A physical interpretation of the collinear reactive scattering resonances in the F+HD, H₂, and D₂ systems

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A simple model is presented that explains the main characteristics of the low energy resonances found in accurate quantum mechanical scattering calculations of collinear reactive collisions of the type F + XY → FX + Y, where X and Y are H or D atoms. The wave function of the resonance complex can be approximately described by a product of a function of the F-XY distance and a vibrationally adiabatic function of the X-Y distance. The corresponding vibrational eigenvalues of the XY diatomic as a function of the F-XY distance form an attractive, effective one-dimensional potential for the F-XY motion that supports a quasibound state. The resulting resonance is broadened by its interaction with the reagent and product scattering states. The resonance energies given by the model are in good agreement with those obtained by exact scattering calculations for the F + HD, H₂, and D₂ systems.

I. INTRODUCTION

There has been a considerable effort in recent years directed toward the understanding of the dynamics of the F + H₂ reaction.1-4 Besides being important technologically for the design of chemical lasers, this reaction is one of the simplest examples of a highly exothermic chemical reaction with a low activation barrier and can be regarded as the prototype model for understanding the dynamics of such reactions. The relatively small number of electrons in the F + H₂ system facilitates the calculation of the ab initio potential surfaces for the nuclear motion. In recent years, several extensive calculations of this type for the ground electronic state potential energy surface have been performed2-5 which determine the surface to within less than 1 kcal/mol. Further refinements to ~0.1 kcal/mol, an accuracy already attained for the H + H₂ reaction in the regions of the surface most important for the dynamics,6,7 can be expected with the advancement in the ab initio techniques in the next few years.

The dynamics of the F + H₂ reaction, and of the isotopic variations in which one or both of the H atoms are replaced by D atoms, have been studied extensively using classical trajectories for both the collinear and the three-dimensional cases.6-18 The results have been used to develop semiempirical potential energy surfaces for which the dynamical studies give rate constants in good agreement with the experimental ones.5,6,10,16 Most such semiempirical surfaces are in good qualitative agreement with the ab initio ones, which gives some confidence in the correctness of the main features of the surface.

Several collinear quantum mechanical scattering calculations have been performed using semiempirical potential energy surfaces for F + H₂ and its isotopic variations18-27 as well as an analytic fit to an ab initio surface.21 The comparison of the quantum with the classical and the semiclassial results18 revealed the existence and gave some indication of the importance of various quantum effects. The large exothermicity of the reaction and the number of accessible quantum states of the products due to it makes accurate three-dimensional (3D) coupled-channel calculations, and the possibility of direct comparison of theory and experiment, difficult at the present time. However, approximate 3D calculations of the integral state-to-state cross sections of F + H₂24 and of the relative vibrational product distributions25 have been performed, which gives some insight into the 3D dynamics of the system.

Crossed molecular beam studies of the detailed dynamics of the F + D₂26 and F + H₂27 reactions have also been performed. For both reactions, vibrationally resolved angular distributions of the products have been obtained which provide detailed data for testing theoretical models.

A number of accurate collinear quantum mechanical scattering calculations on the F + H₂ reactions and its isotopic variations18-23 on different potential energy surfaces have shown the existence of narrow peaks in the state-to-state or total reactive transition probability vs energy curves. These peaks are accompanied by rapid changes in the phases of the scattering matrix as a function of the energy as well as in its eigenphases.28 A recent collision lifetime matrix analysis of accurate coupled-channel calculations, performed in our laboratory, has shown that one single eigenvalue of that lifetime matrix, when plotted as a function of energy, displays a pronounced positive maximum.29 These features have been interpreted28-31 as resonance effects due to the formation of a long-lived, almost bound triatomic complex. Closely related behavior can be found24 in the approximate 3D partial wave results for the F + H₂ reaction on the Muckerman V LEPS surface, indicating that the formation of a long-lived collinear or nearly collinear trilatropic complex is likely to play an important role in the reaction 3D collision dynamics of this system. Exact partial wave calculations have suggested that the formation of a collinear complex plays an important role in the H + H₂ 3D collision dynamics.30 No accurate 3D calculations of the differential state-to-state cross sections for the F + H₂ or any other reaction in the energy region in which resonances can be formed.
have been reported, and it is not known how the details of the angular distribution are affected by the existence and the characteristics of the long-lived complex. The results of the crossed molecular beam experiments for \( \text{F + D}_2 \)\(^{35}\) and \( \text{F + H}_2 \)\(^{37}\) show a broadening of the angular distribution at energies at which the formation of a long-lived complex can be expected. An angular distribution of the reaction products symmetric around a center-of-mass scattering angle of 90° is a consequence of the existence of a quasibound state that exists for a full rotation of the complex or longer.\(^{31}\) More conclusive evidence of the existence of a complex, especially a short-lived one, and more importantly, information about the nature of the interatomic forces that cause it, can be obtained by learning more about the correlation between the resonance characteristics and the resulting observable experimental quantities, especially the collisional energy dependence of the reaction products angular distributions.

