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Molecular hydrogen isotope separation by a graphdiyne membrane: a quantum-mechanical study

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Graphdiyne (GDY) has emerged as a very promising two-dimensional (2D) membrane for gas separation technologies. One of the most challenging goals is the separation of deuterium (D_2) and tritium (T_2) from a mixture with the most abundant hydrogen isotope, H_2 , an achievement that would be of great value for a number of industrial and scientific applications. In this work we study the separation of hydrogen isotopes in their transport through a GDY membrane due to mass-dependent quantum effects that are enhanced by the confinement provided by its intrinsic sub-nanometric pores. A reliable improved Lennard-Jones force field, optimized on accurate *ab initio* calculations, has been built to describe the molecule–membrane interaction, where the molecule is treated as a pseudoatom. The quantum dynamics of the molecules impacting on the membrane along a complete set of incidence directions have been rigorously addressed by means of wave packet calculations in the 3D space, which have allowed us to obtain transmission probabilities and, in turn, permeances, as the thermal average of the molecular flux per unit pressure. The effect of the different incidence directions on the probabilities is analyzed in detail and it is concluded that restricting the simulations to a perpendicular incidence leads to reasonable results. Moreover, it is found that a simple 1D model—using a zero-point energy-corrected interaction potential—provides an excellent agreement with the 3D probabilities for perpendicular incidence conditions. Finally, D_2/H_2 and T_2/H_2 selectivities are found to reach maximum values of about 6 and 21 at ≈ 50 and 45 K, respectively, a feature due to a balance between zero-point energy and tunneling effects in the transport dynamics. Permeances at these temperatures are below recommended values for practical applications, however, at slightly higher temperatures (77 K) they become acceptable while the selectivities preserve promising values, particularly for the separation of tritium.

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1 Introduction

Deuterium (D_2) is an essential substance for several applications in industry, scientific research and medicine. In addition, there is a great interest in the efficient production of D_2 and tritium (T_2) for their use in the emerging nuclear fusion industry. Separation of these scarce isotopes from a mixture with H_2 is a tremendous challenge because standard technologies for gas separation are not efficient for isotope separation, as they rely on physicochemical properties that are nearly identical for these species. A very interesting alternative is quantum sieving, first proposed by Beenakker *et al.*,¹ where the mass-dependent quantum-mechanical behavior of the isotopes is enhanced by

confinement in sub-nanometric spaces. Based on these ideas, there has been an impressive progress in the use of nanoporous materials for isotope separation.^{2–4} An issue—also occurring in the separation of gases in general—is that large selectivities for separation are often accompanied by low permeances (thermal molecular flux per unit pressure) and *vice versa*.⁵ As permeance is usually inversely proportional to the membrane thickness, new one-atomic-thick, two-dimensional (2D) nanoporous membranes offer great prospect for large selectivities without compromising the permeability.^{6–10}

One promising 2D membrane for the separation of hydrogen isotopes is graphdiyne (GDY). Since the first report on its synthesis in 2010,¹¹ there has been a tremendous increase of experimental and theoretical studies on GDY, where several properties have been studied for applications in catalysis, electronics, batteries, solar cells, *etc.*^{12–14} In particular, GDY appears to be an ideal membrane for gas separation applications^{15,16} due to its mechanical stability, chemical inertness and, specially, its porous structure, which is characterized by benzene-like rings linked by diacetylene chains giving rise to a uniform mesh of

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subnanometric triangular pores.¹⁷ Interestingly enough, hydrogen purification has been one of the first applications proposed for this material. Based on Density Functional Theory calculations, Jiao *et al.*¹⁸ predicted that GDY could be very efficient for the separation of H₂ from a mixture with CH₄ and CO. Several other computational studies have appeared^{19–26} proposing, for instance, an improvement of the H₂ permeability/selectivity by means of nitrogen²³ or positive charge^{24,26} doping of the GDY pores. As far as we are aware, hydrogen isotope separation by GDY has not been addressed yet.

We believe that it is important to study the isotope separation by 2D membranes as accurately as possible, with the aim to provide reliable references for simpler models which are often employed in the literature. These processes commonly involve an energy barrier, at the center of the pore, in the molecule-membrane interaction. For light species (within the quantum-mechanical description of the motion of the nuclei), there are two features that are quite important: the tunneling through the barrier and the quantization of the energy levels of the transition state (TS) at the pore center, generally called zero-point energy (ZPE) effects. While tunneling favors the transmission of the lighter isotope, ZPE effects enhance the transport of the heavier species.^{27,28} The selectivity for isotope separation is the result of a delicate balance between these effects, a balance that sometimes simple methods fail to describe correctly. As an example, some of us applied the three-dimensional time-dependent wavepacket (3D-TDWP) propagation method (first designed for atom-surface scattering²⁹) to study the transport and separation of He isotopes through 2D membranes.³⁰ The results were compared with predictions based on tunneling-corrected TS theory (tunn-TST).²⁸ A good agreement was found for the dependence of the ⁴He/³He selectivity with temperature when the separation membrane is GDY. However, for a holey graphene model, tunn-TST overestimates considerably the ⁴He/³He selectivity because it is unable to account for the extent of tunneling as well as other quantum features (resonances) found in the accurate treatment.³⁰ Helium isotopes quantum sieving through a graphtriyne membrane was also studied by means of the 3D-TDWP calculations,³¹ which have served as benchmarks for ring polymer molecular dynamics calculations.³²

In this work, the transport and separation of hydrogen isotopes by GDY are studied by means of 3D-TDWP calculations. A reliable force field describing the interaction potential energy between H₂, treated as a pseudoatom, and GDY is built and optimized on previously reported accurate first principles calculations.³³ For the first time, all relevant directions of incidence of the particles are taken into account in the computations and the results are compared with more commonly employed treatments that only consider a perpendicular approach to the membrane. In addition, we test a simpler model-recently employed in related studies^{34,35} – which involves one-dimensional wave packet (1D-TDWP) calculations of transmission probabilities along a ZPE-corrected potential energy curve^{36,37} in the perpendicular approach. Permeances of H₂ and D₂ are reported as functions of temperature as well as their D₂/H₂ selectivity ratio. Separation of tritium is also addressed within the more approximated treatments.

The paper is arranged as follows. In Section 2, the interaction potential is described and the methods for the calculation of quantum-mechanical probabilities and permeances are summarized. Results are reported and discussed in Section 3, concerning transmission probabilities, permeances, selectivities as well as a comparison with more approximate models. Finally, conclusions are outlined in Section 4. An Appendix is dedicated to provide some computational details.

2 Theoretical approach

We consider the transport of an H₂ molecule or its isotopes through a periodic and rigid GDY membrane. In the present approach, the molecules are treated as point-like particles (pseudoatoms), an approximation usually adequate to describe the molecules in their ground rotational state (*para*-H₂, *ortho*-D₂, *para*-T₂). The *ab initio* study of ref. 33 indicates that this is a reasonable approach, as the computed interaction energies do not differ too much for different orientations of the molecule around the pore region, where the corresponding anisotropy is expected to be the largest. Indeed, it was found that parallel and perpendicular orientations of H₂ with respect to the GDY plane involve similar permeation barriers (a difference of 2 meV for barrier heights of about 50 meV).³³ On the other hand, in the present model there is no need to invoke other processes such as the H₂ dissociation and subsequent formation of C–H bonds in the membrane, since they involve energy barriers much larger than the H₂ translational energies considered here (about 2.1 eV for the hydrogenation of acetylenic carbon atoms³⁸ and larger values for benzenic ones^{39,40}).

In what follows and within the Born-Oppenheimer scheme of separation of the electronic and nuclei motion, we will firstly present the force field employed (representing the solution of the electronic Schrödinger equation), then, the 3D-TDWP method (within the Schrödinger equation for the motion of the nuclei) to compute the transmission probabilities and finally, the calculation of permeances and selectivities and some approximate models for the perpendicular incidence.

