

Cite this: *J. Mater. Chem. A*, 2020, **8**, 17747Efficient Kr/Xe separation from triangular g-C₃N₄ nanopores, a simulation study†Mohammad Tohidi Vahdat,^{ab} Davide Campi,^b Nicola Colonna,^{bc}
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Poly(triazine imide) or PTI is a promising material for molecular sieving membranes, thanks to its atom-thick ordered lattice with an extremely high density (1.6×10^{14} pores per cm²) of triangular-shaped nanopores of ~ 0.34 nm diameter. Here, we investigate the application of PTI nanopores in the purification of Kr from Xe to reduce the storage volume of the mixture of ⁸⁵Kr/Xe. Using van der Waals density-functional theory (vdW-DFT) calculations, benchmarked against the random phase approximation (RPA), we calculate the potential energy profiles for Kr and Xe across the nanopores. For each gas, starting from the RPA potential-energy profile, the force-field parameters to be used in the classical restrained molecular dynamics framework are trained to calculate the Helmholtz free energy barrier as a function of temperature, and therefore, the corresponding entropic loss. Overall, due to the much higher activation energy from the adsorbed state in Xe (17.61 and 42.10 kJ per mole for Kr and Xe, respectively), a large Kr/Xe separation selectivity is postulated from the PTI membrane. Furthermore, the combination of the atom-thick PTI lattice and high pore density leads to extremely large yet selective permeances for Kr. For example, a Kr permeance of 1000 gas permeation units (GPU) accompanying a large Kr/Xe selectivity ($>10\,000$) is calculated at 25 °C, which is significantly better than that of the state-of-the-art membranes for Kr/Xe separation, making PTI-based membranes a leading candidate for processing the hazardous waste of ⁸⁵Kr/Xe mixture.

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1. Introduction

The development of materials to enable energy-efficient and straightforward processes can provide a solution to complex challenges in chemical separations. For example, an important separation challenge is purification of ⁸⁵Kr from a 10/90 mixture of ⁸⁵Kr/Xe, currently stored as hazardous waste.^{1,2} Addressing this would reduce by ten times the storage volume of ⁸⁵Kr, and therefore reduce also associated risks.^{3,4} Moreover, Xe recovered in a pure form or with negligible ⁸⁵Kr contamination can find applications in flash and arc lamps, as propellants for ion thrusters in satellites, and as general anesthetics.^{5–8}

Currently, the separation of non-hazardous Kr/Xe mixtures is carried out predominantly by cryogenic distillation, relying on the relative volatility of liquid Kr and Xe.⁷ When handling hazardous wastes from power plants, such as ⁸⁵Kr/Xe, reducing the process

footprint is desirable, and membrane-based separation becomes very attractive. Membrane separation is one of the simplest industrial processes, requiring minimal maintenance. Moreover, by developing membranes yielding a high Kr permeance (defined as pressure-normalized flux), one can further reduce the membrane area needed. Nanoporous materials, hosting pores with a gap in the electron-density appropriate for sieving Kr (that has a kinetic diameter of 3.60 Å) from Xe (kinetic diameter 3.96 Å) can achieve high separation selectivity.^{9–12} Recent progress in synthesizing atom-thick membranes, for example by using single-layer graphene, has made it possible to realize large but selective gas permeance.^{13–16} However, for effective sieving of Kr from Xe, two-dimensional nanoporous structures hosting uniform nanopores would be highly attractive. Such materials can be identified by screening databases of nanoporous materials.^{17–19} A review of the literature indicates that carbon nitride materials can be very attractive to reach this goal.^{20–22} Among them, g-C₃N₄ is rapidly emerging for application in catalysis,^{23,24} energy storage,²⁵ sensors,²⁶ and molecular separations.^{27,28} The in-plane structure of g-C₃N₄ hosts either a 1,3,5-triazine or a heptazine (1,3,4,6,7,9-b-heptaazaphenylene) building unit.²⁹ In particular, poly(triazine imide) or PTI is quite attractive for gas separation because of its highly ordered nanoporous structure, composed of imide-bridged triazine units which form a triangular sub-nanometer-sized pore (Fig. 1).²⁷ A remarkable aspect of the mesh-like two-dimensional

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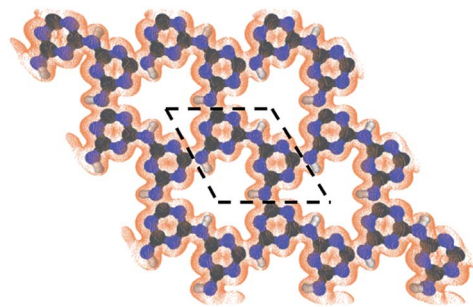


Fig. 1 The structure of the nanoporous PTI lattice and its electron density isosurface (iso-value of $0.03 \text{ e} \text{ \AA}^{-3}$) revealing an electron-density gap of $\sim 3.4 \text{ \AA}$. The dashed lines show the PTI unit cell.