The long-lived complexes responsible for the resonance phenomena obtained from quantum mechanical calculations are normally located in the strong interaction region of the potential energy surface in which all three atoms are close to each other. Consequently, the characteristics of the resonance features are determined mainly by the structure of the potential energy surface in that region of configuration space, and therefore, are capable of providing information about its details. Study of the dynamics of these complexes and of their correlation with the features of the potential energy surface in the strong interaction region and with their experimental manifestation can provide a tool for the experimental determination of interatomic forces at small interatomic distances in systems that form resonant states.

Understanding the nature of a long-lived complex in a collinear collision and developing simple models that are capable of simulating its main dynamical features and of approximating its energy and lifetime is the first step in establishing such a correlation. Since accurate solutions of the Schrödinger equation for a reactive system at energies at which resonances occur are available for collinear collisions, the validity of approximate models can be directly tested for a collinearly restricted complex. An extension to a noncollinearly restricted model for collinearly dominated potential energy surfaces, which seems to describe both the \( \text{H + H}_2 \) and \( \text{F + H}_2 \) reactions, can further be developed.

The dynamics of the formation of quasibound states in a collinear atom–diatomic molecule collision is somewhat simpler in the case when no reaction can occur, i.e., for inelastic collisions. Resonances in collinear inelastic collisions have been studied on both simplified models\(^{35,33}\) and realistic potential energy surfaces,\(^{34,35}\) and highly accurate simple approximate models have been developed for such nonreactive systems.\(^{32–35}\) The long-lived complex has two degrees of freedom and, for the systems for which good approximate models have been developed, its dynamics can be described by that of a two-dimensional oscillator with two weakly coupled modes. One of the modes can be closely approximated by the vibration of the isolated diatomic molecule that participates in the collision.

The dynamics of a reactive collinear system are somewhat more complex. There are two different kinds of diatomic molecule vibrations, those of the reactant and those of the product molecule, and they appear to be strongly mixed in the strong interaction region of the potential energy surface. The quantum mechanical approximate models used for studying resonances in collinear reactive collisions are usually based on some kind of coordinates that transform smoothly from those appropriate for reactants to those appropriate for products.\(^{25,36–38}\)

Significant progress has also been attained in developing semiclassical procedures for locating and describing the dynamics of resonances in both inelastic\(^{35,39}\) and reactive collisions.\(^{40}\)

The model used for describing the resonances in inelastic scattering,\(^{35}\) in which one of the modes of the quasibound complex is assumed to be equivalent to the asymptotic vibration of the colliding molecule, has been used to explain the resonances in the \( \text{I + H}_2 \) system on an approximate (LEPS) potential energy surface.\(^{41}\) A significant feature of this surface is that it has wells in both reactant and product channel regions well separated from each other. The dynamics of the quasibound complexes in those wells is similar to the dynamics of the reactants and of the products, and the model has given excellent approximations for the resonance energies. However, these wells are probably due to an artifact of the LEPS functional form used and are nonphysical.

It was found that resonance peaks in the collinear \( \text{F + H}_2 \) system appear in the vicinity of and are closely related to the opening of vibrational channels.\(^{15}\) The resonance features are strongly influenced by the details of the potential energy surface; slightly different surfaces lead to completely different resonance structures.\(^{21}\) Recently, a detailed analysis of the reaction dynamics in terms of curvilinear coordinates that smoothly change from reactants to products\(^{42}\) showed that no single-channel model in those coordinates can provide a satisfactory explanation for the nature of the resonance.

The proposed surfaces for the \( \text{F + H}_2 \) reaction do not have wells in the reactant or product regions, and since the barrier at the saddle point is low, there appears to be a smooth passage for the system to go from reactants to products, making it possible for the asymptotic reactant and product vibrations to blend into each other in the interaction region and to lose their distinction. The appreciable reactive transition probabilities for most of the isotopic variations of \( \text{F + H}_2 \) reinforce such an interpretation.

On the other hand, the numerical results\(^ {16}\) for the collinear \( \text{F + HD} \rightarrow \text{FH + D} \) reaction on the Muckerman V potential energy surface show a narrow resonance peak in the middle of a wide energy range on which the scattering is almost purely elastic. The obvious conclusion is that despite the smooth shape of the surface...
there is not much mixing between the reactant and the product states during the collision in this energy range and that the vibrational states of the reactants and the products, although probably subject to some distortion, retain their distinction and character throughout the collision. One of the distorted vibrational states of the reactants or of the products, is, hence, likely to be a distinct mode of the motion of the quasibound complex that is responsible for the resonance.

