Inelastic effects in electron tunneling through water layers

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Calculations of tunneling matrix elements associated with electron transfer through molecular environments are usually done for given frozen nuclear configurations, with the underlying assumption being that nuclear motions are slow relative to the time scale of a tunneling event. This paper examines this issue for the case of electron tunneling through water. The motivation for this study is a recent calculation [Peskin et al., J. Chem. Phys. 111, 7558 (1999)] that indicates that electron tunneling through water may be enhanced by tunneling resonances in the range of ~1 eV below the vacuum barrier, and finds that the lifetimes of such resonances are in the 10 fs range, same order as OH stretch periods. Our calculation is based on the absorbing-boundary-conditions-Green’s-function (ABCGF) method and proceeds in two steps. First we consider the effect of a single symmetric OH-stretch mode on electron tunneling in an otherwise frozen water environment, and establish that the inelastic tunneling probability is small enough to justify an approach based on perturbation theory limited to single phonon transitions. Next we note that on the short time scale of a tunneling event, even under resonance conditions, water nuclear dynamics may be represented in the instantaneous normal modes picture. We generalize the ABCGF method to take into account low order inelastic scattering from a continuum of such harmonic normal modes. We find that near resonance the total inelastic transmission probability is of the same order as the elastic one, and may lead to an additional ~20–40% enhancement of the overall transmission in the range of up to 1 eV below the vacuum barrier. The absolute energy exchange is small, of the order of 1% of the incident electron energy. Surprisingly, we find that the main contribution to the inelastic transmission is associated with energy transfer into the rotational–librational range of the water instantaneous normal mode spectrum. © 2001 American Institute of Physics.

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I. INTRODUCTION

Electron tunneling through water is an important element in all electron transfer processes involving hydrated solutes, and in many processes that occur in water based electrochemistry. In the standard theory of electron transfer1–3 the main role played by the solvent is to affect energy matching between donor and acceptor levels by thermal fluctuations. Continuum dielectric theory provides a reasonable description of this effect; however, it obviously cannot describe the effect of solvent structure on the electronic coupling itself. In a recent series of articles4–11 (see also Ref. 12) we have addressed the problem using a molecular level description of the water environment: Water layer structures were sampled from equilibrium molecular dynamics (MD) simulations, and the electron transmission properties of these structures were analyzed numerically using a suitable pseudopotential to describe the electron–water interaction and applying the absorbing boundary condition Green function method to compute electron transmission probabilities. This numerical approach has yielded a comprehensive picture of the interplay between the electron transmission probabilities and the corresponding water structure. In particular, these studies have indicated that one possible source for lowering the effective barrier to tunneling in water is the possible involvement of transient water structures that support resonance tunneling. It was found that such resonances exist within ~1 eV below the bare vacuum barrier, and it was possible to correlate these resonances with particular water nanostructures. Furthermore, the lifetimes of these resonances were estimated to be in the order of 10 fs. The involvement of resonance tunneling in electron transmission through water has been suggested also by other workers.13–15

In the spirit of most calculations of electronic coupling elements in electron transfer theory and of earlier dielectric continuum models that disregard the water structure altogether, we have used in our previous studies of electron transmission probabilities static water structures. This static medium approximation is based on the assumption of short duration of the tunneling process, i.e., it is assumed that any transmission event is completed before substantial nuclear motion takes place. The computation of the transmission probability can therefore be done for individual static water configurations sampled from an equilibrium ensemble, and the results averaged over this ensemble. In turn, this implies that electron transmission through water is essentially an elastic process. Indeed, experimental studies of photoemission through thin water films adsorbed on metals indicate that inelastic processes associated with the water nuclear motion contribute relatively weakly at energies of order ~1 eV.16,17 Numerical simulations of subexcitation electron transmission through one to four water monolayers adsorbed on Pt(111)18 are in agreement with this observation.22 Theoretical calculations of inelastic tunneling23 similarly show...
that when sufficiently far from resonance the overall transmission is only weakly affected by inelastic processes.

The discovery\textsuperscript{11} of transient resonance supporting structures, essentially molecular size cavities, in room temperature water, and the finding that the typical lifetimes of the corresponding resonances are of the same order as vibrational periods of intramolecular water nuclear motions cast some doubt on this static medium approximation. The lifetimes of these cavities are generally long relative to the lifetimes of the corresponding resonances\textsuperscript{24} so from this point of view a static picture is valid. However, the extent to which inelastic processes contribute to the transmission probability has to be reassessed. This is the purpose of the work presented in this paper.

The model used in the present calculation is similar to that used by us in earlier studies, and is briefly described in Sec. II. In Sec. III we relax the static medium approximation in an artificial way, keeping all nuclear coordinates in the water layer static, except for one OH stretch mode associated with a single water molecule. We compute the inelastic transmission probabilities for processes that leave this mode in different vibrational states, and conclude that inelastic tunneling is dominated by single phonon transitions. In Sec. IV we calculate the 1-phonon inelastic electron transmission probability through water, invoking the short duration of the tunneling event by describing water motion in the instantaneous normal mode picture. As expected, we find that inelastic tunneling is important close to resonance energies; however, the overall energy exchange is small. As discussed in the concluding Sec. V, the overall transmission probability is still approximated well by the static medium approximation.

II. THE MODEL SYSTEM

We use the same Pt(100)–H\textsubscript{2}O–Pt(100) model system and the same electron–water pseudopotential that were used in our previous calculations;\textsuperscript{4,11} a polarizable flexible simple point charge (PFSPC) potential for the water–water interaction; Pt(100) surface with water–Pt interaction taken from the work of Spohr and Heinzinger,\textsuperscript{25,26} and an electron–water pseudopotential based on that developed by Barnett et al.,\textsuperscript{27} modified\textsuperscript{7} to include the many body interaction associated with the water electronic polarizability of water. The electron–metal pseudopotential is represented by a square barrier of height 5 eV. The total potential experienced by the electron is assumed to be a superposition of these electron–water pseudopotentials. Here, \(q\) is the electron position vector, \(\mathbf{r}\) the position of the Pt atom, and \(\mathbf{R}\) the position of the water molecule.

For each (static) water configuration obtained in this way, the (elastic) electron transmission probability is computed as follows: a potential field for the electron motion is obtained as a superposition of the square barrier representing the electron–metal interaction and the electron–water pseudopotential. This potential is projected into a cubic grid of dimension 16\(\times\)16\(\times\)400 with corresponding grid spacing 2.7713\(\times\)2.7713\(\times\)0.4 au in the \((x,y,z)\) directions (\(z\) is the tunneling direction). The electronic problem is solved on this grid by using periodic boundary conditions in the \(x\) and \(y\) directions and absorbing boundary conditions in the tunneling \((z)\) direction; see Ref. 11 for details.\textsuperscript{28} The computed transmission probabilities show distinct but configuration dependent resonance tunneling features below the bare (5 eV) work function that were traced\textsuperscript{11} to the existence of transient nanocavities in the water structure. Typical lifetimes of these resonances were found to be in the 10 fs range.

III. TRANSMISSION IN THE PRESENCE OF A SINGLE OH-STRETCH MODE

In order to gain insight on the behavior of inelastic electron tunneling in water we focus first on the dynamics of a single normal mode of water in the otherwise frozen water layer. Expecting that inelastic coupling will be most effective for the high frequency OH stretch modes,\textsuperscript{29} we “unfreeze” the dynamics of a single OH stretch mode of a single water molecule in the layer and consider the resulting inelastic contributions to the tunneling process. For an isolated water molecule whose geometry in the HOH plane is defined in terms of a 2-dimensional Cartesian coordinate system centered on the O atom, with the \(y\) axis coinciding with the bisector of the HOH angle \(\alpha\), standard procedure\textsuperscript{30} yields the relations (see Appendix A) between the OH symmetric stretch mode \(q\) and the deviations \(\delta x_1, \delta y_1\) (for H\textsubscript{11}); \(\delta x_2, \delta y_2\) (for H\textsubscript{22}); \(\delta x_0, \delta y_0\) (for O) of the H and O atoms from their equilibrium positions. If we unfreeze this normal mode for a single water molecule and let it move against the background of all other molecules, it will move in a force field that is affected by this background. In the calculation described below, whose purpose is only to gauge the magnitude of the inelastic effect, we disregard this fact. We therefore consider a Hamiltonian

\[ \hat{H} = \hat{H}_{el} + \hat{V}_q + \hat{V}_{el-W} \]