(2d) structure of PTI is its high pore density, 1.6×10^{14} pores per cm^2 , much larger than those typically realized in other 2d nanoporous membranes (such as nanoporous single-layer graphene),³⁰ making it promising for obtaining large permeances. The chemical and thermal robustness of PTI is also attractive for separation under harsh conditions.³¹

Here, we investigate the performance of PTI nanopores in such useful membrane-based separation processes, employing vdW-DFT calculations, carefully benchmarked against exact exchange (EXX) plus RPA calculations, and molecular dynamics (MD) simulations. We indeed find that PTI nanopores can be very attractive for Kr/Xe separation, with selectivity exceeding 10 000 at room temperature. This is accompanied by a high Kr permeance of 1000 gas permeation units or GPU ($1 \text{ GPU} = 3.35 \times 10^{-10} \text{ mol m}^{-2} \text{ s}^{-1} \text{ Pa}^{-1}$).

2. Computational methods

2.1 DFT

To investigate the potential energy surface of Kr and Xe, vdW-DFT calculations are performed using the Quantum ESPRESSO distribution.^{32,33} A 2×2 supercell is used to ensure the decoupling of in-plane molecule–molecule interactions. Integration over the Brillouin zone is carried out with a uniform $6 \times 6 \times 1$ k -point grid. A vacuum region of 40 \AA is employed in the z direction, to avoid interactions among periodic replicas. An energy cutoff of 60 Ry is used for the plane wave expansion of the wavefunctions. A kinetic energy cutoff of 480 Ry on the charge is used together with ultra-soft pseudopotentials.^{34,35} In order to carefully take into account the influence of the van der Waals (vdW) interactions, five different vdW approximations are tested: vdW-DF2,³⁶ Grimme-D2,³⁷ Grimme-D3,³⁸ the revised version of the Vydrov van Voorhis functional (rVV10),^{39,40} and Tkatchenko–Scheffler (TS).⁴¹ Since TS is not implemented in Quantum ESPRESSO when using ultrasoft pseudopotentials, calculations with this vdW approximation are carried out with norm-conserving pseudopotentials.⁴² In this case, an energy cutoff of 100 Ry is used for the wave functions.

2.2 EXX/RPA

Using the implementation of the ACFDT formalism available in the Quantum ESPRESSO^{33,43,44} distribution, we compute the

total energies within exact-exchange plus the random-phase approximation. From a practical point of view, the main limitation of EXX/RPA is its computational cost, which is considerably higher than that of conventional LDA or GGA calculations. In order to use this computational-demanding method efficiently, some convergence tests are carried out (Fig. S2†); the PES of gases is studied as a function of relaxation of the gas (Fig. S2a†), supercell size (Fig. S2b†), k -point sampling (Fig. S2c†), pseudopotentials (Fig. S2d†), and cutoff values (Fig. S2e†). The PES is not affected significantly when the supercell sizes and cutoff values are reduced, and when the pseudopotential is changed. Overall, it can be concluded that EXX/RPA calculations can be performed for the systems at hand at the Γ point using a PTI unit cell size and norm-conserving pseudopotentials.

2.3 Calculation of free energy profile

All classical MD simulations are carried out with the large-scale atomic/molecular massively parallel simulator (LAMMPS) package.⁴⁵ The interatomic interactions are described by the general AMBER force field (GAFF).⁴⁶ Partial atomic charges on PTI are considered using the AM1-BCC method.⁴⁷ Kr and Xe are modelled as single center Lennard-Jones (LJ) particles. The LJ parameters used to consider non-bonded interactions, are tabulated in ESI Tables 1 and 2.† The particle–particle–particle–mesh (PPPM) is implemented to consider long-range electrostatic interactions⁴⁸ and the vdW interactions are calculated within a cutoff distance of 12 \AA .

The dimensions of the simulation box in the x , y , and z directions are 25.9 , 31.3 , and 40.0 \AA , respectively. The PTI lattice is placed in the x – y plane and its atoms are allowed to move in all directions except the atoms on the external edges, which were kept fixed to avoid any rigid shift of the PTI monolayer due to interactions with gases. In this way, it is ensured that the PTI sheet would not shift due to the interaction with Kr and Xe. Periodic boundary conditions are employed in all directions. For each simulation, the total energy of the system is minimized for 100 000 steps. Then, the system is equilibrated in a constant number, volume and temperature (NVT) ensemble for 1 ns with a timestep of 1 fs . A Nose–Hoover thermostat is employed to control the temperature of the system.^{49,50}

Then, in the restrained MD framework, umbrella sampling⁵¹ is used to gather enough statistics in a reasonable period of time. To overcome the energy barrier and configurations which might be inaccessible in a reasonable amount of time, a harmonic restrain with the force constant of $5 \text{ kcal mol}^{-1} \text{ \AA}^{-1}$ is used every 0.5 \AA . Sampling is performed for 4 ns , and trajectories are collected every picosecond. Finally, the weighted histogram analysis method (WHAM) is used to unbias and combine the distribution of gases along the translocation trajectory function and construct the Helmholtz free energy profile.⁵²