A comparison among the numerical reaction probability results for the $F + HD$, $F + H_2$, and $F + D_2$ systems reveals some common features in the corresponding low energy resonance peaks and indicates that a similar mechanism is likely to be responsible for the lowest energy resonances in all three systems. In this paper, a model of this low-energy resonance based on the adiabaticity of the ground vibrational state of the reactants during a collision between the reactants is developed and tested against coupled-channel calculations on the Muckerman V LEPS potential energy surface, which has no wells. The main reason for this choice is that accurate collinear quantum mechanical collision probability results for four isotopic variations of the reaction are available for this surface.\(^{18}\)

The details of the approach and its mathematical formulation are given in Sec. II. The model for the lowest energy resonance, the results obtained from it, and their comparison with accurate coupled-channel calculations are given in Sec. III. The consequences of the model and their relevance for the study of the actual $F + H_2$ 3D reaction dynamics are discussed in Sec. IV. In Sec. V, we summarize the results and conclusions.

II. MATHEMATICAL MODEL

The Schrödinger equation for the collinear $A + BC - AB + C$ reaction along the $BC$ axis can be written in terms of the Jacobi coordinates for the reactants or products as

$$
\left[ \frac{\hbar^2}{2m_A} \frac{\partial^2}{\partial r_A^2} - \frac{\hbar^2}{2M} \frac{\partial^2}{\partial r_B^2} + V(r_A, R) \right] \psi(R, r_A, r_B) = E \psi(R, r_A, r_B),
$$

$$
\left( \lambda = \alpha, \gamma \right),
$$

(2.1)

where $r_A$ is the BC distance, $R_B$ is the distance of A to the center of mass of BC, $m_A$ is the reduced mass of the BC pair, $M$ is the reduced mass for the motion between $A$ and BC, $V$ is the potential energy function for the system, and $\phi$ its energy eigenfunction. The same symbols with subscript $\gamma$ denote the corresponding quantities for the reverse reaction. In Delves scaled coordinates\(^{42}\)

$$
R = a_1 R_A, \quad \gamma = a_1^2 \gamma_A, \quad a_1 = (M/\sqrt{m_A})^{1/4},
$$

(2.2)

Eq. (2.1) becomes

$$
\left[ \frac{\hbar^2}{2\mu} \frac{\partial^2}{\partial \gamma^2} + V(r_\lambda, R) \right] \psi(R, r_\lambda) = E \psi(R, r_\lambda),
$$

(2.3)

where

$$
\mu = \left( \frac{m_A + m_B + m_C}{m_A + m_B} \right)^{1/2}
$$

is an effective reduced mass for the system, $m_X$ being the mass of atom $X$ ($= A, B, C$).

At large $R$, the potential $V(r_\lambda, R)$ in Eq. (2.3) does not depend on $R$, in the $\lambda$ arrangement channel region of configuration space, and the solutions of that equation in that region are of the form

$$
\psi(R, r_\lambda) = \exp(\pm ikR),
$$

(2.4)

where $\phi_\lambda^k(R)$ is the $\lambda$th vibrational wave function for the unperturbed molecule $AB$ or $BC$, i.e., the solution of

$$
\left[ -\frac{\hbar^2}{2\mu} \frac{\partial^2}{\partial R^2} + V(r_\lambda, R) \right] \phi_\lambda^k(R) = \epsilon_\lambda^k \phi_\lambda^k(R),
$$

(2.5)

and $\epsilon_\lambda^k(R)$ is the free particle wave function for the motion of the center of mass of $BC$ relative to $A$ or $AB$ relative to $C$,\(^{18}\)

$$
\epsilon_\lambda^k(R) = \exp(\pm ikR); \quad k_\lambda^\pm = \left[ 2\mu (E - \epsilon_\lambda^k) \right]^{1/2},
$$

(2.6)

In Eqs. (2.5) and (2.6), $\epsilon_\lambda^k$ is the vibrational energy of the isolated diatomic molecule $AB$ or $BC$.

For smaller values of $R$, the vibrational eigenfunctions of the diatomic molecule are perturbed by the presence of the third atom at a distance $R$. The perturbed wave function $\phi_\lambda^k(r; R)$ can be obtained by solving Eq. (2.3) at a fixed value of $R$,

$$
\left[ -\frac{\hbar^2}{2\mu} \frac{\partial^2}{\partial \gamma^2} + V(r_\lambda, R) \right] \phi_\lambda^k(r; R) = \epsilon_\lambda^k \phi_\lambda^k(r; R),
$$

(2.7)

with the boundary conditions

$$
\phi_\lambda^k(0; R) = \phi_\lambda^k(r_{\max}; R) = 0,
$$

(2.8)

where $r_{\max}$ is the value of $r_\lambda$ that corresponds to the configuration in which $B$ coincides with $A$ (for $\lambda = \alpha$) or $C$ (for $\lambda = \gamma$). These eigenfunctions $\phi_\lambda^k$ form a complete set of real orthogonal functions in the $r_\lambda$ space.