(1)

where \(\hat{H}_{el}\) is the Hamiltonian for the motion of the electron interacting with the frozen water configuration

\[ \hat{H}_{el} = \hat{K} + \hat{U}_B + \hat{V}_{el-W}. \]

(2)

where \(\hat{K}\) is the electron kinetic energy operator, \(\hat{U}_B(r)\) is the bare barrier (taken as a rectangular barrier of height 5 eV and width 10 Å) and \(\hat{V}_{el-W}(r;R)^N\) is the electron–frozen water pseudopotential. Here \(r\) and \(R^N\) denote the electron position and the frozen water configuration, respectively, and \(N\) is the number of the molecules. Also in Eq. (1) \(\hat{H}_q(q^2 + \omega^2q^2)/2\) is the Hamiltonian for the normal mode under
consideration and \( V_{el-q}(r,q) \) is the electron–normal mode interaction. An explicit form for the latter can be derived as the difference
\[
V_{el-q}(r,q) = V_{el-w}(r, R_N^{-1}x_0 + \delta x_0, y_0 + \delta y_0, x_1 + \delta x_1, y_1 + \delta y_1, x_2 + \delta x_2, y_2 + \delta y_2) - V_{el-w}(r, R_N^0),
\]
(3)
where \((R_N^{-1}x_0, y_0, x_1, y_1, x_2, y_2)\) is another notation for the underlying frozen configuration \(R_N\) of the \(N\)-water molecule system, and where the deviations \(\delta x_j, \delta y_j\) \((j = 0, 1, 2)\) of the unfrozen molecular coordinates from their reference positions may be expanded in terms of \(q\) using Eqs. (A9)–(A12) of Appendix A. The resulting interaction obviously depends on the position of the particular water molecule with the unfrozen stretch mode, and on its orientation relative to the tunneling direction, but the general conclusions reached below are independent of these details.

The numerical evaluation of the transmission probability associated with the Hamiltonian (1) is executed as a multichannel scattering problem. The electron motion is described on a spatial grid with absorbing potential set at the edges of the system in the tunneling direction, as in Ref. 11, and the normal mode \(q\) is described in the basis \((\{j\})\) of eigenfunctions of the free harmonic oscillator, defining the different channels. Because of its high frequency this mode starts out in the ground state, and inelastic tunneling therefore involves excitations of this mode to higher levels. Only open channels, i.e., with final nuclear energy not exceeding the total incoming electron energy, \(^{31}\) should be included in the calculation. There is a finite number \(n\) of such channels, with \(n\) of order \(5–10\). Denoting the total number of grid points by \(N_G\), and the number of vibrational states by \(n\), the Hamiltonian \(H\) in this representation is a matrix of order \(n \times N_G\) that can be viewed as an \(n \times N\) block matrix where each block is of order \(N_G\). The diagonal blocks correspond to the elastic motion of the electron in the different channels defined by the quantum states of the oscillator, while the nondiagonal blocks define for each grid point the coupling \(V_{\nu,\nu'}(r) = \langle \nu | V_{el-q}(r,q) | \nu' \rangle\) between the different channels \(\nu, \nu' = 0, ..., n-1\). In the same representation the wave function of the electron–oscillator system is a column consisting of \(n\) subcolumns of order \(N_G\). Each subcolumn corresponds to the electron wave function in the corresponding channel.

We focus on the one-to-all transmission problem, where the electron is incident normal to the water layer, and the total tunneling probability to exit in all final directions is calculated. The initial state that corresponds to the incoming electron is \(\Phi_{in,\nu=0} = \phi_{in}(r)|\nu=0\rangle \sim e^{ikz}|\nu=0\rangle\) (\(z\) is the transmission direction), with energy \(E_{in} = \hbar^2 k^2/2m_e\). In the grid/oscillator basis, the column vector that represents this state contains zeros everywhere except in the first subcolumn \((\nu = 0)\), which contains the grid representation of the wave function \(\exp(ikz)\). The relevant absorbing-boundary-conditions Green’s operator is
\[
\hat{G} = \frac{1}{E - \hat{H} + i \delta}
\]
(4)
where \(\hat{\varepsilon}(z) = \hat{\varepsilon}_{in}(z) + \hat{\varepsilon}_{out}(z)\) is the sum of absorbing potentials defined at the incoming and outgoing edges of the numerical grid. The total transmission probability is obtained from \(^{32,33}\) (see also Appendix B)
\[
P = \hat{\varepsilon}(\phi_{in}) \hat{G}(E_{in}) \hat{\varepsilon}_{out}(E_{in}) \hat{\varepsilon}_{in}(\phi_{in}),
\]
(5)
and the transmission probability into individual channels is given by
\[
P_\nu = \langle \psi_\nu | \hat{\varepsilon}_{out} | \psi_\nu \rangle,
\]
(6)
where \(|\psi_\nu\rangle\) is the \(\nu\)th block of scattering state supervector \(|\psi\rangle = i \hat{G}(E_{in}) \hat{\varepsilon}_{in}(\phi_{in})\).

Figure 1 displays results obtained from this calculation for one particular water configuration sampled from an equilibrium ensemble at 300 K. Two cases with different water molecules from the middle water layer with a “thawed” symmetric OH stretch were considered. In one case (“a”) the thawed molecule is chosen such that one of its OH bonds is approximately parallel to the tunneling direction. In the other (“b”) a molecule with the molecular plane perpendicular to the tunneling direction is chosen. The top full line in Fig. 1 shows the total transmission probability as a function of incident electron energy for both cases. This line coincides with the elastic transmission components in both cases because the inelastic contributions are extremely small. Results for inelastic transmission probabilities with transitions into levels \(j = 1, 2\) of the oscillator are given by the lower full lines for case \(a\) and by the lower dashed lines for case \(b\). The important observation is that, per mode, the inelastic contribution to the transmission probability in water is small, with the transmission probability for the \(0\rightarrow1\) transition smaller by about five orders of magnitudes than its elastic counterpart. Figure 2 compares similar results for the \(0\rightarrow1\) transition for water, \(D_2O\) and \(T_2O\). We find, as expected, that the inelastic coupling is stronger for the lighter isotope.
IV. TRANSMISSION THROUGH WATER DESCRIBED
BY INSTANTANEOUS NORMAL MODES

The results of the previous section indicate that the interaction of the tunneling electron with a single water molecule can be taken as a weak perturbation. The excitation of a single quantum of the symmetric OH stretch mode was seen to be a very low probability event, and the probability to induce more than a single phonon transition during the tunneling event is extremely small even near the transmission resonance. Still, since the electron interacts with a large number of modes (the normalization volume used in the calculation of Sec. III corresponds to an electron interacting with a sample of \( \sim 200 \) water molecules, i.e., with \( \sim 2000 \) nuclear degrees of freedom), the total inelastic contribution may be appreciable. In this section we provide an approximate estimate of this contribution.

The calculation described below relies on the fact, discussed above, that the time scale for the interaction of the tunneling electron with the water environment is short. The upper limit for this timescale is provided by the lifetimes of the transmission resonances, estimated to be in the order of 10 fs.\textsuperscript{11} In addition, a calculation of the traversal time for electron tunneling through water layers similar to those used in the present calculation\textsuperscript{34} yields times of the order \( \sim 1 \) fs away from the resonance energy, and again times of order \( \sim 10 \) fs near resonance. These short times are consistent with the low probability for inelastic transitions observed in the calculation of Sec. III. Still, as discussed in the introduction, nuclear motion on this time scale cannot be disregarded altogether. Instead we use previous work that indicates that on time scales shorter than \( \sim 100 \) fs, the nuclear dynamics of different solvents, including water, can be reliably described in the instantaneous normal mode picture.\textsuperscript{35–38}

In this approach, each static liquid configuration sampled from the equilibrium ensemble is used as a reference for expanding the nuclear potential up to second order in the deviations of the nuclear coordinates from this reference configuration. Diagonalizing the Hessian matrix associated with the second order derivatives of the nuclear potential about this reference point then yields a set of “instantaneous normal modes” (INMs) whose dynamics describe the short time evolution of the liquid about that configuration. In a similar spirit we assume that the inelastic contribution to electron transmission through a thin (a few molecular monolayers) water layer can be accounted for by the interaction of the tunneling electron with the instantaneous normal modes associated with the configuration encountered at each tunneling event. Previous studies have indicated that taking into account only stable (real frequency) INMs does not lead to appreciable errors in the very short time dynamics of the fluid considered, and provides a better representation of the exact fluid dynamics than an all-mode treatment at longer times.\textsuperscript{36–38}

Here we follow the same practice and represent the water dynamics using only the subset of stable INMs. In the discussion below, we refer to these modes as “phonons,” keeping in mind the limited applicability of this term in the present context.