2.1 Interaction potential

The total H₂–GDY interaction potential is described as a sum of H₂–C pairwise contributions (H₂ as a point-like particle), each one represented by a potential term V^i :

$$V = \sum_i V^i, \quad (1)$$

and the above sum runs over all carbon atoms composing the GDY plane. For the representation of each V^i term, the Improved Lennard-Jones (ILJ) formulation⁴¹ is used:

$$V^i(R) = \varepsilon \left[\frac{6}{n(x) - 6} \left(\frac{1}{x} \right)^{n(x)} - \frac{n(x)}{n(x) - 6} \left(\frac{1}{x} \right)^6 \right], \quad (2)$$

where x is a reduced pair distance $x = R/R_m$ (between C and the center of mass of H₂), and ε and R_m represent the well depth and equilibrium distance of the pair interaction, respectively.



Table 1 ILJ optimized parameters for the H₂-C and H₂-H pairs involved in the interaction between H₂ and the GDY molecular precursor. The parameters for the H₂-C pairs were the ones employed to describe the H₂-GDY interaction. R_m and ε are in Å and in meV, respectively, while β is dimensionless

Pair	R_m	ε	β
H ₂ -C	3.503	4.569	6.5
H ₂ -H	3.279	2.301	9.0

Moreover, $n(x)$ is expressed by $n(x) = \beta + 4.0x^2$ where β is a parameter defining the shape and stiffness of the potential.

The ILJ parameters that have been used are reported in Table 1. Zeroth-order values of ε and R_m were obtained from data of the polarizabilities of the interacting partners,⁴² following expressions reported in ref. 43. In this way, these parameters maintain a physical meaning and, in turn, are transferable to other systems involving H₂ interacting with graphene derivatives. These values, together with the remaining β parameter, were subsequently optimized on the basis of comparisons of the ILJ energies with reference *ab initio* energies for a set of selected positions of the center of mass of H₂ with respect to the GDY pore (shown in Fig. 1). Within the adopted pseudoatom approximation, the ILJ force field represents the interaction for an average over all possible orientations of the molecule. The reference *ab initio* energies adequate to this purpose were obtained by averaging the *ab initio* interaction potential profiles of ref. 33 corresponding to two parallel and one perpendicular orientations of the molecule with respect to the GDY plane. In addition, it is noted that those *ab initio* estimations were obtained for the interaction between H₂ and a specific (rigid) GDY molecular precursor, shown in the upper panel of Fig. 1. This molecular prototype represents the smallest approximation of a GDY pore and it has been previously used to assess the features of the interaction between GDY and different atomic and molecular species.^{33,44,45} It is worth pointing out that this precursor includes H atoms which saturate some C atoms of the benzenic rings, so it was necessary to determine ILJ parameters also for the H₂-H interaction (included in Table 1). However, these H₂-H parameters are not needed to compute the interaction with the extended GDY membrane, which is only formed by carbon atoms.

In Fig. 1 a comparison between the ILJ force field and the reference *ab initio* calculations is presented for different interaction profiles, corresponding to out-of-plane (z axis) and in-plane (x and y axes) displacements of the center of mass of the H₂ molecule with respect to the center of the pore. A quite good agreement can be observed for both out-of-plane and in-plane interaction potential features, which validate the reliability of the current force field. Notice that the final optimized ε and R_m parameters reported in Table 1 differ from their zeroth-order estimates by just about 1%. In the upper panel, a low barrier (of about 50 meV) for H₂ permeation through the pore can be noticed, which is in agreement with the previous theoretical prediction reported in ref. 33 (50 meV). From the intermediate and lower panels a quite large interaction anisotropy is also

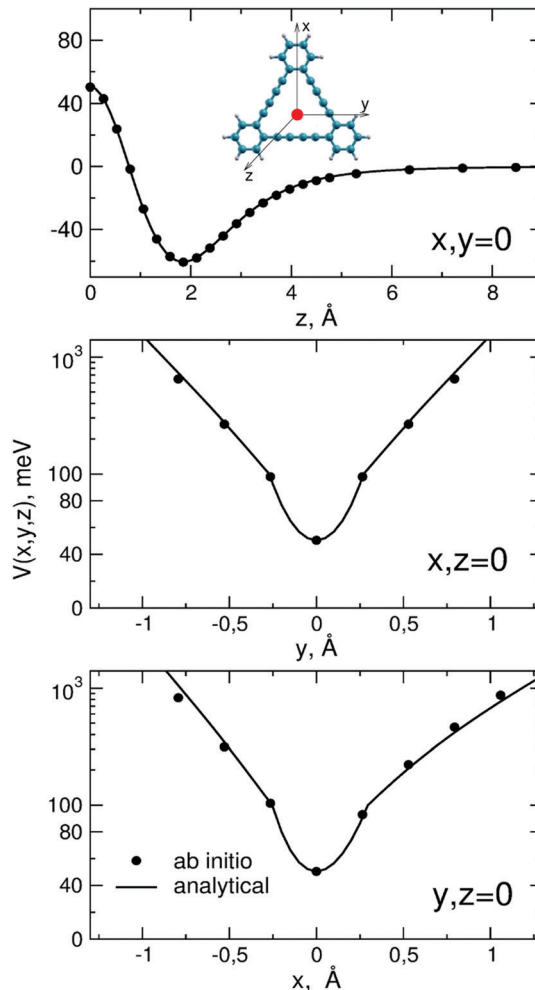


Fig. 1 Potential energy for the interaction between H₂ (described as a pseudoatom) and the smallest precursor of the GDY pore. Upper panel: Planar molecular prototype used to represent the GDY pore; interaction profile for the H₂ out-of-plane perpendicular approach towards the pore center (z axis). Intermediate and lower panels: interaction profile for the H₂ in-plane displacements with respect to the pore center along x and y , respectively. Full circles have been obtained by averaging the *ab initio* energies for two parallel and one perpendicular orientations of H₂ onto the GDY plane.³³ The analytic ILJ force field is shown with solid lines.

evidenced when the H₂ molecule is out of the pore geometric center: even for displacements as short as 0.25–0.30 Å in both directions the potential is more than twice and then it increases rapidly reaching very repulsive values of about 1 eV at distances of around 1 Å from the pore center.

The GDY unit cell used in this work, set on the $z = 0$ plane, is illustrated in the upper panel of Fig. 2. The interaction potential between H₂ and GDY has been obtained by applying eqn (1) to as many carbon atoms in the membrane until the desired convergence is reached (including up to ten “shells” of unit cells around the central one). Dependence of V with x and z , for $y = 0$, is depicted in the lower panel of Fig. 2. The center of the pore coincides with the origin of the coordinate system and at this point, $V = 47.75$ meV; this value equals the barrier height since the zero of energy is set for an infinite separation of the



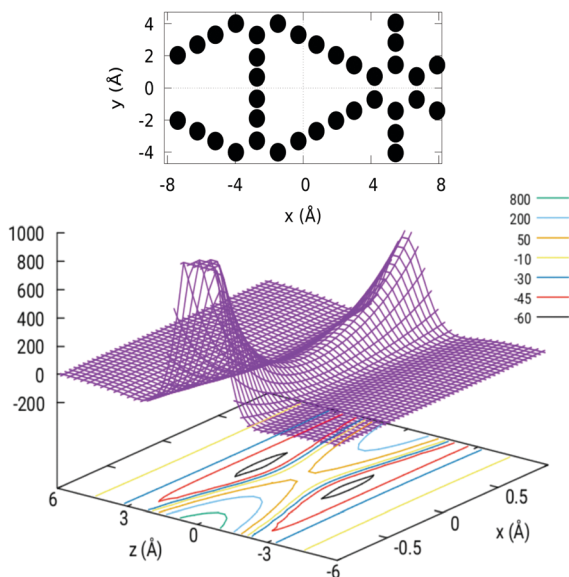


Fig. 2 Upper panel: GDY unit cell employed in the 3D-TDWP calculations, with indication of the coordinate system used. Lower panel: H_2 -GDY interaction potential (vertical axis, in meV) as a function of x and z (in Å), around the center of the pore, with $y = 0$.

molecule from the membrane. At this point, the potential exhibits a saddle-point topography while for larger distances from the membrane (about 2 Å) there is a well (63 meV) formed by van der Waals attractive forces (physical adsorption well).