2.4 Calculations of the concentration of occupied pores

In order to calculate the concentration of occupied pores, $[C_{\text{ox}}]$, a simulation box with the dimensions of 25.9 , 31.3 , and 120.0 \AA



in the x , y , and z directions, respectively, is defined (Fig. S3†). We position the PTI lattice in the x - y plane at $z = 60$ Å, dividing the simulation box into two chambers (feed and permeate) of equal volume. In order to prevent the escape of the gas into the vacuum region due to the periodicity, a graphene layer is positioned at $z = 0$ as a physical barrier. To accelerate the MD simulations and gather enough statistics in the 7 ns-simulations, a gas pressure of 50 bar is used corresponding to 56 Kr or Xe atoms in the feed chamber while the permeate chamber is initially empty. In order to prevent vertical displacement of graphene and PTI, their coordinates are fixed. The trajectories of gas atoms are collected every ten picoseconds to keep track of the number of adsorbed atoms. At the end of simulations, the average number of atoms in the adsorption region is obtained to calculate $[C_{\text{ox}}]$ (Fig. S3†).

3. Results and discussion

First, the PTI lattice is relaxed to its minimum energy configuration. The optimized lattice parameters are 8.64, 8.65, 8.65, 8.64, and 8.67 Å with vdW-DF2, Grimme-D2, TS, Grimme-D3, and rVV10, respectively, in good agreement with each other. To give a heuristic representation of the electron-density gap, the iso-surface of PTI at a small value of 0.03 e Å^{-3} is generated (Fig. 1).

Table 1 The adsorption energies, the adsorption heights, and the activation barriers of Kr and Xe on the PTI lattice calculated with five different van der Waals approximations

	E_{ads} (eV)		Adsorption height (Å)		E_{act} (eV)	
	Kr	Xe	Kr	Xe	Kr	Xe
vdW-DF2	−0.06	−0.07	2.0	2.5	0.22	0.46
Grimme-D3	−0.15	−0.18	1.7	2.0	0.19	0.43
TS	−0.14	−0.17	1.8	2.0	0.18	0.41
Grimme-D2	−0.16	−0.21	1.7	1.9	0.17	0.40
rVV10	−0.17	−0.22	1.6	1.8	0.14	0.37

An equilateral-triangle-shaped nanopore can be observed, with each side measuring 9.15 Å. The nanopore can host a circle with a diameter of 3.40 Å. This pore size is attractive for the separation of light gases based on their size, especially Kr from Xe by the size-sieving mechanism because the kinetic diameter of Kr is closer to the nanopore size. To investigate the adsorption behavior of gases on the porous PTI monolayer, a constrained relaxation is carried out. The gas is placed at several positions away from the sheet and the mass center of the gas is fixed in the z direction (perpendicular to the sheet); the gas still has the freedom to relax in x and y directions to minimize the potential energy at a given distance from the PTI monolayer in the z direction. Since the potential energy surface (PES) for Kr and Xe are simple enough (Kr and Xe are just one single atom), it is not required to perform Nudge elastic band calculations. The convergence threshold on the total energy is 2×10^{-6} Ry and this criterion on forces is equal to 10^{-4} Ry/a.u. The minimum PES for different gases passing through the PTI monolayer is explored by calculating the interactions energy with the PTI film as:

$$E_{\text{int}} = E_{\text{molecule+PTI}} - E_{\text{molecule}} - E_{\text{PTI}} \quad (1)$$

where $E_{\text{molecule+PTI}}$ is the total energy of the system when the gas interacts with PTI at different distances. E_{molecule} and E_{PTI} are the energy of the gas and the PTI monolayer, respectively, when they are isolated and do not interact.

Fig. 2 shows the PES for Kr and Xe obtained using the five considered vdW approximations. In order to make sure that the PTI monolayer does not shift rigidly due to the interactions with gases, the atoms on the edge of the PTI plane are kept fixed. It is worth emphasizing that during these calculations all the PTI monolayer atoms (except the atoms on the edge of the plane) are free to move and relax. When the PTI lattice is treated as a rigid lattice, the PES profile did not change by a considerable amount (Fig. S1†), indicating that the PTI lattice is reasonably rigid. The results indicate that the Kr adsorption energy in the middle of the pore (E_{ads} , defined as the maximum attractive interaction energy when $z \neq 0$) is in the range of -0.06 to -0.17 eV, and the