The inelastic contribution to the transmission probability is computed as the average of results obtained in this way over configurations sampled from an equilibrium ensemble of layer configurations. Furthermore, based on the results of Sec. III we assume that the inelastic process is dominated by single phonon transitions that may be evaluated using perturbation theory. This use of perturbation theory to evaluate the inelastic contribution to electron transmission through water, combined with the description of the water nuclear dynamics as the motion of noninteracting normal modes, is what makes this problem tractable.

For any given water configuration \( \mathbf{R}_0 \) our starting point is the Hamiltonian

\[
\hat{H} = \hat{H}_{\text{el}}(\mathbf{R}_0) + \hat{H}_{\text{ph}}(\mathbf{R}_0) + \hat{V}_{\text{el-ph}}(\mathbf{r}, \mathbf{R}_0),
\]

\[
\hat{H}_{\text{el}} = \hat{K} + \hat{U}_b(\mathbf{r}) + \hat{V}_{\text{el-w}}(\mathbf{r}, \mathbf{R}_0),
\]

where \( \mathbf{r} \) is the electron coordinate, \( \hat{H}_{\text{el}} \) is the Hamiltonian associated with the motion of the electron across the static metal–water–metal junction, \( \hat{H}_{\text{ph}} \) is the phonon Hamiltonian [see Appendix B, Eq. (B5)] and \( \hat{V}_{\text{el-ph}} \) is the electron–phonon interaction that we discuss below. \( \hat{H}_{\text{el}} \) may be further written as a sum of electron kinetic energy \( \hat{K} \), the bare barrier \( \hat{U}_b \) that is modeled here as a rectangular barrier of height 5 eV, and the electron–water pseudopotential \( \hat{V}_{\text{el-w}} \) discussed in Sec. II.
For a given instantaneous water configuration, \( \hat{V}_{\text{el-ph}} \) is obtained from expanding the electron–water interaction in powers of the deviations of the water-atoms positions from this instantaneous configuration. Denoting the water configuration close to a reference configuration \( \mathbf{R}_0 \) by \( \mathbf{R} = \mathbf{R}_0 + \sum_a C_a \mathbf{q}_a \) where \( \{ \mathbf{q}_a \} \) are the instantaneous normal modes coordinates associated with that reference configuration (see Appendix B), the electron–water interaction is expanded up to linear order in \( \{ \mathbf{q}_a \} \)

\[
\hat{V}_{\text{el-w}}(\mathbf{r}, \mathbf{R}) = \hat{V}_{\text{el-w}}(\mathbf{r}, \mathbf{R}_0) + \sum_a U_a(\mathbf{r}, \mathbf{R}_0) \hat{\mathbf{q}}_a ,
\]

with \( U_a(\mathbf{r}, \mathbf{R}_0) = (\partial V_{\text{el-w}}(\mathbf{r}, \mathbf{R}_0)/\partial \mathbf{R}_0) \cdot \partial \mathbf{R}_0/\partial \mathbf{q}_a \), so that in Eq. (7)

\[
\hat{V}_{\text{el-ph}} = \sum_a U_a(\mathbf{r}, \mathbf{R}_0) \cdot \hat{\mathbf{q}}_a .
\]

In Eqs. (9) and (10) we have assumed that the order linear in the INM coordinates is sufficient. This is consistent with our expectation that the inelastic contribution to the tunneling process is dominated by single phonon transitions. One more approximation was made in the actual application of this procedure: As already discussed in Sec. II the electron–water pseudopotential used in the present work is based on that developed by Barnett \textit{et al.}\textsuperscript{27} modified to incorporate the many-body nature of the effect of the water electronic polarizability. We have found that including this aspect of the electron–water interaction is essential for getting the correct effective barrier for electron tunneling through water. However, the effect of these terms on the dynamics of inelastic tunneling is expected to be small relative to the stronger shorter-range parts of the pseudopotential, and are disregarded in the evaluation of the \( U_a \) terms. This simplifies the calculation of these terms considerably.

Next we consider the needed scattering formalism. The Hamiltonian (7) describes an electron that is moving as a free particle to the right and left of a target, i.e., \( \hat{U}, \hat{\mathbf{V}}_{\text{el-w}} \) and \( \hat{V}_{\text{el-ph}} \) are taken to vanish for \( \mathbf{r} \) outside the target. This target is a water layer superimposed on the vacuum barrier. In the process under discussion, the electron tunnels through this barrier, while interacting with the INMs bath localized in the barrier region. We denote by \( | l \rangle = | \phi_l \rangle | \chi_l \rangle \) and \( | r \rangle = | \phi_r \rangle | \chi_r \rangle \) the initial and final states of the electron–water system, where \( | \phi_l \rangle \) and \( | \phi_r \rangle \) are respectively incoming and outgoing states on the left and right sides of the barrier, while \( | \chi_l \rangle \) and \( | \chi_r \rangle \) are the initial and final nuclear states of the water target. The corresponding energies are \( E_{l,r} = E_{l,r}^{\text{el}} + E_{l,r}^{\text{ph}} \) and the relevant \( S \) matrix element is

\[
S_{l,r} = -2 \pi i T_{l,r}(E_i) \delta(E_r - E_i) .
\]

For the scattering geometry considered here, the corresponding \( T \) matrix element may be written in the form

\[
T_{l,r}(E) = \langle r | \hat{H}_{\text{RM}} \hat{G}(E) \hat{H}_{\text{LM}} | l \rangle
\]

by dividing space into three regions: The right (\( R \)) and left (\( L \)) regions of free particle motion, and a target region (\( M \)) that encloses the molecular layer. With this in mind we will sometimes write the electronic Hamiltonian in the form

\[
\hat{H}_{\text{el}} = \hat{H}_{\text{el}}^{0} + \hat{H}_{\text{LM}} + \hat{H}_{\text{RM}} ,
\]

where \( \hat{H}_{\text{el}}^{0} \) represents the electronic motion in the disconnected subspaces \( R, L, \) and \( M \), and where \( \hat{H}_{\text{RM}} \) and \( \hat{H}_{\text{LM}} \) are the couplings between these subspaces. Eq. (12) assumes that no direct coupling exists between the \( R \) and \( L \) regions. In the spatial grid representation used, \( \hat{H}_{\text{RM}} \) and \( \hat{H}_{\text{LM}} \) originate from the kinetic energy operator, and direct \( R-L \) coupling may indeed be disregarded if the target encompasses more than a few grid spacings in the tunneling direction. From the structure of Eq. (12) it follows that the Green function \( \hat{G} \) enters only through its projection on the \( M \) subspace, \( \hat{G}_{MM}(E) = \left[ E - \hat{H}_{MM} - \Sigma(E) \right]^{-1} \), where \( \hat{H}_{MM} \) is the projection of the Hamiltonian (7) on the \( M \) subspace and where \( \Sigma \) is the corresponding self-energy. \( \hat{G}_{MM} \) is a matrix of the order of the number of grid points in the \( M \) region. Note that, since \( \hat{V}_{\text{el-ph}}(\mathbf{r}, \mathbf{R}_0) \) vanishes for \( \mathbf{r} \) outside the molecular layer; \( \Sigma \) is an operator in the electronic space only.