2.2 Three-dimensional time-dependent wave packet (3D-TDWP) propagation

The 3D-TDWP method applied to the transmission of atoms through a 2D membrane has been described in ref. 30, therefore here we will just present a brief summary and discuss some aspects that are specific to the present study. A comprehensive exposition of the time-dependent approach of quantum mechanics can be found elsewhere.^{46,47}

The Hamiltonian for the interaction of a point-like H_2 isotope of mass m with a GDY membrane can be written as

$$H = \frac{-\hbar^2}{2m} \left[\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2} \right] + V(x, y, z), \quad (3)$$

where V is the interaction potential presented above.

A wave packet representing the molecule is discretized in a grid of evenly spaced (x, y, z) points. Following the work of Yinon and Kosloff,²⁹ the periodicity of the system is exploited by limiting the ranges of coordinates x and y to those of a unit cell (Fig. 2). The range in the perpendicular coordinate, z , should be large enough to describe the evolution of the wave packet from the initial approach to the membrane up to the final distancing from it after the interaction. The initial wave packet is given by

$$\psi(x, y, z, t = 0) = G(z; z_0, k_{z_0}, \alpha) \frac{\exp[i\mathbf{K} \cdot \mathbf{R}]}{(A_x A_y)^{1/2}}, \quad (4)$$

where G is a normalized Gaussian wavepacket⁴⁸ (see Appendix) and the other function is a plane wave with a wave vector \mathbf{K} ,

A_x and A_y being the lengths of the unit cell ($A_x = \sqrt{3}A_y$, $A_y = 9.45 \text{ \AA}$). The values of \mathbf{K} must be chosen such that the plane wave is periodic with the lattice,²⁹ a condition achieved when

$$\mathbf{K} = 2\pi \left(\frac{l_x}{A_x}, \frac{l_y}{A_y} \right), \quad (5)$$

l_x, l_y being integer numbers. In this way, the initial wave packet has well-defined parallel velocities

$$\mathbf{v}_{\parallel} = (v_{l_x}, v_{l_y}) = \frac{2\pi\hbar}{m} \left(\frac{l_x}{A_x}, \frac{l_y}{A_y} \right), \quad (6)$$

and energy,

$$E_{\parallel}(l_x, l_y) = \frac{(2\pi\hbar)^2}{2m} \left(\left[\frac{l_x}{A_x} \right]^2 + \left[\frac{l_y}{A_y} \right]^2 \right), \quad (7)$$

while in the perpendicular direction it has a distribution of velocities v_z (and perpendicular energies $E_{\perp} = mv_z^2/2$) centered around $v_{z0} = \hbar k_{z0}/m$. Likewise, the incidence polar angle, $\theta = \arctan(v_{\parallel}/v_z)$, is not well defined in the wave packet of eqn (4) (except for the perpendicular incidence, $v_{\parallel} = 0$); instead it is distributed around $\arctan(v_{\parallel}/v_{z0})$. The azimuthal angle, $\phi = \arctan(v_y/v_x)$, does have a definite value. The total translational energy, $E = E_{\parallel} + E_{\perp}$, coincides with the total energy since at the start of the propagation the potential energy $V \approx 0$.

The solution of the time-dependent Schrödinger equation, $\psi(t) = \exp(-iHt/\hbar)\psi(0)$, will allow us to extract the probability of transmission through the membrane, $\mathcal{P}_{l_x, l_y}(v_z)$. The propagation of the wavepacket is carried out employing the Split-Operator method⁴⁹ where the evolution operator, $\exp(-iHt/\hbar)$, is approximately splitted as a product of potential and kinetic evolution operators, which are successively applied to the wavepacket. Propagation with the potential and kinetic terms is performed in the space and momentum representations, respectively. The fast Fourier transform is used to transform the wavepacket between both representations. The error associated to the Split-Operator method is $\mathcal{O}(\Delta t^3)$.⁴⁷

The stationary wave function, $\Psi_E^+(x, y, z)$, can be obtained from $\psi(x, y, z, t)$ using the Fourier transform from the time to the energy space.^{46,50} Finally, the transmission probability is obtained from the flux of $\Psi_E^+(x, y, z)$ through a surface $z = z_f$ separating transmitted from incident and reflected waves,^{30,51}

$$\mathcal{P}_{l_x, l_y}(v_z) = \frac{2\pi\hbar^2}{m} \text{Im} \left(\int dx dy \Psi_E^{+*}(x, y, z_f) \left[\frac{\partial \Psi_E^+}{\partial z} \right]_{z=z_f} \right), \quad (8)$$

where $\left[\frac{\partial \Psi_E^+}{\partial z} \right]_{z=z_f}$ indicates the derivative of $\Psi_E^+(x, y, z)$ with respect to z , computed at $z = z_f$.

Details of these calculations are given in the Appendix.

2.3 Calculation of permenances and selectivities

The flux of molecules with a certain velocity $\mathbf{v} = (v_x, v_y, v_z)$ crossing a membrane placed in the (x, y) plane can be defined as the product of the flux of molecules hitting the membrane, Z , and the probability \mathcal{P} that these molecules actually cross the



membrane

$$J(v_x, v_y, v_z) = Z \mathcal{P}(v_x, v_y, v_z). \quad (9)$$

As it is known,⁵² $Z = \rho v_z$, where ρ is the number density of the molecular gas. The permeating flux at a temperature T is obtained by the thermal averaging:

$$J(T) = \langle Z \mathcal{P} \rangle_T = \rho \int_{-\infty}^{\infty} dv_x f(v_x, T) \int_{-\infty}^{\infty} dv_y f(v_y, T) \int_0^{\infty} dv_z f(v_z, T) v_z \mathcal{P}(v_x, v_y, v_z), \quad (10)$$

where $\langle \dots \rangle_T$ indicates thermal averaging and $f(v_x, T)$, $f(v_y, T)$ and $f(v_z, T)$ are the Maxwell-Boltzmann velocities distributions in cartesian coordinates. In eqn (10), the integral in v_z only involves positive v_z values as only forward directions are effective for the transmission.

The permeance S of the membrane for the molecular transmission is defined as the thermal flux $J(T)$ divided by ΔP , the pressure difference across the membrane.⁹ The gas is assumed to be sufficiently diluted in the feed side of the membrane at a pressure P and temperature T . Despite existing a molecular flow through the membrane pores, the gas-membrane system can be considered in equilibrium if the pores are sufficiently small compared to the mean free path of the gas,⁵² $\lambda = K_B T / (\sqrt{2} \sigma P)$, where σ is the molecular cross sectional area. Taking $\sigma_{H_2} = 27 \text{ \AA}^2$ and for $T = 40 \text{ K}$ and $P = 10 \text{ bar}$, $\lambda = 15 \text{ \AA}$, much larger than the pore size, of less than 1 \AA . Therefore the system can be safely regarded as in equilibrium for pressures below 10 bar. In these conditions, the ideal gas law is valid and it is used to substitute ρ in eqn (10) by $P/K_B T$. In this way, the permeance writes

$$S(T) = \frac{\langle v_z \mathcal{P} \rangle_T}{K_B T} \cdot \frac{P}{\Delta P}. \quad (11)$$

We assume that the pressure in the permeate side is negligible ($\Delta P \approx P$), so $\frac{P}{\Delta P} = 1$ and the permeance becomes independent of pressure (a very weak dependence of the permeance on pressure has been confirmed by different molecular dynamics calculations⁵³⁻⁵⁵). Some other specific values of $P/\Delta P$ could be set, as done in ref. 20 and 56.