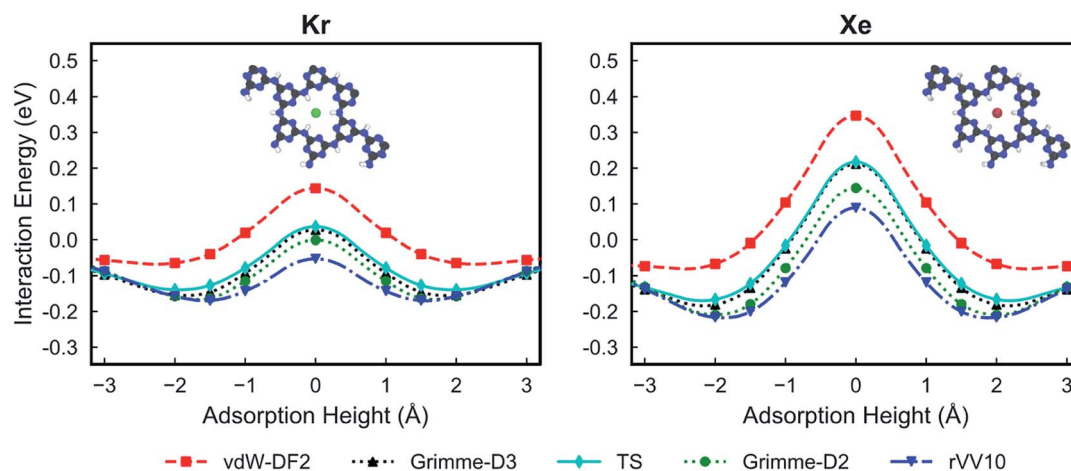


Fig. 2 Potential energy surfaces of Kr (left) and Xe (right) for the PTI lattice using different van der Waals functionals: vdW-DF2,³² Grimme-D2,³³ Grimme-D3,³⁴ rVV10,^{35,36} and TS.³⁷ A height of 0 Å corresponds to the plane of PTI.



adsorption height from the PTI plane is in the range of 1.60–2.00 Å. Xe adsorption energy is in the range of –0.07 to –0.22 eV, and the adsorption height in the range of 1.80 to 2.50 Å. The resulting activation barriers to translocate the nanopore (E_{act} , defined as the difference in the interaction energy at $z = 0$ and E_{ads}) for Kr and Xe are in the range of 0.14 to 0.22 eV, and 0.37 to 0.46 eV, respectively. In all cases, E_{act} for Xe is higher than that for Kr (Table 1). The discrepancy between the vdW approximations is more apparent in E_{ads} than in E_{act} .

To understand the effects of the vdW approximations on the sieving performance, the Arrhenius relationship for activation energies is used to calculate Kr/Xe selectivities, assuming that the translocation of the PTI nanopore would be the rate limiting step after the gases are adsorbed on the PTI lattice.⁵³ The selectivity based on the Arrhenius equation is defined in eqn (2):

$$S_{\text{Kr/Xe}} = \frac{r_{\text{Kr}}}{r_{\text{Xe}}} = \frac{A_{\text{Kr}} \exp(-E_{\text{Kr}}/RT)}{A_{\text{Xe}} \exp(-E_{\text{Xe}}/RT)} \quad (2)$$

where r is the rate of translocation and A_{Kr} and A_{Xe} are the prefactors for the Arrhenius relationship for Kr and Xe,

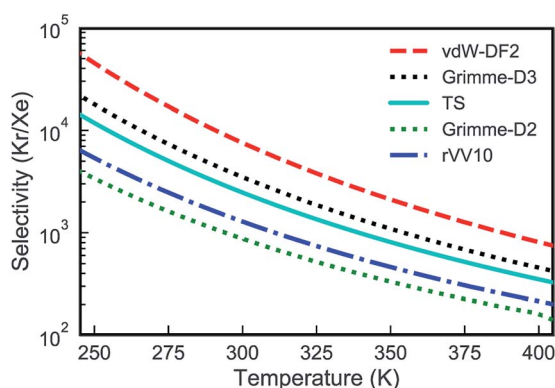


Fig. 3 Kr/Xe selectivities from the PTI lattice as a function of temperature calculated with several van der Waals approximations.

respectively. E_{Kr} and E_{Xe} refer to the apparent activation energies (defined as $E_{\text{ads}} + E_{\text{act}}$) for Kr and Xe, respectively; R is the universal gas constant and T is the temperature. Assuming that A_{Kr} and A_{Xe} are within the same order of magnitude,⁵⁴ an approximate gas pair selectivity within the temperature range of 250–400 K can be calculated (see Fig. 3).

Overall, all vdW approximations predict a high Kr/Xe separation selectivity, above 100 in all cases in the temperature range of 250–400 K (Fig. 3). Selectivities increase when the temperature is reduced, since Xe encounters a higher activation energy than Kr.

Despite the fact that all the vdW-DFT calculations give the same qualitative result, selectivities obtained with different functionals can differ quite dramatically, with room-temperature values spanning a wide range that goes from 875 for Grimme-D2 to 7550 for vdW-DF2 (Fig. 3). Properly capturing vdW interactions with density-functional theory is complex and a relatively recent endeavor that still represents a challenging problem.⁵⁵ In order to have a higher-level description of van der Waals interactions, crucial to determine the gas-PTI monolayer interaction, and to have a qualitative estimation of the selectivity, for the first time, we evaluated the energetics of the PTI complexes in the framework of adiabatic-connection fluctuation dissipation theory (ACFDT),^{56,57} allowing a better understanding of the performances of different vdW approximations for these systems. The EXX/RPA method has been proved to correctly capture vdW correlations,^{58,59} and *e.g.*, the results of physisorption binding energies between different types of molecules and surfaces show an excellent agreement with the values measured by experiments.^{60–66}

The RPA/EXX results reveal that the Grimme-D2, Grimme-D3 and TS approximations yield a good agreement PES (Fig. 4). The calculations show that vdW-DF2 overestimates the adsorption energy and on the other hand rVV10 underestimates the adsorption energy.