To first order in the electron–phonon interaction we may write

\[
\hat{G}(E) = \hat{G}^{0}(E) + \hat{G}^{0}(E) \hat{V}_{\text{el-ph}} \hat{G}^{0}(E) ,
\]

\[
\hat{G}^{0}(E) = \left[ E - \hat{H}_{\text{el}} - \hat{H}_{\text{ph}} + i \eta \right]^{-1} ; \ (\eta \rightarrow 0).
\]

From the discussion above it is clear that we need only the projected part of \( \hat{G}^{0} \)

\[
\hat{G}_{MM}^{0}(E) = \left[ E - (\hat{H}_{\text{el}})_{MM} - \hat{H}_{\text{ph}} - \Sigma(E) \right]^{-1} .
\]

Of course, \( \hat{G}_{MM}^{0} \) can only couple between states \( | l \rangle \) and \( | r \rangle \) with \( | \chi_l \rangle = | \chi_r \rangle \) and, since \( \hat{H}_{\text{ph}} \) commutes with the electronic operators \( \hat{H}_{\text{el}} \) and \( \Sigma \), it can be replaced in (15) by the corresponding phonon energy. For this reason we will also encounter below the electronic Green’s function

\[
\hat{G}_{el}(E) = \left[ E - \hat{H}_{\text{el}} + i \eta \right]^{-1} ,
\]

and its \( M \)-projected part

\[
\hat{G}_{el}^{M}(E) = \left[ E - (\hat{H}_{\text{el}})_{MM} - \Sigma(E) \right]^{-1} .
\]

With these preliminaries, we are now ready to calculate the matrix element \( T_{l,r}(E) \) of Eq. (12). Below we use interchangeably the notations \( | l \rangle = | l_{\text{el-ph}} \rangle = | \phi_l \chi_l \rangle \) and (similarly for \( r \)). Using Eq. (14) we may write this coupling as a sum of elastic and inelastic components

\[
\hat{T}_{l,r}(E) = \hat{T}^{\text{elas}}_{l,r} + \hat{T}^{\text{inelas}}_{l,r} ,
\]

\[
T^{\text{elas}}_{l,r}(E) = \langle \phi_l \chi_l | \hat{H}_{LM} \hat{G}^{0}(E) \hat{H}_{RM} | \phi_r \chi_r \rangle
\]

\[
= \langle \phi_l | \hat{H}_{LM} \hat{G}_{el}(E) \hat{H}_{RM} | \phi_r \rangle \delta_{l,r} \delta_{\text{ph}} ,
\]

\[
T^{\text{inelas}}_{l,r}(E) = \sum_a \langle \phi_l | \hat{H}_{LM} \hat{G}_{el}(E) \hat{V}_{\text{el-ph}} \hat{G}^{0}(E) \hat{H}_{RM} | \phi_r \rangle \chi_a \chi^*_a
\]

In Eq. (19) Kronecker \( \delta \) term denotes that the initial (\( l_{\text{ph}} \)) and final (\( r_{\text{ph}} \)) states of the phonons are identical in the elastic process, and \( E_{\text{ph}} = E_{l_{\text{ph}}}^{\text{ph}} = E_{r_{\text{ph}}}^{\text{ph}} \) is the corresponding phonon energy. In Eq. (20) \( E_{\text{el}}^{\text{el}} = E_{l_{\text{el}}} - E_{r_{\text{el}}} \), and in each term of the \( \alpha \) summation \( E_{\text{el}}^{\text{el}} \) and \( E_{\text{el}}^{\text{el}} \) differ from each other by one
quantum of the mode $\alpha$. In addition, the structures of Eqs. (19) and (20) imply that only the M-projected parts of the Green’s operators in these equations, i.e., $\hat{G}^{(0)}_{MM}$ and $(\hat{G}_d)_{MM}$ respectively, should be considered. For simplicity of notation we drop here and below this projected notation, keeping in mind that the Green’s functions appearing below are all projected onto the M subspace.

We will focus on a particular “one-to-all” transmission probability whereupon the electron is incident from the left in the direction normal to the water layer, and the sum over all final states is considered. The one-to-all transmission probability is given by

$$P_t = \frac{2\pi}{\hbar} \sum_r |T_{tl}|^2 \delta(E_l - E_r).$$

(21)

In the presence of phonons this should be averaged over their initial distribution and summed over their final states to yield the total averaged transmission probability.

$$\langle P_t \rangle = \frac{2\pi}{\hbar} \sum_r \frac{1}{Z_L} \sum_{\ell_i} e^{-\beta E_{\ell_i}} \sum_{\ell_f} e^{-\beta E_{\ell_f}} |T_{tl}|^2 \delta(E_{\ell_f} + E_{\ell_i} - E_{\ell_f} - E_{\ell_i}).$$

(22)

Here $Z_L$ is the phonon partition function in the initial state. Note that the summations in (22) are done for a given incoming state $l_i$ of the electron, and therefore at constant $E_{\ell_i}$. Also, in what follows we shall omit the explicit average $\langle \rangle$, writing merely $P_t$ for the averaged transmission probability when the meaning is clear from the text.

Consider first the elastic tunneling component. The sum over final phonon states eliminates the $\delta_{\ell_i,\ell_f}$ term in the square of Eq. (19), and the average over initial phonon states is trivial: $Z_L \sum_{\ell_i} \exp(-\beta E_{\ell_i}) = 1$, so effectively the $|T|^2$ term becomes purely electronic. It leads (see Appendix C) to the following expression for the one-to-all transmission probability

$$P_t^{\text{elastic}}(E_{\ell_i}) = \frac{2\pi}{\hbar} \langle \phi_l | \hat{H}_{\text{eff}} | \hat{\Gamma}_M \hat{G}_{\text{el}} \hat{\Gamma}_R \hat{G}_{\text{el}} \hat{H}_{\text{ML}} | \phi_l \rangle,$$

(23)

where \( \hat{\Gamma}_R = \hat{f} [\hat{\Sigma} - \hat{\Sigma}^{(R)}] \hat{f} \). In Eqs. (21)–(23) the wave function is normalized to unit flux in the incoming wave

$$\phi_E(x,y,z) \to \sqrt{\frac{m}{\hbar^2 k_T}} e^{i k_T z} f(x,y),$$

$$E = \hbar^2 k_T^2 z^2 m + E_{xy},$$

(24)

where $E_{xy}$ is the energy in the direction normal to $z$ and $f(x,y)$ is normalized to 1. In the absorbing boundary conditions (ABC) Green’s function method the boundaries separating the left and right free electron regions $L$ and $R$ from the “molecular” region $M$ are taken far enough from the target to allow the replacement of the self energy matrix $\Sigma$ by simple position dependent imaginary potential terms $iE_{L}(r)$ and $iE_{R}(r)$ that rise smoothly towards the boundaries of the $M$ system and insure the absorption of outgoing waves at these boundaries. This leads to (see Appendix C)

$$P_t^{\text{elastic}}(E_{\ell_i}) = \frac{2\pi}{\hbar} \langle \phi_l | \hat{\Sigma} \hat{G}_{\text{el}}^d \hat{G}_{\text{el}}^e | \phi_l \rangle.$$

(25)

where $\hat{G}_{\text{el}}(E) = [E - \hat{H} + i(\hat{\varepsilon}_L + \hat{\varepsilon}_R)]^{-1}$. The numerical evaluation of this expression requires (a) evaluating the Hamiltonian matrix on the grid, and (b) evaluating the operator of the corresponding Green’s operator on the vector $| \phi_l \rangle$. In our implementation seventh order finite-differencing representation is used to evaluate the kinetic energy operator on the grid. This results in a sparse matrix representation of the Hamiltonian, suggesting the applicability of Krylov space based iterative methods for such calculations. In the present work we have used the PETSc package.

In Appendix C we develop an equivalent procedure for evaluating inelastic transmission probabilities. The result equivalent to (25) is

$$P_t^{\text{inelastic}}(E_{\ell_i}) = \sum_{\alpha=1}^{N} \left[ g_{\alpha}^+(E_{\ell_i}) h_{\alpha}^+ + g_{\alpha}^-(E_{\ell_i}) h_{\alpha}^- \right].$$

(26)

where

$$g_{\alpha}^\pm(E_{\ell_i}) = \frac{2\pi}{\hbar} \langle \phi_l | \hat{\Sigma} \hat{G}_{\text{el}}^d \hat{G}_{\text{el}}^e | \hat{U}_\alpha(r) \hat{\Sigma} \hat{G}_{\text{el}}^d \hat{G}_{\text{el}}^e | \phi_l \rangle,$$

(27)

$$h_{\alpha}^+ = \frac{\hbar}{2m_{\alpha}\omega_{\alpha}} [1 + \tilde{n}(\omega_{\alpha})],$$

$$h_{\alpha}^- = \frac{\hbar}{2m_{\alpha}\omega_{\alpha}} [1 + \tilde{n}(\omega_{\alpha})],$$

(28)

with $\tilde{n}(\omega) = \exp(\beta\omega) - 1$. Note that in the position representation $\hat{U}_\alpha(r)$ is real and diagonal, i.e., $\hat{U}_\alpha^\dagger(r) = \hat{U}_\alpha(r)$. Also, since we used mass weighted coordinates, $m_{\alpha} = 1$.