Finally, when two different molecular species A and B are hitting the membrane, the A/B selectivity or separation factor is given by the ratio of the permeances of A and B,

$$R_{A/B}(T) = \frac{S_A(T)}{S_B(T)}. \quad (12)$$

In this work we have run 3D-TDWP calculations for a set of incident conditions (l_x, l_y) , where $l_{x,y}$ varies from $-l_{x,y}^{\max}$ to $l_{x,y}^{\max}$ with $\Delta l_{x,y}$ increments. In this way, the permeance is actually computed using a trapezoidal rule for the integration

in v_x and v_y ,

$$S(T) = \frac{\Delta v_{x,y} \Delta v_y}{K_B T} \sum_{l_x=-l_x^{\max}}^{l_x^{\max}} \sum_{l_y=-l_y^{\max}}^{l_y^{\max}} C_{l_x} C_{l_y} f(v_{l_x}, T) f(v_{l_y}, T) \int_0^{\infty} dv_z f(v_z, T) v_z \mathcal{P}_{l_x, l_y}(v_z), \quad (13)$$

where $\Delta v_{x,y} = 2\pi\hbar\Delta l_{x,y}/(m\Delta_{x,y})$ and $C_{l_{x,y}} = 1/2$ for $|l_{x,y}| = l_{x,y}^{\max}$ or 1, otherwise. In the calculations, we have chosen $l_x^{\max} = 16$, $\Delta l_x = 4$, $l_y^{\max} = 12$, $\Delta l_y = 4$ for H_2 and $l_x^{\max} = 24$, $\Delta l_x = 6$, $l_y^{\max} = 16$, $\Delta l_y = 8$ for D_2 . After several tests where $l_{x,y}^{\max}$ and $\Delta l_{x,y}$ values were varied, the chosen values were found to be sufficient for the convergence of the permeances.

It should be noted that a different (less exact) definition of permeance has been used in the literature,^{20,31,32,35,56,57} where the molecular flux Z and the transmission probability are separately averaged over the Maxwell-Boltzmann distribution:

$$S'(T) = \frac{\langle Z \rangle_T \langle \mathcal{P} \rangle_T}{P} = \frac{\langle v \rangle_T \langle \mathcal{P} \rangle_T}{4K_B T}, \quad (14)$$

where $\langle v \rangle_T = (8K_B T/\pi m)^{1/2}$. In the present study we have found that although permeances obtained from eqn (14) are about 10 times smaller than those computed using eqn (11), their general dependence with temperature is rather similar.

2.4 The case of perpendicular incidence: 3D and 1D calculations

We aim to study the effect on the permeances of just computing the 3D transmission probability (Section 2.2) along the perpendicular incidence, an approach employed in some previous works^{30,31} and that considerably reduces the computational cost. For this approximation, the permeances are computed by setting the equation above that $\mathcal{P}_{l_x, l_y}(v_z) = \mathcal{P}_{0,0}(v_z)$, $\forall l_x, l_y$, leading to

$$S_p(T) = \frac{1}{K_B T} \int_0^{\infty} dv_z f(v_z, T) v_z \mathcal{P}_{00}(v_z). \quad (15)$$

The validity of this approximation is discussed in the following Section.

In addition, we have considered a low-cost 1D model for the transmission probability at perpendicular incidence, which has recently been employed in related works.^{34,35} This model, related to reaction path theories,^{36,37} involves an adiabatic separation between the coordinates parallel to the membrane (x, y) and the perpendicular one (z). In a first step, the Schrödinger equation is solved for the parallel degrees of freedom at various fixed distances z ,

$$\left\{ \frac{-\hbar^2}{2m} \left[\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} \right] + V(x, y; z) \right\} \phi_n(x, y; z) = W_n(z) \phi_n(x, y; z). \quad (16)$$

The eigenvalues $W_n(z)$ are the so-called adiabatic potential energies. The ground state energies, $W_0(z)$, correspond to the ZPE-corrected potential along the perpendicular direction. In other words, the ZPE along the perpendicular incidence, associated to the ground-state vibrations of the center of mass of the molecule along the x and y directions, is the difference



$W_0(z) - V(0,0,z)$. This correction will differ for the various isotopes studied due to their different mass m . Next, the transmission probability is obtained by solving a 1D Schrödinger equation along z , using $W_0(z)$ as effective potentials. This task has been carried out by adapting the 3D-TDWP approach described above to a 1D problem. Then, these probabilities are scaled using a factor γ that takes into account the fraction of the membrane that is effective for the transmission,^{30,58}

$$\gamma = n_p \frac{A_{\text{eff}}}{A_{\text{uc}}}, \quad (17)$$

where A_{eff} is the effective pore area, $A_{\text{uc}} = \Delta_x \Delta_y$ is the unit cell area and n_p is the number of pores in the unit cell (four in the present case, see Fig. 2).

Eqn (16) is also used to compute a set of energies and wave functions at $z = 0$, a task that helps to understand—within the ideas of the TS theory—various features of the transmission probabilities, as shown below.

The bound-state problem of eqn (16) has been solved following the procedures described in ref. 28.

3 Results

In Fig. 3 we present the quantum-mechanical probabilities (eqn (8)) for the transmission of H_2 and D_2 through GDY, obtained for different incidences (l_x, l_y) of the initial wave packets. First of all, we discuss the “staircase” structure of these probabilities aided by some concepts of the TS theory.⁵⁹ In this scheme, the different energy levels and functions of the TS are obtained by diagonalizing the Hamiltonian of eqn (16) for $z = 0$, where the bottleneck for the transmission is. These energy levels $W_n(0)$ are given in Table 2 for H_2 and D_2 and the corresponding wave functions for D_2 are depicted in Fig. 4. It can be seen that the thresholds of the different “steps” of the probabilities (see vertical arrows in Fig. 3) occur at energies very close to the TS energies of Table 2. In this way, these energies can be considered as the effective (and successive) energy barriers for permeation. It can be seen that the transmission probabilities for the perpendicular incidence $(l_x, l_y) = (0,0)$ (black lines in Fig. 3) lack the “second” threshold (the rise of probability at the energy of the $n = 2, 3$ states). This is due to the relationship between the symmetries of the initial wave packet and those of the TS wave functions: an initial wave packet with $(l_x, l_y) = (0,0)$ does not populate states antisymmetric with respect to the y or x axis, as the $n = 2, 3$ states (Fig. 4). Apart from that, it is worth noting the large difference between the first energy thresholds (100.63 for H_2 and 84.85 for D_2) and the potential energy barrier (47.75 meV), in other words, the large ZPEs involved (52.88 and 37.1 meV for H_2 and D_2 , respectively). This is due to the light masses of the molecules as well as the confinement imposed by the interaction potential along the x and y directions (Fig. 1). Moreover, the lower energy threshold of D_2 with respect to that of H_2 will cause a preferred permeation of the heavier isotope over a wide range of temperatures, as will be seen below. Note that this ZPE appears in an intrinsic manner when solving the 3D time-dependent Schrödinger

