{\text{B}}T}}} \quad (6)$$

where V represents the droplet volume, Φ is the fraction area of the water and substrate, $K(T)$ denotes a factor of the diffusion flux of water molecules through the ice–water interface, usually its value is $10^{25} \text{ cm}^{-2} \text{ s}^{-1}$, K_{B} denotes the Boltzmann constant, and ΔG_{homo} is the nucleation barrier, which can be expressed as

$$\Delta G_{\text{homo}} = -\frac{4\pi r^3}{3}\Delta G_{\text{v}} + 4\pi r^2\gamma_{\text{iw}} \quad (7)$$

where $\Delta G_{\text{v}} = \Delta H_{\text{m,f}}(T_{\text{f}} - T)/T_{\text{f}}$, $\Delta H_{\text{m,f}}$ and T_{f} denote melting enthalpy and melting temperature, respectively. $f(m,x)$ denotes the interfacial correlation factor, which can be expressed as

$$f(m,x) = 1 + \left(\frac{1-mx}{g}\right)^3 + x^3 \left[2 - 3\left(\frac{x-m}{g}\right) + \left(\frac{x-m}{g}\right)^2\right] + 3mx^3\left(\frac{x-m}{g} - 1\right) \quad (8)$$

$$x = R/r_{\text{c}}, m = (\gamma_{\text{sw}} - \gamma_{\text{si}})/\gamma_{\text{iw}}, g = (1 + x^2 - 2mx)^{\frac{1}{2}} \quad (9)$$

where R and r_{c} are the mean surface roughness and the radius of the critical nucleus, respectively. Additionally, the minimum

specific interfacial free energy $\gamma_{\text{si}}(\alpha_0)$ of a given substrate is expressed as³⁴

$$\gamma_{\text{si}}(\alpha_0) = |\varepsilon_0|n/2 \quad (10)$$

where ε_0 is the interaction between ice and a solid, and n is the number of neighboring molecules. Thus, the regression analyses of experimental τ_{c} , simulated τ_{c} and predicted τ_{c} are displayed in Fig. 3. Since δ_1 and δ_2 are similar, we take δ_1 as an example for analysis.

In all cases, δ describes the mismatch between the substrates and the ice (I_{h}). We report only absolute values ($|\delta|$) of the mismatch. Only one value in the table indicates that the lattice parameters of the corresponding crystal planes are equal. For example, a and b of the basal plane in β -AgI are equal. Y means that ice nucleation occurred spontaneously on the substrate, and the nucleation plane is given. N means that no ice nucleation was observed within the simulation time scale. Nucleation occurs at both the basal and prism planes of β -AgI.

Fig. 3 shows the regression analyses of δ_{d} than δ_1 or $\delta_{2\text{D}}$. In panels (a), (c), and (e), the y-axis signifies times ascertained through experimental measurements, whereas in panels (b), (d), and (f), it delineates the freezing delay time derived from MD simulations^{15,16} as reported in publications. The x-axis embodies the theoretical nucleation time, which is deduced by integrating nucleation temperatures and specific surface characteristics into eqn (6). The line $y = x$ symbolizes instances where experimental or simulation values equal theoretical predictions, while the positions of points within the figure elucidate deviations between experimental or simulated outcomes and their corresponding theoretical expectations. Fig. 3 demonstrates that the predicted freezing delay time (τ_{c}) based on δ_{d} has an error of less than 20% when compared to