In order to compute the matrix element in Eq. (27), one needs to evaluate the vector $\hat{G}_{\text{el}}^d(E_{\ell_i}) \pm \hbar \omega_{\alpha} \hat{U}_\alpha(r) \hat{G}_{\text{el}}^e(E_{\ell_i}) \hat{\varepsilon}_L | \phi_l \rangle$. This involves two operations of Green’s matrices on a vector that are carried out as outlined below Eq. (25). Obviously, an exact calculation of the sum (26) involves too many operations of this kind to be practical. Instead we resort to a coarse-graining approximation. First rewrite Eq. (26) in the form

$$P_t^{\text{inelastic}}(E_{\ell_i}) = \int_0^\infty d\omega \rho(\omega) \left[ g^+(E_{\ell_i},\omega) h^+(\omega) + g^-(E_{\ell_i},\omega) h^-(\omega) \right],$$

(29)

where

$$g^\pm(E_{\ell_i},\omega) = \rho^{-1}(\omega) \sum_{\alpha} g_{\alpha}^\pm(E_{\ell_i}) \delta(\omega - \omega_{\alpha}),$$

(30)

and where $\rho(\omega) = \sum_{\alpha} \delta(\omega - \omega_{\alpha})$ is the density of phonon modes. The approximation is to use the following expression for $g^\pm(E_{\ell_i},\omega)$:
vollving probability for inelastic tunneling. Obviously, the term in-
approximation to $U^2(\mathbf{r},\omega)$ is obtained by dividing the $\omega$ axis into a finite number of segments $\{i\}$ of sizes $\Delta\omega_i$ and taking the average of $U^2_a$ over all modes in a segment, i.e.,
\begin{equation}
U(\mathbf{r},\omega) = \left( \frac{1}{N(\omega)} \sum_{\omega - \Delta\omega/2 < \omega < \omega + \Delta\omega/2} U^2_a(\mathbf{r}) \right)^{1/2},
\end{equation}
where $N(\omega) = \rho(\omega)\Delta\omega$ is number of modes in the segment considered.

Eq. (29) yields, under our approximations, the total probability for inelastic tunneling. Obviously, the term involving $g^+$ in this equation corresponds to processes in which the electron gained energy, while the term involving $g^-$ is associated with processes in which the electron lost energy during the transmission. These terms are integrals of the corresponding differential transmission probabilities
\begin{equation}
\mathcal{T}^{\text{inelast}}_i(E^\text{cl}_i, \pm \hbar \omega) = \rho(\omega) g_\pm(E^\text{cl}_i, \omega) \hbar^\pm(\omega).
\end{equation}
To summarize, our calculation proceeds along the following steps.

(a) Generate an equilibrium trajectory of the water layer confined between the two Pt(100) surfaces and use it to sample a desired number of water layer configurations.

(b) For any such water configuration and for an appropriate spatial grid we evaluate the grid representation of the inverse Green function $E - \hat{H}_a + i(\hat{E}_L + \hat{E}_R)$. This Green’s function can be used to calculate the elastic tunneling probability as described in Refs. 4–11.

(c) For a given water configuration we find the corresponding set of instantaneous normal modes using the potential and procedure described in Appendix B. We proceed with the subset of stable modes of real frequency.

(d) The electron–water pseudopotential is expanded in the normal modes about the given water configuration. The linear terms in this expansion yield the coupling parameters $U_a(\mathbf{r})$ according to Eqs. (9) and (10).

(e) The coarse-grained representation $U(\mathbf{r},\omega)$ of $U_a(\mathbf{r})$ is obtained using Eq. (32).

(f) For the given incoming state $|\psi_i^\text{cl}\rangle$, represented as a grid vector, we calculate the vector $\hat{G}^\text{cl}(E^\text{cl}_i, \pm \hbar \omega) \hat{U}(\mathbf{r},\omega) \hat{G}^\text{cl}(E^\text{cl}_i, \pm \hbar \omega) |\psi_i^\text{cl}\rangle$ and use it to obtain $g_\pm(E^\text{cl}_i, \omega)$, Eq. (31). The latter is then used in Eqs. (29) and (33).

V. RESULTS AND DISCUSSION

Figure 3(a) shows the density of instantaneous normal modes for the system described in Sec. II: three monolayers of normal water and of D$_2$O confined between two Pt surfaces at 300 K. Figure 3(b) show similar results for “bulk” water at 300 and 60 K, together with those obtained for the water layer. Following usual convention, the density of imaginary frequency modes is displayed on the negative $\omega$ axis. These results are averaged over 20 water configurations sampled from the corresponding equilibrium trajectories. The results for bulk water are in close agreement with those obtained previously by other workers. The results obtained for the water layer are very similar to those obtained for the bulk system of the same temperature except at the low frequency regime $\omega \approx 500$ cm$^{-1}$. In this regime the normal mode spectrum of the confined layer is seen to shift to somewhat higher frequencies, and at the same time the number of imaginary frequency modes is reduced relative to the corresponding spectral features of bulk water. Both observations reflect the increased local binding in the confined layer. The absence of appreciable effects in the higher frequency regime may be understood by noting the frequencies associated with the O–Pt stretch and the C–Pt–Pt bend of a single...
water molecule adsorbed on Pt(100) which, based on the
potential of Refs. 25 and 26 are 97 and 504 cm\(^{-1}\) respectively. Note that the relatively small density of imaginary
frequency modes justifies the approximation that disregards
them in Sec. IV.

The elastic and integrated inelastic one-to-all (incident
perpendicular direction) transmission probabilities for one
randomly chosen configuration of the 3-monolayer water
film are shown in Fig. 4. Figure 4(a) shows the elastic and
inelastic components of the transmission probabilities as well
as the total transmission probability—their sum—as func-
tions of the incident electron energy. Fig. 4(b) shows the
elastic transmission probability as well as the corresponding
inelastic components obtained for identical H\(_2\)O, D\(_2\)O, and
T\(_2\)O layers. The resonance enhancement of the transmission
probability near 4.4 eV (for the studied configuration) is seen
to be accompanied by a substantial increase in the inelastic
component. The results of Fig. 4(b) also indicate that
the importance of inelastic electron transmission through iso-
pically substituted water layers decreases in the order H\(\text{D}\)>T, as expected from the relative masses.\textsuperscript{42}

It is important to keep in mind the transient nature and the variability of the water layer structures that support such resonances. The ratio \(P_{\text{inelast}}/P_{\text{elast}}\) between the (integrated) inelastic and the elastic components of the transmission is shown for several different water configurations in Fig. 5. This figure shows the variability in the nature of the resonance supporting structure in the water configuration already discussed in Ref. 11. At the same it shows again the resonance enhancement of the inelastic tunneling component. Resonance enhancement of inelastic transition is a common feature in many scattering processes, e.g., in Raman scattering.