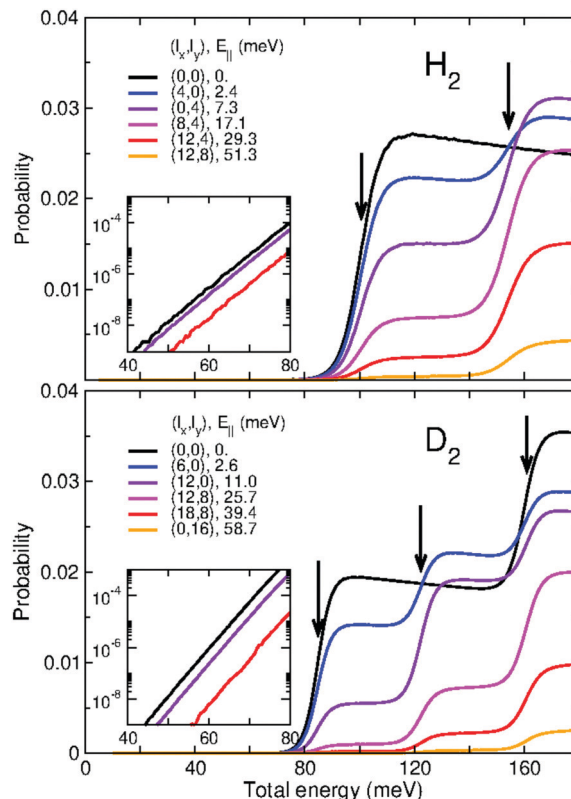


Fig. 3 3D-TDWP transmission probabilities for H_2 -GDY (upper panel) and D_2 -GDY (lower panel) as functions of the total translational energy (meV) for different incidences (l_x, l_y) . The corresponding parallel energies E_{\parallel} (l_x, l_y) (eqn (7)) are indicated in the legends. Arrows show the energy levels of the transition state (Table 2). Insets depict the behavior of some of the probabilities in the low energy range. See text for discussion.

Table 2 Energy levels of the H_2/D_2 -GDY transition state ($W_n(0)$ of eqn (16))

n	Energy (meV)	
	H_2	D_2
0	100.63	84.85
1	154.35	122.36
2	154.35	122.36
3	208.57	160.11
4	210.16	160.95
5	210.16	160.95

equation, producing an effective permeation barrier, whereas it can be clearly identified as the difference $W_0(z = 0) - V(0,0,0)$ in the solution of eqn (16) within the TS theory.

In the insets of Fig. 3 we present the low-energy behavior of some of the transmission probabilities, specifically, the probabilities for $(l_x, l_y) = (0,0)$, $(0,4)$ and $(12,4)$, for H_2 , and $(0,0)$, $(12,0)$ and $(18,8)$, for D_2 . For these energies, tunneling effects play an important role in the behavior of the probabilities. It can be clearly seen that the probabilities decrease exponentially as energy decreases. This decay occurs at a smaller rate in the case of H_2 , because tunneling is much easier for this light molecule



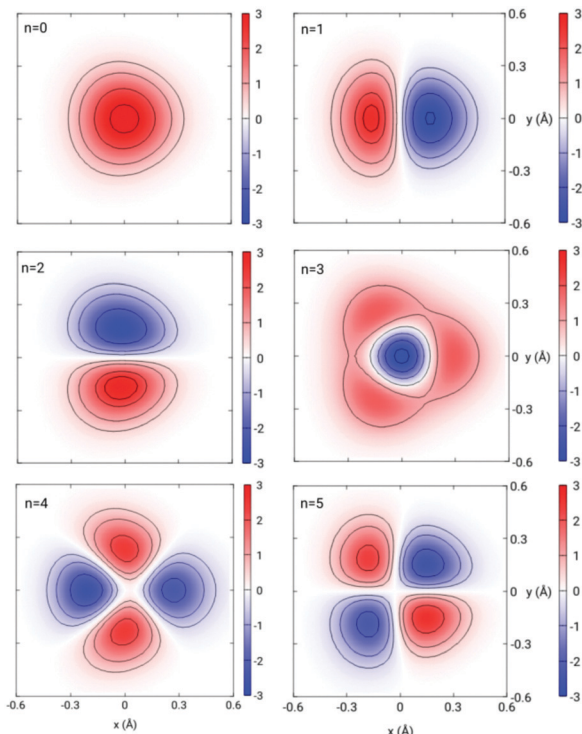


Fig. 4 Contour plots of the wave functions $\phi_n(x, y; z = 0)$ (in arbitrary units) of the D_2 -GDY transition state for levels $n = 0-5$.

than for the heavier D_2 . Indeed, at the lowest translational energies (40 meV), the transmission probabilities for H_2 become larger than those for D_2 . This behavior is at the origin of the maximum value found for the D_2/H_2 selectivity as a function of the temperature, as described below.

It is worth discussing in more detail the dependence of the transmission probabilities with the different incidence conditions (l_x, l_y) of the wave packet. In Fig. 3 it is seen that all the probabilities rise at the same energy thresholds, however the height of the “plateau” that they reach rapidly decreases with the parallel energy, $E_{\parallel}(l_x, l_y)$. Interestingly, we have found that for the various combinations of l_x, l_y that lead to the same $E_{\parallel}(l_x, l_y)$, the transmission probabilities are equal. For instance, $(l_x, l_y) = (12, 4)$ and $(0, 8)$ have identical E_{\parallel} (because $\Delta_x = \sqrt{3}\Delta_y$, see eqn (7)) and the associated probabilities are found to be identical too. This property allows us to plot the H_2 and D_2 probabilities as functions of E_{\parallel} for various fixed values of E , as shown in Fig. 5. For total energies below the second permeation threshold ($E < 154$ and < 122 meV for H_2 and D_2 , respectively), it is quite clear that the probabilities decrease exponentially with E_{\parallel} . This behavior is modified for higher energies due to the rise of the probabilities with $(l_x, l_y) \neq (0, 0)$ at the second threshold, discussed above. A decrease of the probabilities with E_{\parallel} was reasonably expected since the wave packets that approach the membrane in directions distant from the perpendicular one do not access too directly to the pore region. However, it is difficult to foresee the specific dependence (as it is determined by intricate details of the dynamics), so it is noteworthy to have found a simple exponential decay. Notice

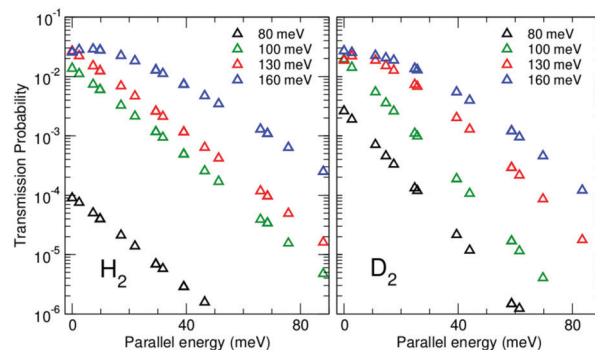


Fig. 5 3D-TDWP transmission probabilities for H_2 -GDY (left panel) and D_2 -GDY (right panel) as functions of parallel energy E_{\parallel} (meV) for total translational energies equal to 80, 100, 130 and 160 meV.

also that the decay rate is stronger for the D_2 probabilities than for the H_2 ones.