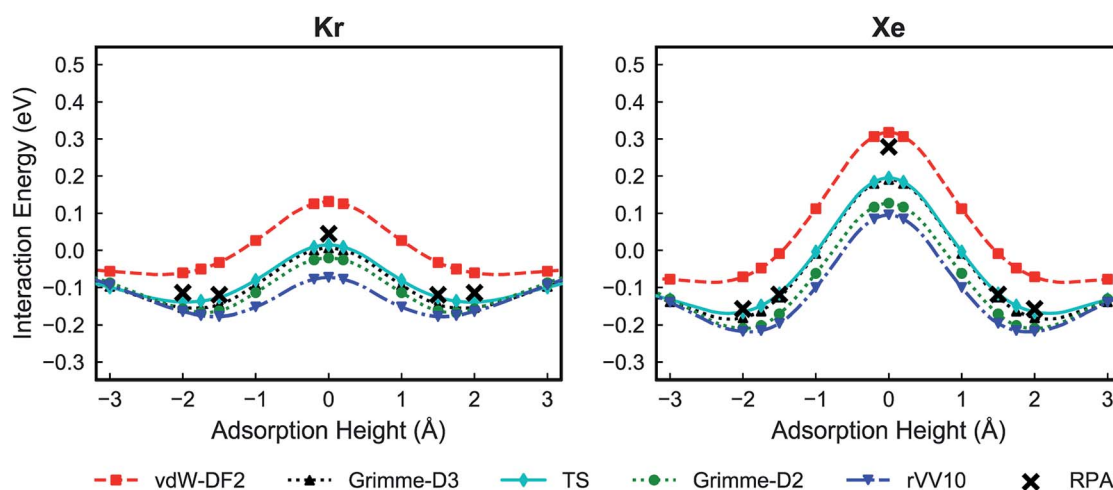


Fig. 4 Potential energy surfaces of Kr (left) and Xe (right) with several van der Waals approximations in comparison to the results calculated by EXX/RPA. Here, potential energy surfaces calculated with vdW-DFT are obtained with same parameters as EXX/RPA (sampling at Γ point and using a PTI unit cell size and norm-conserving pseudopotentials).



In absolute terms, the discrepancies with RPA appear to be larger for the two non-local functionals (rVV10 and vdW-DF2). This might suggest that the models for the effective polarizability of the two functionals, sharing an isotropic dependence on the charge density gradient, might give unprecise results in highly anisotropic configurations like gas-membrane interacting systems. However, further study is needed in order to elucidate this point. A minor role could also be played by the many-body interactions, fully included in RPA, partially included in Grimme-D3 through a three-body term and absent in the other approximations.⁶⁷

The transport of gases through two-dimensional nanopores can take place *via* direct gas phase transport or adsorbed-phase transport.^{13,68} Attributing to the small nanopores of PTI, the gas transport is expected to proceed predominantly *via* the adsorbed-phase transport mechanism, where gas first adsorbs on the

lattice, transport to the pore mouth in the adsorbed state, and finally arrive at the rate-limiting transition state at the center of the pore. Overall the gas pair selectivity is determined by the adsorption energy and relative size of the atoms/molecules with respect to the size of the nanopore. In the present case, based on the relative size of Kr (kinetic diameter of 3.6 Å) and Xe (kinetic diameter of 3.96 Å) with respect to PTI nanopore (3.4 nm), it is expected that size-sieving is the predominant separation mechanism.⁶⁹ However, while the vdW-DFT calculations reveal that PTI nanopores are promising for sieving Kr from Xe by the size-sieving mechanism, the selectivity calculations do not take entropic effects into account (loss of entropy of the gas in the adsorbed phase and in the transition state with respect to that in the gas phase). The calculations of entropic loss, for example using an enhanced method like umbrella sampling in the framework of classical MD simulations, allow one to accurately