It is also interesting to examine the relative importance of different phonon-frequency ranges in affecting inelastic transmission. The water normal mode spectrum contains modes associated with intramolecular motions [stretching and bending modes of frequencies larger than \(1500\text{ cm}^{-1}\) (for H\(_2\)O)], and intermolecular modes of frequencies smaller than \(\approx1000\text{ cm}^{-1}\), which may be further subdivided to libration/rotation modes at the higher frequency end, and translation-dominated modes at the lower frequency end of this range. Time scale arguments would suggest that the electron couples more efficiently to the higher frequency modes associated with the intramolecular stretching and bending vibrations; however, additional factors should be taken into account. First, because the terms \(\hbar\omega\) (for electron energy loss) and \(\hbar\omega\) (for electron energy gain) in Eq. (29) are proportional to \(\bar{n}(\omega)+1\) and to \(\bar{n}(\omega)\), respectively, tunneling accompanied by electron energy loss is more important than tunneling with energy gain, particularly for high frequency modes with \(\hbar\omega>k_BT\). On the other hand, the transmission probability for an energy loss process is reduced relative to the corresponding elastic process because of the larger barrier encountered effectively in the latter process. For the combination of these reasons the low frequency regime associated with water rotations and librations is more important in affecting inelastic tunneling than the higher fre-

\[\text{FIG. 7. The energy loss/gain spectrum for the same process depicted in Fig. 6(a) at resonance (}\mathcal{E}_l^\text{el}=4.4\text{ eV) and (b) below resonance (}\mathcal{E}_l^\text{el}=3.5\text{ eV). The gain spectrum is shown on the positive}\ \omega \text{ axis (right vertical scale) while the loss spectrum is displayed on the negative}\ \omega \text{ side (left vertical scale). Dashed and dotted lines are for H}_2\text{O and D}_2\text{O respectively.}\]
for identical configurations of H$_2$O and D$_2$O layers for two values of the incident energy $E_l$. Dashed line—the energy loss integrated over all phonon frequencies $\omega$. Dotted line—same for the range that spans the intramolecular bending frequency, 1500 cm$^{-1}<\omega<2000$ cm$^{-1}$. Dashed–dotted line—same for the intramolecular stretch frequencies range, $\omega>3000$ cm$^{-1}$.

is displayed as a function of the phonon frequency $\omega$. Shown are the spectral distributions of the net energy loss computed for identical configurations of H$_2$O and D$_2$O layers for two values of the incident energy $E_l$: at resonance (4.4 eV for the configuration used), Fig. 6(a) and substantially below resonance (3.5 eV), Fig. 6(b). In both cases, the largest contribution to inelastic scattering is seen to arise from modes in the range $\omega<1000$ cm$^{-1}$, i.e., in the libration/rotation range of the normal mode spectrum of water. Figures 7(a) and 7(b) show for the same configuration and the same incident energies the two components, $(d/d\omega)(\Delta E)_{\text{gain}}$, (on the negative frequency axis) and $(d/d\omega)(\Delta E)_{\text{loss}}$, (on the positive frequency side) as functions of the phonon frequency. As expected, the contribution of the different frequency regimes to energy gain by the transmitted electron decreases rapidly with increasing mode frequency, reflecting the activated nature of the corresponding mode occupation numbers at the given (room) temperature.

Finally, Fig. 8 depicts the total energy loss, as well as some of its spectral components, for the same water configuration used in Figs. 6 and 7. The prominence of the low frequency regime ($\omega<1000$ cm$^{-1}$) in affecting inelastic tunneling is clearly seen. It should be emphasized that, while this spectral regime contains also contributions from oxygen translational modes, the dominance of hydrogenic motions is evident from the strong sensitivity to hydrogen isotope substitution seen in Figs. 6 and 7. The actual net energy loss is quite small, substantially less than 1% of the incident electron energy.

**VI. CONCLUSIONS**

In this work we have assessed the relative importance of inelastic effects in electron tunneling through water layers. It was prompted by previous studies that indicated that electron tunneling through water is enhanced by resonances associated with transient water structures characterized by molecular cavities in the water structure. A generalization of the absorbing boundary conditions Green’s function methodology makes it possible to compute first order (single phonon) corrections to the elastic tunneling probability. We have combined this with an instantaneous normal mode representation of the short time scale nuclear dynamics of the water layer in order to compute the effect of inelastic tunneling on the water transmission properties.

The results described above indicate that indeed near resonance energies inelastic tunneling cannot be disregarded. For particular configurations we find inelastic currents that exceed the elastic component. A rough average over several configurations using data such as in Fig. 5 leads to a modest increase of the total tunneling probability by 20–40% in the range of $\sim 1$ eV below the vacuum barrier due to inelastic contributions. In the deep tunneling regime (incident energy lower than 1 eV below the vacuum barrier) inelastic contributions can be disregarded as far as their effect on the overall transmission is concerned.

We have also examined the separate contributions of different parts of the instantaneous water phonon spectrum to the inelastic tunneling process. We have found that relatively low frequency modes involving hydrogen rotations and librations are the main nuclear motions affecting inelastic electron tunneling through water.

Finally, we should keep in mind that the calculation described above has focused only on transmission properties of the water layer. In a calculation of the actual tunneling current the differential transmission probability should be summed over all initial and final energies with proper account given to the Fermi population factors that determine the availability of electrons with the corresponding initial energy and the accessibility of final energy states. The differential transmission probability, Eq. (33), which is a function of both initial and final electron energies, provides, in principle, the input for such a calculation.

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**APPENDIX A**

We denote by 0, 1 and 2 the oxygen atom and the two hydrogen atoms, respectively. Using standard procedure, the relationship between $q$, the symmetric OH-stretch mode, and the deviations $\delta x_1, \delta y_1$ (for $H_{(1)}$); $\delta x_2, \delta y_2$ (for $H_{(2)}$); $\delta x_0$, respectively.
\[ \delta y_0 = \text{for O) of the H and O atoms from their equilibrium positions in the molecular plane xy can be written as follows:} \]
\[ \kappa_{11} = \frac{1}{m_H} \left( k_{\text{OH}} \sin^2 \alpha + \frac{2k_a}{r^2} \cos^2 \alpha \right), \tag{A1} \]
\[ \kappa_{22} = \frac{1}{m_H} \left( 1 + \frac{2m_H}{m_O} \right) \left( k_{\text{OH}} \cos^2 \alpha + \frac{2k_a}{r^2} \sin^2 \alpha \right), \tag{A2} \]
\[ \kappa_{12} = \kappa_{21} = \frac{1}{2} m_H \sqrt{1 + \frac{2m_H}{m_O} \left( k_{\text{OH}} - \frac{2k_a}{r^2} \right) \sin 2 \alpha}, \tag{A3} \]
\[ m_1 = 2m_H, \tag{A4} \]
\[ m_2 = 2m_H \left( 1 + \frac{2m_H}{m_O} \right), \tag{A5} \]
\[ \omega = \sqrt{\frac{1}{2} \kappa_{11} + \frac{1}{2} \kappa_{22}} \]
\[ + \sqrt{\left( \kappa_{11} + \kappa_{22} \right)^2 - 4 \left( \kappa_{11} \kappa_{22} - \kappa_{12} \right)^2} \right]^{1/2}, \tag{A6} \]
\[ c = \frac{\omega^2 - \kappa_{22}}{\kappa_{12}}, \tag{A7} \]
\[ N = \sqrt{1 + c^2}. \tag{A8} \]
where \( m_H \) and \( m_O \) are the atomic masses of the H and O atoms, \( r \) is the equilibrium OH bond length, \( \alpha \) is the equilibrium HOH angle, \( k_{\text{OH}} \) is the force constant for the OH bond, and \( k_a \) is the force constant for the angle \( \alpha \) [so that the potential for harmonic deviations of the water configuration from equilibrium is \((1/2)k_{\text{OH}}(\delta r_{12})^2 + (1/2)k_{\text{OH}}(\delta \theta_\alpha)^2 + (1/2)k_a(\delta \alpha)^2\)]. Then
\[ \delta x_0 = 0, \tag{A9} \]
\[ \delta y_0 = - \frac{2m_H}{m_O \sqrt{m_1}} \frac{1}{N} q \tag{A10} \]
\[ \delta x_1 = - \delta x_2 = \frac{C}{N \sqrt{m_1}} q, \tag{A11} \]
\[ \delta y_1 = \delta y_2 = - \frac{1}{N \sqrt{m_2}} q. \tag{A12} \]
In the preliminary study discussed in Sec. III we take the Hamiltonian for the thawed normal mode \( q \) to be the same as in a free water molecule, \( H_q = (\dot{q}^2 + \omega^2 q^2)/2 \).