Permeances of GDY for the transport of H_2 and D_2 , in gas permeance units (GPU), are reported in Fig. 6 (upper panel) for temperatures between 40 and 150 K. D_2/H_2 selectivities are presented in the lower panel of the same Figure. We first discuss the permeances obtained from the calculations that include all the incidence conditions (l_x, l_y) considered (eqn (13), solid lines in the Figure). It can be seen that the permeances increase by several orders of magnitude as temperature increases and that the D_2 permeances are clearly larger than the H_2 ones at all temperatures. Hence, the resulting D_2/H_2 selectivity is always larger than one. This behavior is mainly due to the ZPE effects, discussed above, which make the D_2 probability threshold to be at a lower energy than the H_2 one (Fig. 3), with a consequent increase of the integral of eqn (13). However, the D_2/H_2 selectivity does not monotonously increase as temperature decreases, as it would be expected if ZPE effects were prevalent,²⁸ but reaches a maximum value of 5.6 at 47 K. This maximum is due to the increasing role of tunneling effects:²⁸ as temperature decreases, the H_2 permeances do not decrease as fast as the D_2 ones because H_2 tunnels more effectively than D_2 (see insets of Fig. 3), so at some temperature the D_2/H_2 selectivity eventually decreases. The maximum selectivity reached is close to the acceptable reference value of 6 for separation applications.⁶⁰ However, the D_2 permeance at 47 K is ≈ 0.1 GPU, rather small compared to the minimum accepted value for industrial applications, of 20 GPU.⁶⁰ It would probably be more convenient to design applications at higher temperatures, where permeances are higher and operations are easier. For instance, at the temperature of liquid nitrogen (77 K), deuterium permeance rises to ≈ 70 GPU and the selectivity still has a reasonable value of about 3.

Permeances obtained by restricting the calculations to the perpendicular incidence (eqn (15)) are also shown in Fig. 6. This approximation, here named as “3D-perp”, has been the approach followed in some of our previous works.^{30,31} It is interesting to explore its range of validity since it involves a considerable saving of computational time. It can be seen that the approximation provides a reasonable estimation of the



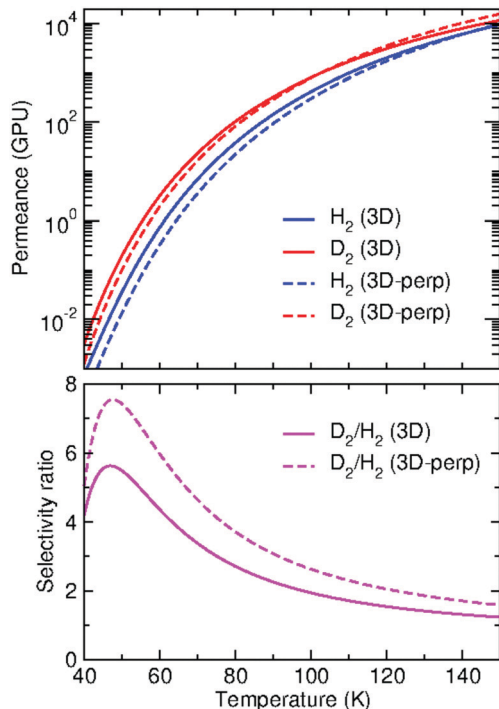


Fig. 6 Upper panel: Permeances of GDY for H₂ (blue) and D₂ (red) transport as functions of temperature (in K), obtained from the full 3D-TDWP calculations (3D, solid lines). Permeances from calculations restricted to the perpendicular incidence are also shown (3D-perp, dashed lines). 1 GPU = 3.35×10^{-10} mol m⁻² s⁻¹ Pa⁻¹. Lower panel: Ratio of the D₂/H₂ permeances (selectivity) in the same temperature range, for the 3D (solid line) and the 3D-perp (dashed line) calculations. See text for discussion.

permeances. The approximation works slightly worse for H₂, for which we find a larger underestimation of the permeances at most of the considered temperatures. As a result, the corresponding selectivity ratio predicts a more favorable separation of D₂, with a maximum value of 7.6 at 47.5 K (lower panel of Fig. 6). Still, the 3D-perp approximation gives a good estimation of the overall temperature dependence of the selectivity.

In addition, we have performed 1D-TDWP calculations of the transmission probabilities for the perpendicular incidence within the approach described in Section 2.4. The transmission of T₂ molecules has been additionally studied within this model, and the 3D-TDWP probability for the perpendicular incidence has been also computed for the sake of comparison. As mentioned above, the 1D model probabilities are scaled using the factor given by eqn (17), which requires the estimation of the effective size of the pore, A_{eff} . As in a previous work,³⁰ this quantity has been obtained by computing the area where the squared modulus of the TS wavefunction, $|\phi_0(x, y; 0)|^2$ is larger than a cutoff set to 10^{-4} . This has led to $A_{\text{eff}} = 1.08, 0.78$ and 0.65 \AA^2 for H₂, D₂ and T₂, respectively. Such probabilities, named as “1D-perp”, are compared with the 3D-perp probabilities in Fig. 7. It can be seen that the agreement between both calculations is outstanding for all energies up to the first energy threshold, which shows the importance of correcting the interaction potential along the perpendicular direction with the ZPE of the parallel degrees of

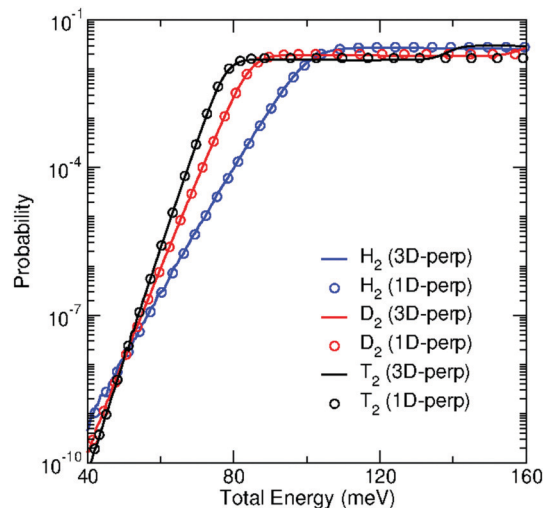


Fig. 7 Transmission probabilities through GDY of H₂ (blue), D₂ (red) and T₂ (black) for perpendicular incidence within the 3D-TDWP calculations (full lines) and the 1D model (open circles) as functions of total energy (meV).

freedom. It is specially noticeable how the 1D-perp probabilities perfectly replicate the 3D ones all along the tunneling region. The following “steps” in the probabilities at higher energies are not reproduced since higher adiabatic potential energies ($W_n(z), n > 0$) were not taken into account in the model. Nevertheless, we have checked that probabilities at these higher energies do not play a significant role in the calculation of the permeances, so 3D-perp and 1D-perp permeances are almost identical in the studied temperature range.

As a summary, the selectivities for D₂/H₂, T₂/D₂, and T₂/H₂ separation, obtained from 3D-perp calculations, are presented in Fig. 8. As expected from the relative extent of quantum effects in the three isotopes, the T₂/D₂ ratio is rather small at all temperatures whereas the T₂/H₂ selectivity becomes the largest, reaching a maximum of 21 at 45 K. Similarly to the case of the D₂/H₂ separation, the T₂ permeance is quite small at this temperature (≈ 0.05 GPU) but it rapidly increases reaching acceptable values at temperatures where the selectivity is still very promising (e.g. at 77 K, the permeance is about 90 GPU and the selectivity ratio, 7).

4 Concluding remarks

In this work we have carried out a quantum-mechanical study of the transport of hydrogen molecules through graphdiyne and the capacity of this 2D carbon membrane to separate their isotopes. A wave packet propagation method has been applied where the motion of the molecules, within the pseudoatom approximation, is simulated in the 3D space. The calculations were performed using a realistic Improved Lennard-Jones force field, which has been built on the basis of previous accurate electronic structure calculations. For the first time, transmission probabilities have been computed for a complete set of different incidence directions of the wave packets, which in turn are used



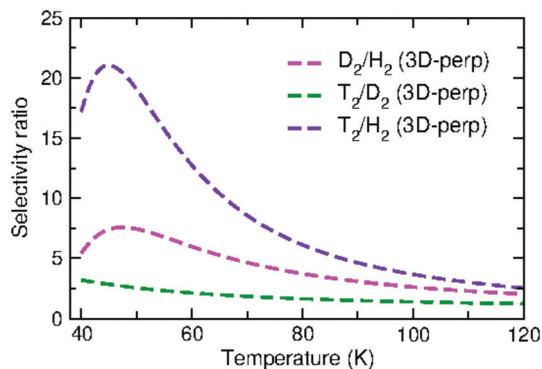


Fig. 8 Selectivity ratios for D₂/H₂, T₂/D₂, and T₂/H₂ as functions of temperature (K), as estimated from the 1D model.

to adequately estimate permeances. In this way, these calculations have served to test the reliability of less expensive computations. Firstly, it has been found that restricting the simulations to the perpendicular incidence direction gives, despite some quantitative differences, a reasonable estimation of the temperature dependence of the permeances and selectivities. Secondly, an even simpler model, using 1D effective potential energy curves, gives an excellent agreement with the 3D probabilities for the perpendicular approach, regarding both tunneling and zero-point energy effects. It would be very interesting to further explore the validity of this 1D model for other 2D nanoporous materials and molecules.