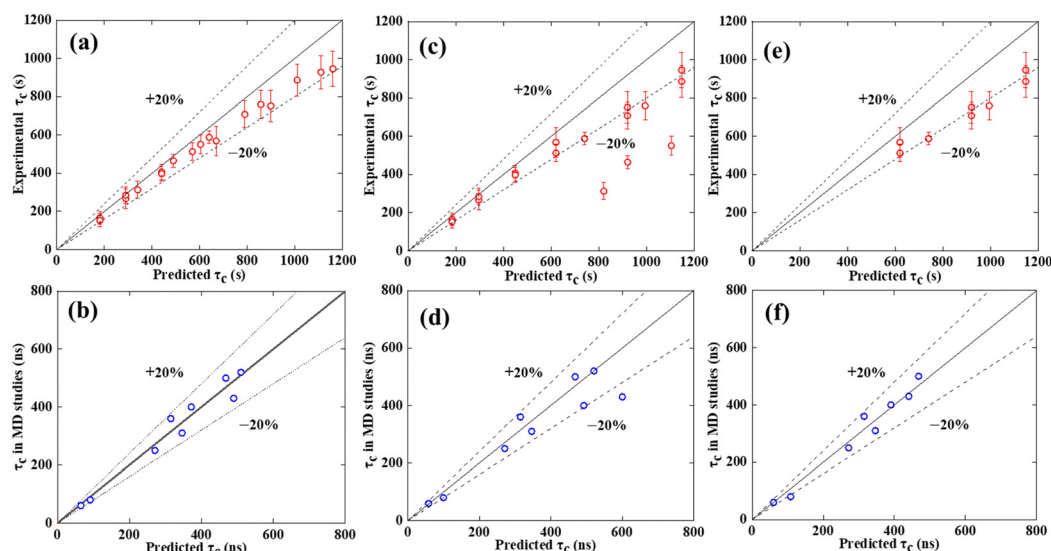


Fig. 3 Regression analyses. (a), (c) and (e) Show regression analyses based on experimental data. (b), (d) and (f) Show regression analyses based on simulated data. The data from the simulation is listed in Table S4 (ESI†). (a) and (b) Show regression analyses based on δ_{d} . (c) and (d) Show regression analysis based on δ_1 . (e) and (f) show regression analyses based on $\delta_{2\text{D}}$. The mean freezing delay time is more accurately predicted across a wide range of substrates by δ_{d} than δ_1 or $\delta_{2\text{D}}$. The red points represent the mean values of experimental measurements, with error bars indicating the standard error of the mean; the blue points derive from the freezing delay time of MD simulations^{15,16} in existing studies.

experimental and simulated τ_c . However, when using δ_1 for prediction, the error exceeds 20% in several trials, indicating poor accuracy. Although the predicted τ_c based on δ_{2D} is within the 20% error range, determining the analogous lattice distance proves challenging, especially for complex crystal structures like $\alpha\text{-Al}_2\text{O}_3$. Consequently, τ_c cannot be reliably predicted using δ_{2D} . Hence, the mismatch measure δ_d is more accurate in predicting the freezing delay time and exhibits good universality.

In summary, the analysis establishes the relationship between different representations of δ and nucleation from a crystal structure perspective. The extension of the mismatch measure δ_d from one-dimensional lattice parameters (a , b , and c) to crystal planes with various Miller indices has demonstrated its superior accuracy in both experimental and numerical simulations. This approach only requires basic information such as crystal type, lattice parameters, and Miller indices of the crystal plane to obtain interplanar spacings. By comparing the mismatch between the substrate and ice, it becomes possible to rapidly select and design structures for promoting or inhibiting ice nucleation in specific materials. However, heterogeneous ice nucleation is a complex interplay between ice and the substrate, involving atomic-level insights. Achieving a more accurate description of heterogeneous ice nucleation would require further systematic and microscopic analysis.

4. Conclusions

In this study, we performed a comparison of δ_1 (based on lattice parameters), δ_2 (based on lattice parameters), δ_{2D} (based on analogous lattice distances), and δ_d (based on interplanar spacings) through experiments and simulations. Our results demonstrate that δ_d , which takes into account the lattice parameter and Miller index, is more accurate than δ_1 and δ_2 , as well as more universal than δ_{2D} . δ_d provides a more comprehensive description of the mismatch in heterogeneous ice nucleation. It is crucial to have a proper match between the crystal structure and ice for successful ice nucleation. However, it should be noted that heterogeneous ice nucleation is a complex phenomenon requiring further systematic and microscopic analysis. The methods utilized in this paper are expected to contribute to a broader understanding of heterogeneous nucleation.

Conflicts of interest

There are no conflicts to declare.

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