**APPENDIX B**

**A. Water potential and instantaneous normal modes (INM)**

A set of instantaneous normal modes associated with a given water configuration is obtained by expanding the water potential to second order in the deviations from the given configuration
\[ V_w(R) - V_w(R_0) = -F(R_0) \cdot (R - R_0) + \frac{1}{2} (R - R_0) \cdot \mathbf{D}(R_0) \cdot (R - R_0), \tag{B1} \]
where \( R \) are mass weighted coordinates and where
\[ F_j(R) = - \frac{\partial V_w(R)}{\partial R_j}, \tag{B2} \]
\[ D_{jk}(R) = \frac{\partial^2 V_w(R)}{\partial R_j \partial R_k}, \tag{B3} \]
and diagonalizing the Hessian matrix \( \mathbf{D} \). Denoting by \( \mathbf{M} \) the corresponding transformation matrix, the squared normal modes frequencies, \( \omega_{\alpha}^2 \), are the elements of the diagonal matrix \( \Omega^2 = \mathbf{MDM}^\dagger \) and the corresponding shifted coordinates are \( \mathbf{q} = \mathbf{M} \cdot (\mathbf{R} - \mathbf{R}_0) - (\Omega^2)^{-1}\mathbf{f} \), where \( \mathbf{f} = \mathbf{M} \cdot \mathbf{F} \) is the transformed force. The short time evolution about the configuration is then determined by the Hamiltonian
\[ \mathbf{H}_{\text{ph}}(t) = \sum_a \left( (1/2) \omega_a^2 q_a^2 (1/2) \omega_a^2 q_a^2 \right), \tag{B4} \]
where \( \omega_a^2 \) are the diagonal elements of \( \Omega^2 \). In general, some of the INM’s frequencies are imaginary, and in the present tunneling calculation we have taken into account only stable modes with real frequencies that we refer to as "phonons." The starting point of the calculation outlined in Sec. IV is the second quantization representation of the Hamiltonian (B4)
\[ \mathbf{H}_{\text{ph}} = \sum_a \hbar \omega_a \left( a_a^\dagger a_a + \frac{1}{2} \right), \tag{B5} \]
where \( a_a^\dagger \) and \( a_a \) are phonon creation and annihilation operators, and where the sum is over stable INMs.

The nuclear potential for the water system is
\[ V_w = V_{wb} + V_{\text{Pt-w}}. \tag{B6} \]
Here \( V_{wb} \) is the bulk water potential while \( V_{\text{Pt-w}} \) is the water–platinum wall interaction. For the former we use the flexible simple point charge (FSPC) model\(^9,43,44\) where the potential is written as a sum of intermolecular and intramolecular contributions
\[ V_{wb} = V_w^{\text{inter}} + V_w^{\text{intra}}. \tag{B7} \]
The forms of these potentials used in the present calculation are as follows:

(a) The intermolecular water term is a sum of a pair of interactions between water atoms not belonging to the same molecule, as in the SPC water model
\[ V_w^{\text{inter}} = \sum_{i<j} V_w^{\text{inter}}(i,j), \quad \text{(sums over water pairs)} \tag{B8} \]
\[ V_w^{\text{inter}}(i,j) = \frac{q_i^2}{R_{H_iH_j}} + \frac{q_i^2}{R_{H_iH_j}} + \frac{q_i^2}{R_{H_iH_j}} + \frac{q_i^2}{R_{H_iH_j}} + \frac{q_iq_i}{R_{O_iH_j}} \]
\[ + \frac{q_iq_i}{R_{O_iH_j}} + \frac{q_iq_i}{R_{O_iH_j}} + \frac{q_iq_i}{R_{O_iH_j}} + \frac{q_iq_i}{R_{O_iH_j}} + 4 \varepsilon \left( \frac{\sigma}{R_{O_iO_j}} \right)^6 \left( \frac{\sigma}{R_{O_iO_j}} \right)^6 - 1, \tag{B9} \]
where, e.g., $R_{H_{i}H_{j}} = |R_{H_{i}} - R_{H_{j}}|$, with $R_{H_{i}}$ being the position of the hydrogen “1” on molecule $i$. The parameters used in Eq. (B9) are $q_{G} = -0.82$ a.u., $q_{H} = 0.41$ a.u., $\sigma = 3.16554 \text{ Å}$, and $\epsilon = 0.1554$ kcal/mol.

(b) The intramolecular water term is a sum over all water molecules

$$V_{W}^{\text{intr}} = \frac{1}{2} \sum_{i} n_{H_{2}O}(i)$$

(B10)

of terms of the form

$$\nu_{W}^{\text{intr}}(i) = K_{1}(d_{1i}^{2} + d_{2i}^{2} + K_{5}d_{3i}^{2}) + 2K_{3}(d_{1i} + d_{2i})d_{ai} + K_{4}(d_{1i}^{2} + d_{2i}^{2})$$

$$+ K_{6}(d_{1i} + d_{2i})d_{1i} + K_{7}(d_{1i}^{2} + d_{2i}^{2})d_{ai}$$

$$+ K_{8}d_{1i}d_{2i}d_{ai} + K_{9}(d_{1i} + d_{2i})d_{ai}^{2} + K_{10}d_{3i}^{3}$$

$$+ K_{11}(d_{1i}^{2} + d_{2i}^{2}) + K_{12}d_{1i}d_{2i}^{2} + K_{13}d_{1i}^{2}d_{2i}$$

$$+ K_{14}(d_{1i}^{2} + d_{2i}^{2})d_{ai}^{2} + K_{15}d_{1i}d_{2i}d_{ai}^{2},$$

(B11)

where

$$d_{1i} = |R_{H_{i}} - R_{O}| - R_{eq}$$

$$d_{2i} = |R_{H_{i}} - R_{O}| - R_{eq}$$

$$d_{ai} = R_{eq}\left\{\arccos\left(\frac{(R_{H_{i}} - R_{O}) \cdot (R_{H_{i}} - R_{O})}{|R_{H_{i}} - R_{O}|^2||R_{H_{i}} - R_{O}|^2 - \alpha_{eq}}\right)\right\}.$$ (B12)

The parameters used in Eqs. (B11) and (B12) are

$$K_{1}(i = 1...3) = 8.4319269, -100.73624, 759.01307 \times 10^{-11} \text{ J/Å}^2,$$

$$K_{2}(i = 4...10) = 262.1435, -159.16137, 159.58416, -658.27683, 149.60711,$$

$$-139.63423 \times 10^{-21} \text{ J/Å}^3,$$

and

$$K_{3}(i = 11...15) = 15359.793, 797.88605, 1296.5996,$$

$$-1695.5426, -498.64404 \times 10^{-21} \text{ J/Å}^4, R_{eq} = 1 \text{ Å}$$

and

$$\alpha_{eq} = 109.5^\circ.$$

(c) The water–Pt(100) interaction is taken from Refs. 25 and 26. It takes the form

$$V_{Pt-H_{2}O} = \sum_{j} \sum_{j} \nu_{Pt-H_{2}O}^{(i,j)} + \sum_{j} \nu_{Pt-H_{j}}^{(i,j)},$$

(B13)

with

$$\nu_{Pt-H_{2}O}^{(i,j)} = [ae^{-\alpha r_{ij}} - be^{-\beta r_{ij}}]f(\rho_{ij}) + ce^{-\gamma r_{ij}}[1 - f(\rho_{ij})],$$

(B14)

$$R_{ij} = |R_{i} - R_{j}|$$

(B15)

$$\rho_{ij} = |(R_{i} - R_{j}) - \hat{n}((R_{i} - R_{j}) \cdot \hat{n})|$$

(B16)

$$f(\rho) = \exp(-\rho^2),$$

(B17)

where $\hat{n}$ is a unit normal vector to the Pt wall and pointing in the outward (i.e., into the water) direction and $i$ and $j$ refer to the oxygen and Pt atoms respectively. Also

$$\nu_{Pt-H_{2}O}^{(i,j)} = g \exp(-\eta r_{ij}).$$

(B18)

The parameters used in these equations are $a = 1.8942 \times 10^{-16}$ J, $b = 1.8560 \times 10^{-16}$ J, $c = 10^{-13}$ J, $\alpha = 1.1004 \text{ Å}^{-1}$, $\beta = 1.0966 \text{ Å}^{-1}$, $\gamma = 5.3568 \text{ Å}^{-1}$, $\delta = 5.2082 \text{ Å}^{-2}$, $g = 1.7142 \times 10^{-19}$ J, and $\eta = 1.2777 \text{ Å}^{-1}$. In the equations above energy is given in J while $R$ and $\rho$ are expressed in Å.