It has been found that the D₂/H₂ and T₂/H₂ selectivities reach rather high values at low temperatures (about 6 at $T \approx 50$ K and 21 at 45 K, respectively). However, at these temperatures the corresponding permeances are too small for practical experiments. It would rather be interesting to design applications at somewhat higher temperatures where the permeances values become acceptable and the selectivities are still favorable for separation of deuterium or tritium from hydrogen.

An important improvement of the present approach would be to explicitly incorporate the dependence of the interaction potentials on the relative orientation of H₂ and its rotational degrees of freedom in the wave packet dynamics (including rotationally excited states, *e.g.* *ortho*-H₂), a project planned for the near future. Another relevant aspect regards the impact of the motion of the carbon atoms of GDY on the permeances of the studied isotopes. This effect could be estimated by calculations where the positions of the C atoms are relaxed at each relative position of H₂; in such a case, a reduction of the permeation barrier is expected due to a pore deformation during molecular penetration, as found for He-/Ne-GDY by means of DFT periodic calculations.⁴⁴ An explicit account of the vibrations of the membrane as well as other interesting aspects (such as the role of different mixture ratios, thermodynamic conditions, physical adsorption, *etc.*) could be studied by classical molecular dynamics simulations.^{53–55}

This study can be extended along various interesting directions. For example, the present procedure for obtaining realistic force fields can be used to study the transport of hydrogen and other gases through other graphynes, especially graphtetrayne, which

has been recently synthesized.⁶¹ Also, in relation with recent proposals of coherent diffraction in the transmission of atoms through graphene,⁶² it could be worth exploring the diffraction pattern of atoms transmitted through graphynes, estimated by the projection of the transmitted wave packet onto well-defined wave vector states.⁶³ Work in some of these directions is in progress.

Conflicts of interest

There are no conflicts to declare.

Appendix

Here we present the computational details of the 3D-TDWP calculations outlined in Section 2.2. All calculations were carried out using an in-house code developed in our group.

The Gaussian wavepacket of eqn (4) can be written as

$$G(z; z_0, k_{z_0}, \alpha) = \left(\frac{2\text{Im}(\alpha)}{\pi\hbar} \right)^{1/4} \exp[i\alpha(z - z_0)^2/\hbar + ik_{z_0}(z - z_0)], \quad (18)$$

where z_0 and k_{z_0} are the central position and wave vector of the wave packet and α is a pure imaginary number related to the width σ of the squared modulus of G : $\sigma = (2\hbar\ln 2/\text{Im} \alpha)^{1/2}$. Typically used values are $z_0 = 17\text{--}25$ Å (a sufficiently large initial separation from the membrane) and $\sigma = 1\text{--}2$ Å. For each incidence condition we have run calculations for various k_{z_0} values to adequately represent the transmission probabilities in the chosen energy range ($E = 5\text{--}180$ meV). The wave packets were represented in the 3D space where the parallel coordinates x and y range along the unit cell and $z \in [-30:45]$ Å. The number of points in the discretization of the wave packet vary depending on the molecule: for H₂, (n_x, n_y, n_z) they are typically (96,64,512) whereas for D₂ and T₂, (128,96,768) and (192,128,1024), respectively. A time step $\delta t = 0.04$ fs was used in the application of the Split-Operator algorithm.⁴⁹ The wave packets were propagated up to a total of 3–5 ps. Larger total times were considered when the wave packet involved very low energy components of the wave packet, particularly for H₂ (up to 25 ps). To obtain the transmission probability (eqn (8)), the flux surface was placed at $z_f = -3$ Å. To prevent unphysical behavior of the wave packets at the edges of the z interval (the non-periodic coordinate) and following the method of ref. 64 and 65 they were splitted every 0.5 fs into “interaction” and “product” wave packets using the damping function $f_{\pm}(z) = \exp[-\beta(z - z_{\pm})^2]$, where f_+ and f_- are defined in the asymptotic regions $z < z_- = -10$ Å and $z > z_+ = 25$ Å, respectively, and $\beta = 3 \times 10^{-3} \text{Å}^{-2}$. After splitting, propagation is resumed using the interaction portion of the wave packet.

Many convergence checks were carried out by varying the time step, the number of grid points, the grid size along coordinate z , the absorption regions, *etc.* The computed transmission probabilities barely changed upon these modifications. In particular, it was checked in most of the calculations that halving the time step did



not produce any appreciable change in the probabilities. Slightly different probabilities (differences of about 1%) were found when enlarging the number of grid points (from (96,64,512) to (144,96,768), for instance) in the range of the largest translational energies.