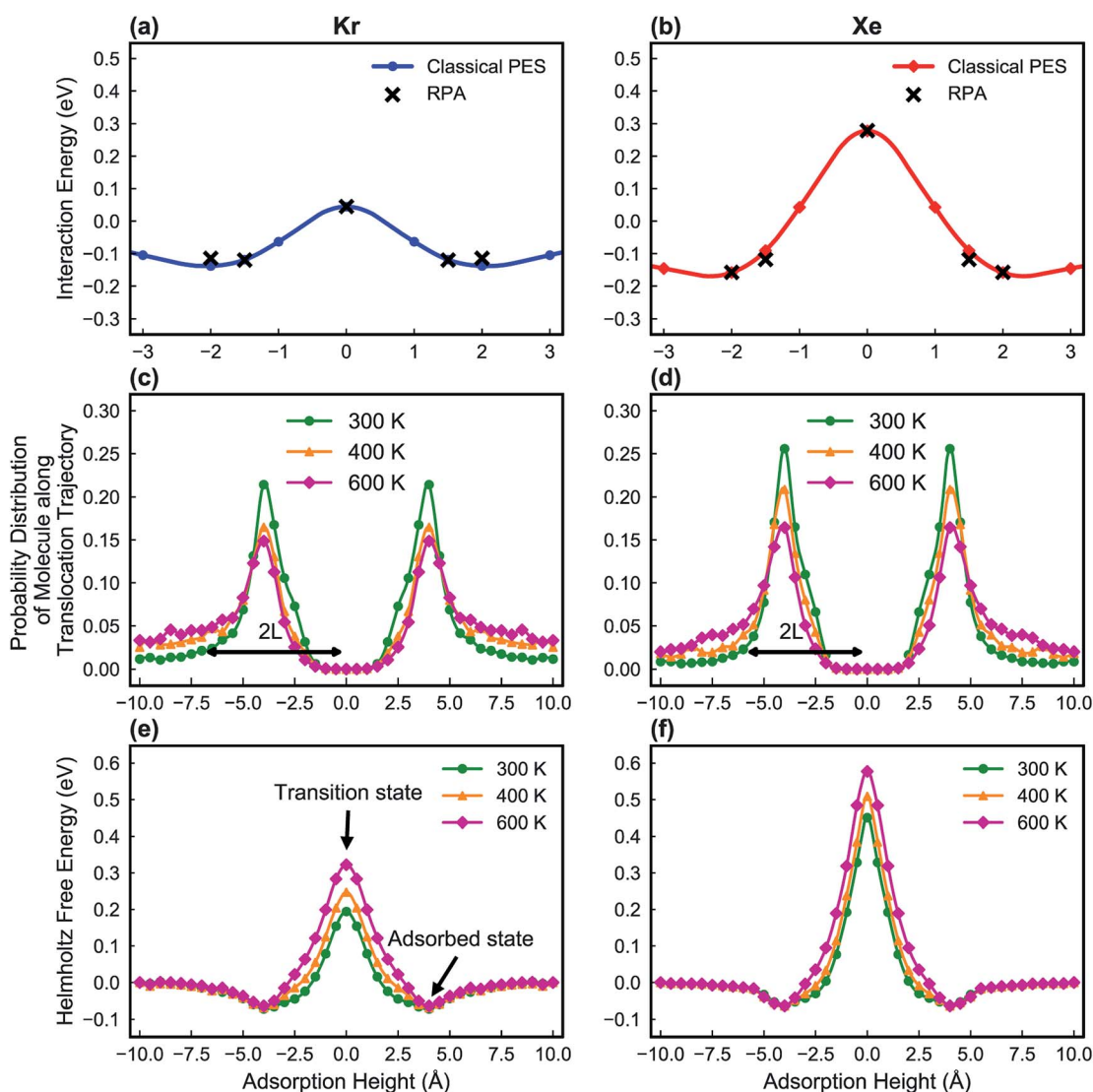


Fig. 5 The potential energy surfaces of Kr (a) and Xe (b) on a PTI monolayer, obtained with classical MD simulations to find out the best LJ parameters for Kr and Xe. The LJ parameters are trained in such a way that the potential energy surface shows a good agreement with the RPA results. The probability distribution of Kr (c) and Xe (d) along the translocation trajectories. Corresponding L (half-peak width of the probability distribution function) is shown. The Helmholtz free energy profile of Kr (e) and Xe (f) on a PTI lattice calculated with the umbrella sampling method at three different temperatures (300, 400 and 600 K) in the classical MD framework.



predict the gas permeance using a gas transport model across 2d nanopores, as *e.g.*, recently proposed by Strano and co-workers,^{53,68} and more recently by Blankschtein and co-workers.⁶⁹ A unique advantage of this route is that one can avoid unaffordable long MD simulations to track the passage of gases. This is especially relevant when the activation barriers are high, making translocation of gases a rare event.

In this study, the LJ parameters are trained such that the resulting PES in the classical MD framework agrees with those from the RPA calculations (Fig. 5a and b).

Fig. 5e and f show the calculated Helmholtz free energy profiles for Kr and Xe interacting with the PTI lattice at three different temperatures. The free energy barrier for both gases to translocate PTI nanopores increases with temperature. The barrier for Xe is larger than that for Kr, consistent with earlier observation from the vdW-DFT calculations.