**APPENDIX C**

Here we provide some details pertaining to the derivations of Eqs. (23), and (25)–(27). Consider first the elastic transmission rate out of electronic state $l$

$$k_{l}^{\text{elast}} = 2\pi \hbar \sum_{r} \langle \phi_{l}\hat{H}_{LM}\hat{G}_{el}(E_{l}^{cl})\hat{H}_{RM}\phi_{r} \rangle |\delta(E_{l}^{cl} - E_{r}^{cl})|^{2}.$$ (C1)

Expanding in the basis of $m$ states that span the M subspace, this can be written in the form

$$k_{l}^{\text{elast}} = 2\pi \hbar \sum_{m_{1}} \sum_{m_{2}} \sum_{m_{3}} \sum_{m_{4}} \sum_{r} \langle \phi_{l}\hat{H}_{LM}\phi_{m_{1}} \rangle \langle \hat{G}_{el}(E_{l}^{cl}) \rangle_{m_{1}m_{2}} \langle \hat{H}_{RM}\phi_{r}\phi_{m_{3}} \rangle \delta(E_{l}^{cl} - E_{r}^{cl}).$$ (C2)

Next use the identity

$$\Gamma_{m_{1}m_{2}}^{R}(E_{l}^{cl}) = i[\Sigma^{R} - \Sigma^{R}T]$$

$$= 2\pi \sum_{r} \langle \hat{H}_{RM}\phi_{r}\rangle \langle \phi_{l}\hat{H}_{LM}\phi_{r} \rangle \delta(E_{l}^{cl} - E_{r}^{cl})$$

(C3)

and do the formal sums over the $m$ states to get

$$k_{l}^{\text{elast}}(E_{l}^{cl}) = 2\pi \langle \phi_{l}\hat{H}_{LM}\hat{G}_{el}(E_{l}^{cl})\hat{H}_{RM}\phi_{l} \rangle$$

$$\langle \hat{G}_{el}(E_{l}^{cl})\hat{H}_{LM}\phi_{l} \rangle.$$ (C4)

This rate can be converted to transmission probability by dividing it by the incoming flux, $j_{i} = \hbar k_{z}/m$, where $m$ is the electron mass. Alternatively we may write the transmission probability as

$$P_{l}^{\text{elast}}(E_{l}^{cl}) = 2\pi \langle \phi_{l}\hat{H}_{LM}\hat{G}_{el}(E_{l}^{cl})\hat{H}_{RM}\phi_{l} \rangle$$

$$\langle \hat{G}_{el}(E_{l}^{cl})\hat{H}_{LM}\phi_{l} \rangle.$$ (C5)
providing that the wave function \( \phi \) is normalized to unit incoming flux, i.e.,

\[
\phi_{E}(x,y,z) \rightarrow \sqrt{\frac{m}{\hbar k}} e^{i k z} f(x,y),
\]

\[
E = \hbar^2 k^2 / 2m + E_{cv}
\]

\( E_{cv} \) is the energy in the direction normal to \( z \) and \( f(x,y) \) is normalized to 1. In the absorbing boundary conditions (ABC) Green’s function method the boundaries separating the left and right free electron regions \( L \) and \( R \) from the “molecular” region \( M \) are taken far enough from the target to allow the replacement of the self energy matrix \( S \) by a simplified term \( i \varepsilon R \) (or \( i \varepsilon_{L} \)) that rise smoothly towards the boundaries of the system and ensure the absorption of outgoing waves at these boundaries. In this case \( \hat{\Gamma}^{-1}(E) \) in Eq. (C5) may be replaced by \( 2 \varepsilon R \). A further simplification may be achieved if we use the fact that the incoming electronic state \( \phi_{l} \) is localized in the \( L \) subspace and may be taken to vanish near the RM boundary. This implies that \( \varepsilon_{l} \phi_{l} = 0 \) and \( \hat{H}_{RM} \phi_{l} = 0 \). Using these equalities we can write

\[
\hat{\varepsilon}_{l} \hat{G}_{el} \hat{H}_{LM} \phi_{l} = \hat{\varepsilon}_{l} \hat{G}_{el} \left( \hat{G}_{el}^{-1} + \hat{H}_{LM} + \hat{H}_{RM} \right) \phi_{l}.
\]

Together with \( \left[ \hat{G}_{el}^{-1}(E_{l}) \right]^{-1} = E_{l} - \hat{H}_{0}^{\dagger} - \hat{H}_{LM} - i \varepsilon \) and \( (E_{l}^{\dagger} - \hat{H}_{0}^{\dagger}) \phi_{l} = 0 \) this yields

\[
\hat{\varepsilon}_{l} \hat{G}_{el} \hat{H}_{LM} \phi_{l} = \hat{\varepsilon}_{l} \hat{G}_{el} \hat{G}_{el}^{-1} \phi_{l}
\]

so that

\[
P_{l}^{\text{elastic}}(E_{l}^{\dagger}) = \frac{2}{\hbar} \sum_{\alpha} \frac{h}{2m \omega_{\alpha}} \left\{ \bar{n}_{\alpha} \langle \phi_{l} | \hat{H}_{LM} \hat{G}_{el} (E_{l}^{\dagger}) \hat{U}_{\alpha}^{\dagger}(r) \hat{G}_{el} (E_{l}^{\dagger}) \phi_{l} \rangle \right\}.
\]

Consider next the inelastic transmission probability. Inserting Eq. (20) into (22) yields the inelastic rate

\[
k_{l}^{\text{inelastic}}(E_{l}^{\dagger}) = \frac{2 \pi}{\hbar} \sum_{\alpha} \frac{h}{2m \omega_{\alpha}} \left\{ \bar{n}_{\alpha} \langle \phi_{l} | \hat{H}_{LM} \hat{G}_{el} (E_{l}^{\dagger}) \hat{U}_{\alpha}^{\dagger}(r) \hat{G}_{el} (E_{l}^{\dagger}) \phi_{l} \rangle \right\}.
\]

Finally, using the ABC Green’s function methodology and the arguments that lead to Eq. (C9) we now get

\[
P_{l}^{\text{elastic}}(E_{l}^{\dagger}) = \sum_{\alpha=1}^{N} \left[ g_{\alpha}^{+}(E_{l}^{\dagger}) h_{\alpha}^{+} + g_{\alpha}^{-}(E_{l}^{\dagger}) h_{\alpha}^{-} \right],
\]

where

\[
g_{\alpha}^{+}(E_{l}^{\dagger}) = \frac{2}{\hbar} \sum_{\alpha} \frac{h}{2m \omega_{\alpha}} \left\{ \bar{n}_{\alpha} \langle \phi_{l} | \hat{H}_{LM} \hat{G}_{el} (E_{l}^{\dagger}) \hat{U}_{\alpha}^{\dagger}(r) \hat{G}_{el} (E_{l}^{\dagger}) \phi_{l} \rangle \right\},
\]

and where in Eq. (C14) the functions \( \langle \phi_{l} \rangle \) are normalized to unit flux, as in Eq. (C6). Note that in our mass-weighted coordinate system \( m_{\alpha} = 1 \) for all \( \alpha \).
It should be kept in mind that in energy transfer from the transmitting electron to water nuclear degrees of freedom, the mechanism responsible for capturing and localizing the electron as a solvated species must play an important role for thicker layers (Refs. 19–21).


22. This grid was used in the calculation described in Sec. IV. The preliminary calculation described in Sec. III is more computationally demanding and we have therefore sacrificed some accuracy by using a smaller grid: $16 \times 16 \times 151$ points with grid spacings $2.78 \times 2.78 \times 0.375$ Å corresponding to grid dimensions $23.54 \times 23.54 \times 29.96$ Å.

23. While this is probably true, we find in Sec. V that it does not imply that high frequency internal modes are the main promoters of inelastic electron tunneling through water.


25. In actual calculations of tunneling in a metal–molecule–metal junction the number of open channels at zero temperature is determined by the maximum energy that the electron can release, which in turn depends on the potential drop (difference in Fermi energies) between the two sides.


35. The bulk water configurations were obtained from a simulation of a system with 256 molecules in a cubic box of volume $7.6 \times 10^{-21}$ cm$^3$ using periodic boundary conditions with the minimum image convention.

36. The ratios computed are of the order of the square roots corresponding mass ratios, but depend to some extent on the incident energy and on the chosen molecule.