Acknowledgements

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Notes and references

- J. J. M. Beenakker, V. D. Borman and S. Y. Krylov, *Chem. Phys. Lett.*, 1995, **232**, 379–382.
- J. Cai, Y. Xing and X. Zhao, *RSC Adv.*, 2012, **2**, 8579–8586.
- J. Y. Kim, H. Oh and H. R. Moon, *Adv. Mater.*, 2019, **31**, 1805293.
- L. G. Gao, R. M. Zhang, X. Xu and D. G. Truhlar, *J. Am. Chem. Soc.*, 2019, **141**, 13635–13642.
- L. M. Robeson, *J. Membr. Sci.*, 2008, **320**, 390–400.
- D.-e Jiang, V. R. Cooper and S. Dai, *Nano Lett.*, 2009, **9**, 4019–4024.
- S. P. Koenig, L. Wang, J. Pellegrino and J. S. Bunch, *Nat. Nanotechnol.*, 2012, **7**, 728–732.
- Y. Jiao, A. Du, M. Hankel and S. C. Smith, *Phys. Chem. Chem. Phys.*, 2013, **15**, 4832–4843.
- L. Wang, M. S. H. Boutilier, P. R. Kidambi, D. Jang, N. G. Hadjiconstantinou and R. Karnik, *Nat. Nanotechnol.*, 2017, **12**, 509–522.
- C. Owais, A. James, C. John, R. Dhali and R. S. Swathi, *J. Phys. Chem. B*, 2018, **122**, 5127–5146.
- G. Li, Y. Li, H. Liu, Y. Guo, Y. Li and D. Zhu, *Chem. Commun.*, 2010, **46**, 3256–3258.
- C. Huang, Y. Li, N. Wang, Y. Xue, Z. Zuo, H. Liu and Y. Li, *Chem. Rev.*, 2018, **118**, 7744–7803.
- A. James, C. John, C. Owais, S. N. Myakala, S. Chandra Shekar, J. R. Choudhuri and R. S. Swathi, *RSC Adv.*, 2018, **8**, 22998–23018.
- R. Sakamoto, N. Fukui, H. Maeda, R. Matsuoka, R. Toyoda and H. Nishihara, *Adv. Mater.*, 2019, **31**, 1804211.
- H. Qiu, M. Xue, C. Shen, Z. Zhang and W. Guo, *Adv. Mater.*, 2019, **31**, 1803772.
- J. Xu and S. Meng, *J. Phys. D: Appl. Phys.*, 2020, **53**, 493003.
- R. H. Baughman, H. Eckhardt and M. Kerte, *J. Chem. Phys.*, 1987, **87**, 6687–6699.
- Y. Jiao, A. Du, M. Hankel, Z. Zhu, V. Rudolph and S. C. Smith, *Chem. Commun.*, 2011, **47**, 11843–11845.
- S. W. Cranford and M. J. Buehler, *Carbon*, 2011, **49**, 4111–4121.
- H. Zhang, X. He, M. Zhao, M. Zhang, L. Zhao, X. Feng and Y. Luo, *J. Phys. Chem. C*, 2012, **116**, 16634–16638.
- W.-h Zhao, L.-f Yuan and J.-l Yang, *Chin. J. Chem. Phys.*, 2012, **25**, 434–440.
- H. Zhang, X. Zhao, M. Zhang, Y. Luo, G. Li and M. Zhao, *J. Phys. D: Appl. Phys.*, 2013, **46**, 495307.
- Y. Jiao, A. Du, S. C. Smith, Z. Zhu and S. Z. Qiao, *J. Mater. Chem. A*, 2015, **3**, 6767–6771.
- X. Tan, L. Kou, H. A. Tahini and S. C. Smith, *Mol. Simul.*, 2016, **42**, 573–579.
- P. Sang, L. Zhao, J. Xu, Z. Shi, S. Guo, Y. Yu, H. Zhu, Z. Yan and W. Guo, *Int. J. Hydrogen Energy*, 2017, **42**, 5168–5176.
- Q. Liu, L. Cheng and G. Liu, *Membranes*, 2020, **10**, 286.
- J. Schrier and J. McClain, *Chem. Phys. Lett.*, 2012, **521**, 118–124.
- M. I. Hernández, M. Bartolomei and J. Campos-Martínez, *J. Phys. Chem. A*, 2015, **119**, 10743–10749.
- A. T. Yinnon and R. Kosloff, *Chem. Phys. Lett.*, 1983, **102**, 216–223.
- A. Gijón, J. Campos-Martínez and M. I. Hernández, *J. Phys. Chem. C*, 2017, **121**, 19751–19757.
- M. I. Hernández, M. Bartolomei and J. Campos-Martínez, *Nanomaterials*, 2021, **11**, 73.
- S. Bhowmick, M. I. Hernández, J. Campos-Martínez and Y. V. Suleimanov, *Phys. Chem. Chem. Phys.*, 2021, **23**, 18547–18557.
- M. Bartolomei, E. Carmona-Novillo and G. Giorgi, *Carbon*, 2015, **95**, 1076–1081.
- Y. Qu, F. Li and M. Zhao, *Sci. Rep.*, 2017, **7**, 1483.
- M. Li, L. Wang, H. Lei, Y. Yang, Y.-Q. Li, M. Zhao, J. Guan, W. Li and Y. Qu, *Adv. Theory Simul.*, 2021, **n/a**, 2100327.
- W. H. Miller, N. C. Handy and J. E. Adams, *J. Chem. Phys.*, 1980, **72**, 99–112.
- J. L. Bao and D. G. Truhlar, *Chem. Soc. Rev.*, 2017, **46**, 7548–7596.
- G. M. Psfogiannakis and G. E. Froudakis, *J. Phys. Chem. C*, 2012, **116**, 19211–19214.
- F. Costanzo, P. L. Silvestrelli and F. Ancilotto, *J. Chem. Theory Comput.*, 2012, **8**, 1288–1294.
- M. Bartolomei, M. I. Hernández, J. Campos-Martínez, R. Hernández-Lamonedá and G. Giorgi, *Carbon*, 2021, **178**, 718–727.
- F. Pirani, S. Brizi, L. Roncaratti, P. Casavecchia, D. Cappelletti and F. Vecchiocattivi, *Phys. Chem. Chem. Phys.*, 2008, **10**, 5489–5503.
- A. Gavezzotti, *J. Phys. Chem. B*, 2003, **107**, 2344–2353.
- R. Cambi, D. Cappelletti, G. Liuti and F. Pirani, *J. Chem. Phys.*, 1991, **95**, 1852–1861.
- M. Bartolomei, E. Carmona-Novillo, M. I. Hernández, J. Campos-Martínez, F. Pirani and G. Giorgi, *J. Phys. Chem. C*, 2014, **118**, 29966–29972.
- M. Bartolomei, E. Carmona-Novillo, M. I. Hernández, J. Campos-Martínez, F. Pirani, G. Giorgi and K. Yamashita, *J. Phys. Chem. Lett.*, 2014, **5**, 751–755.
- J. Z. H. Zhang, *Theory and Application of Quantum Molecular Dynamics*, World Scientific, Singapore, 1999.



- 47 R. Gerber, R. Kosloff and M. Berman, *Comput. Phys. Rep.*, 1986, **5**, 61–113.
- 48 E. J. Heller, *J. Chem. Phys.*, 1975, **62**, 1544–1555.
- 49 M. D. Feit, J. A. Fleck and A. Steiger, *J. Comput. Phys.*, 1982, **47**, 412–433.
- 50 D. Zhang and J. Z. H. Zhang, *J. Chem. Phys.*, 1991, **101**, 1146–1156.
- 51 W. H. Miller, *J. Chem. Phys.*, 1974, **61**, 1823–1834.
- 52 S. Blundell and K. M. Blundell, *Concepts in Thermal Physics*, Oxford University Press, 2006.
- 53 C. Sun, M. S. H. Boutilier, H. Au, P. Poesio, B. Bai, R. Karnik and N. G. Hadjiconstantinou, *Langmuir*, 2014, **30**, 675–682.
- 54 H. Liu, S. Dai and D.-E. Jiang, *Nanoscale*, 2013, **5**, 9984–9987.
- 55 Y. B. Apriliyanto, N. Faginas Lago, A. Lombardi, S. Evangelisti, M. Bartolomei, T. Leininger and F. Pirani, *J. Phys. Chem. C*, 2018, **122**, 16195–16208.
- 56 S. Blankenburg, M. Bieri, R. Fasel, K. Müllen, C. A. Pignedoli and D. Passerone, *Small*, 2010, **6**, 2266–2271.
- 57 A. W. Hauser, J. Schrier and P. Schwerdtfeger, *J. Phys. Chem. C*, 2012, **116**, 10819–10827.
- 58 Y. Wang, J. Li, Q. Yang and C. Zhong, *ACS Appl. Mater. Interfaces*, 2016, **8**, 8694–8701.
- 59 M. Hankel, H. Zhang, T. X. Nguyen, S. K. Bhatia, S. K. Gray and S. C. Smith, *Phys. Chem. Chem. Phys.*, 2011, **13**, 7834–7844.
- 60 Z. Zhou, *J. Membr. Sci.*, 2006, **281**, 754–756.
- 61 Q. Pan, S. Chen, F. Wu, C. Shao, J. Sun, L. Sun, Z. Zhang, Y. Man, Z. Li, Y. He and L. Zhao, *CCS Chem.*, 2020, **2**, 1368.
- 62 C. Brand, M. Debiossac, T. Susi, F. Aguilon, J. Kotakoski, P. Roncin and M. Arndt, *New J. Phys.*, 2019, **21**, 033004.
- 63 M. I. Hernández, J. Campos-Martínez, S. Miret-Artés and R. D. Coalson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **49**, 8300–8309.
- 64 R. Heather and H. Metiu, *J. Chem. Phys.*, 1987, **86**, 5009–5017.
- 65 P. Pernot and W. A. Lester, *Int. J. Quantum Chem.*, 1991, **40**, 577–588.