In molecular transport across a nanoporous layer, it is well-established that when the interaction of a molecule at the center of the pore is repulsive, the corresponding transition state is the rate limiting step. Therefore, the translocation rate can be determined by measuring the rate of molecules crossing the transition state ($k_{\text{trans}}^{\text{TST}}$). To calculate the rate of gases crossing the transition state, an Arrhenius-type equation can be used, and parameters of such a model can be obtained using the transition state theory:^{70,71}

$$k_{\text{trans}}^{\text{TST}} = A_{\text{trans}}^{\text{TST}} \exp\left(-\frac{\Delta E}{RT}\right) \quad (3)$$

where ΔE is the energy barrier which corresponds to the activation energy needed for the translocation of gases from the adsorbed state above the pore to the center of the pore or to the transition state (as illustrated in Fig. 5e). Based on Fig. 5a and b, ΔE values for Kr and Xe are calculated to be 17.6 and 42.1 kJ mol⁻¹, respectively. The translocation prefactor, A_{trans} , can be expressed as follows:⁶⁹

$$A_{\text{trans}}^{\text{TST}} = \frac{1}{L} \sqrt{\frac{RT}{2\pi M}} \exp\left(\frac{\Delta S}{R}\right) \quad (4)$$

where L is the half-peak width of the probability distribution of gases along the translocation trajectory function (Fig. 5c and d),

M is the molecular weight of gases, and ΔS is the corresponding entropy changes for the pore translocation. ΔS can be calculated by taking the first derivative of the Helmholtz free energy changes with respect to the temperature:

$$\Delta F = \Delta E - T\Delta S \quad (5)$$

where ΔF is the Helmholtz free energy changes when the gas moves from the adsorbed state to the transition state (Fig. 5c and d). Plotting ΔF as a function of temperature and by measuring the slope of the curve, the entropy loss is calculated (Fig. 6). The entropy losses are 38.0 and 39.7 J mol⁻¹ K⁻¹ for Kr and Xe, respectively. The entropy changes are referred to as loss because the degree of freedom of the activated complex at the transition state is lower. Interestingly, in the context of separation, these calculations point that the entropic losses are similar for Kr and Xe, perhaps attributing to the fact that both of them have a similar shape (spherical atoms). Since the entropic losses for Kr and Xe are comparable, their effect on the Kr/Xe selectivity is not dramatic (~15% at room temperature). Based on the entropic loss, we calculated the translocation prefactor, A_{trans} , and the rate of gas translocation through the center of the pore, $k_{\text{trans}}^{\text{TST}}$, which are reported in Table 2 and discussed later.

Once $k_{\text{trans}}^{\text{TST}}$ is known, the gas flux can be obtained as:

$$J = k_{\text{trans}}^{\text{TST}} [C_{\text{OX}}] \quad (6)$$

Table 2 Calculated parameters for adsorption and translocation of Kr and Xe at 300 K and a feed pressure of 50 bar

	Kr	Xe
Adsorption		
$[C_{\text{OX}}]$ (mol m ⁻²)	9.6×10^{-7}	1.1×10^{-6}
Translocation through nanopore		
ΔE (kJ mol ⁻¹)	17.6	42.1
ΔS (J mol ⁻¹ K ⁻¹)	38.0	39.7
$A_{\text{trans}}^{\text{TST}}$ (s ⁻¹)	2.0×10^9	1.5×10^9
$k_{\text{trans}}^{\text{TST}}$ (s ⁻¹)	1.7×10^6	72.0
Permeance (GPU)	1000	0.05

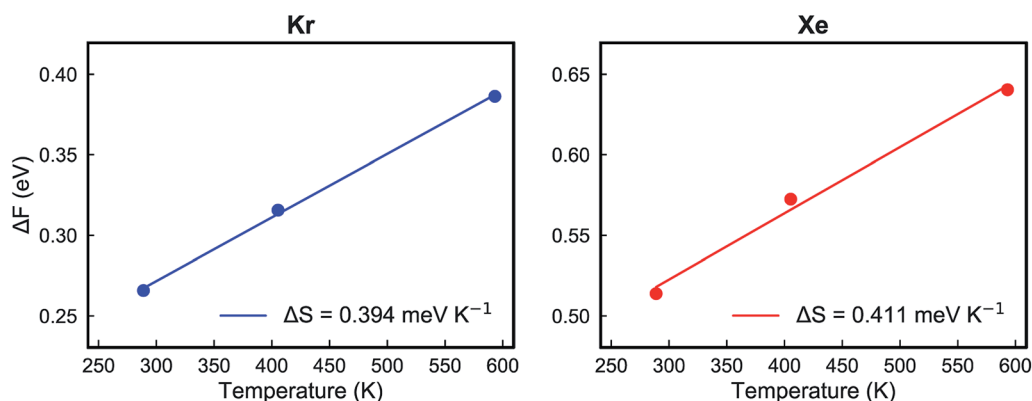


Fig. 6 The Helmholtz free energy changes of Kr (left) and Xe (right) at different temperatures when the gas arrives at the transition state from the adsorbed state.



where $[C_{\text{ox}}]$ is the concentration of occupied nanopores and can be obtained from the equilibrium relationship with respect to the concentration of unoccupied nanopores, $[C_o]$, and the gas pressure. This equilibrium can be expressed as follows:



where X, O, and O_X represent the gas phase, unoccupied nanopore and occupied nanopores, respectively. The results show that on average, 4.5 Kr and 5.0 Xe atoms adsorb on top of the PTI lattice in the simulation box. Considering the pore density in PTI, the corresponding $[C_{\text{ox}}]$ are equal to 9.6×10^{-7} and $1.1 \times 10^{-6} \text{ mol m}^{-2}$ for Kr and Xe, respectively (Table 2), based on which the flux and permeance are calculated. Finally, the Kr/Xe selectivity can be extracted by taking the ratio of the respective fluxes.

$$S_{\text{Kr/Xe}} = \frac{J_{\text{Kr}}}{J_{\text{Xe}}} \quad (8)$$

The key parameters related to adsorption and translocation of Kr and Xe are summarized in Table 2. Briefly, the concentration of adsorbed atoms corresponds to site occupancy of 37 and 41% for Kr and Xe, respectively. This sub-monolayer adsorption at a high pressure of 50 bar is expected for the noble gases. The adsorption of gas atoms sets the stage for the translocation event during which the adsorbed gas loses entropy. Since the pre-exponential factor, $A_{\text{trans}}^{\text{TST}}$, is an exponential function of entropy loss (eqn (3)), it decreases from the typical 10^{13} s^{-1} to 2.0×10^9 and $1.5 \times 10^9 \text{ s}^{-1}$ for Kr and Xe, respectively. Despite this, thanks to the presence of a single transition state in the translocation of the gases across the atom-thick PTI nanosheets, a high Kr permeance of 1000 GPU at 300 K is obtained. Owing to the relatively high activation energy for Xe to translocate the PTI nanopore, Xe permeance is low (0.05 GPU), leading to a selectivity of 20 000.

Overall, the calculated Kr permeance (1000 GPU) and Kr/Xe selectivity (20 000) from PTI at room temperature is significantly better than those from the state-of-the-art membranes demonstrated for Kr/Xe separation. Typically, selectivities in the range of 6–45 and Kr permeance in the range of 20–450 GPU have been reported. For example, 2–5 μm thick silicoaluminophosphate-34 (SAPO-34, chabazite zeolite) films hosting a nominal pore size of 0.38 nm could separate Kr/Xe with a selectivity of 23 and Kr permeance up to 26 GPU.¹ Another study using 3- μm -thick SAPO-34 membranes yielded Kr permeance of 360 GPU and selectivity of 45 which is the best reported membrane performance to date.⁹ The lower selectivity in the case of SAPO-34, with respect to that predicted from PTI membrane, can be attributed to relatively lower energy barrier of Xe to transport across 0.38 nm sized SAPO-34 pores compared to 0.34 nm sized PTI pores. Similarly, using another nanoporous material with a pore size of 0.38 nm (aluminophosphate-18 or AlPO-18), only a small Kr/Xe selectivity of 6.4 could be obtained.¹⁰ The separation of Kr/Xe with nanoporous materials hosting smaller pores (0.34 nm) has also been attempted, involving zeolitic imidazolate framework-8 or ZIF-8 membranes. However, owing to the lattice flexibility of

ZIF-8, which tends to accommodate larger molecules in ZIF-8, a small selectivity of 14.2 was realized.¹¹

The extremely high Kr/Xe selectivity from PTI nanopore makes it especially attractive for the separation of ^{85}Kr from a 10/90 mixture of $^{85}\text{Kr/Xe}$, targeting near pure ^{85}Kr and Xe streams using a PTI membrane. This will allow one to safely use the purified Xe for commercial applications. The recent development in obtaining single-layer PTI nanosheets⁷² will allow one to fabricate a few nanometer thick films, similar to what has been achieved in the case of graphene and metal-organic framework (MOF) nanosheet-based membranes.^{73,74}

4 Conclusions

This study reveals the high potential of PTI-based membrane, thanks to $\sim 0.34 \text{ nm}$ -sized nanopores, for separation of Kr/Xe mixtures. In order to ensure accurate predictions on gas selectivity, which are critically influenced by small changes in the adsorption and the activation energies, we performed an extensive comparison of several vdW-corrected functionals within DFT and benchmarked them against accurate, reference EXX/RPA calculations. We highlight that differences in commonly used vdW functionals can lead to dramatic changes in selectivity measurements, raising accuracy concerns for some of these that require further care of practitioners in the field. In order to consider finite-temperature effects, the entropic contribution in the transition state has been determined from Helmholtz free energy calculations at different temperatures, obtained with restrained MD simulations where the force-field parameters have been trained on the EXX/RPA potential-energy profile. Finally, gas permeances are predicted by calculating the translocation coefficient from single-layer PTI using transition-state theory. We obtain, from our simulations, a Kr/Xe separation factor exceeding 10 000 at room temperature, which would allow one to obtain nearly pure ^{85}Kr and Xe streams.

Overall, the method developed here, vdW-DFT calculations benchmarked against EXX/RPA calculations to predict the energy barrier and entropic penalties for gas translocation across a nanopore, can be applied to a diverse set of two-dimensional nanoporous films, *e.g.*, nanoporous graphene, C_2N , polyheptazine imide, *etc.*, for calculating the molecular separation performance. Apart from the high accuracy, the method used here to predict permeance is also attractive in comparison to the classical MD simulations when the energy barrier for the translocation event of at least one component is high enough to make the permeation a rare event, in which case, one has to conduct time-consuming and expensive long MD simulation trials.

Conflicts of interest

There are no conflicts to declare.

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