At low collision energies, the vibrational motion of BC (or AB) along $r_\lambda$ in the $R_\lambda$, $r_\lambda$ configuration space may be much faster than the relative motion of $A$ and BC (or AB and C) along $R$, and an effective adiabatic separation of these two modes of motion can be expected.\(^{37,43-44}\) In such cases, one can approximate the solution of Eq. (2.3) at any $R$ by an expression that reduces to Eq. (2.4) at large $R$, namely,

$$
\psi(r_\lambda, R) \approx \phi_\lambda^k(r_\lambda; R) \phi_\lambda^k(r_\gamma; R),
$$

(2.9)

where $\phi_\lambda^k(R)$ is the wave function for the motion along $R_\lambda$ on the effective potential determined by the $\epsilon_\lambda^k(R)$ of Eq. (2.7).

In the regions in which the potential is nearly separable in terms of $R_\lambda$ and $r_\lambda$, such as the reactant (or the product) valley of the potential (see Fig. 1) outside the interaction region, Eq. (2.9) will obviously be valid. In the strong interaction region where the potential is not nearly separable, Eq. (2.9) will be valid only if there is a good separation of the velocities along $r_\lambda$ and $R_\lambda$, and if none of the $\lambda' \neq \lambda$ states $\phi_\lambda^k(r_\lambda; R)$ is nearly degenerate with the $\lambda$ state $\phi_\lambda^k(r_\lambda; R)$ that appears in that equation. The separation of velocities condition is usually satisfied if $M/\sqrt{m_A}$ is much larger than unity.

The nature of the approximations used in assuming Eq. (2.9) can be examined more systematically by using
FIG. 1. Contour plot of the Muckerman V potential energy surface for the F + HD, F + H₂, F + D₂, and F + DH systems in the Delvsa scaled coordinates $R_a$, $r_a$. $\theta$ represents the skew angle between the $R_a$ and $R_\alpha$ axes and $\mu$ the effective mass of Eq. (2.3) in units of the H atom mass. The zero of energy is the bottom of the isolated hydrogenic molecule well. The energies of the contours are from −1.154 to 2.446 eV, uniformly spaced by 0.3 eV. The x mark indicates the position of the saddle point. The vertical lines on the F + H₂ plot denote the cuts for which the profile of the surface is shown on Fig. 2.

the $\phi^j_i$ from Eq. (2.7) as a basis set for expanding the solution of Eq. (2.3)

$$\psi(r_\alpha, R_\alpha) = \sum_j g^j_i(R_\alpha) \phi^j_i(r_\alpha; R_\alpha).$$  \hspace{1cm} (2.10)

Substituting Eq. (2.10) in Eq. (2.3), premultiplying by $\phi^j_i(r_\alpha; R_\alpha)$ and integrating over $r_\alpha$, one gets a set of coupled ordinary differential equations for the coefficient wave functions $g^j_i(R_\alpha)$

$$\frac{-\hbar^2}{2\mu} \frac{d^2 g^j_i}{dR_\alpha^2} = \frac{-\hbar^2}{2\mu} \sum_j \left[ 2\langle \phi^j_i | \frac{d}{dR_\alpha} | \phi^j_i \rangle \frac{d}{dR_\alpha} + \langle \phi^j_i | \frac{d^2}{dR_\alpha^2} | \phi^j_i \rangle \right]$$

$$\times g^j_i = (E - \epsilon^j_i) g^j_i \quad i = 0, 1, 2, \ldots.$$  \hspace{1cm} (2.11)

Equations (2.11) constitute an infinite set of coupled ordinary differential equations fully equivalent to Eq. (2.3). By solving them and imposing the correct scattering boundary conditions, one can, in principle, obtain the full scattering solution of Eq. (2.3).

More importantly, however, Eq. (2.11) can be used as a starting point for developing simple approximations. If the elements that couple the state $i$ with all the others are small in or near the $R_\alpha$ region of classical motion in state $i$ [i.e., where $\epsilon^j_i(R_\alpha) \approx E$], the $g^j_i(R_\alpha)$ can be viewed as a weakly perturbed solution of, and successfully approximated by, the solution of the equation

obtained from Eq. (2.11) by neglecting the coupling terms.

The potential energy function $V(r_a, R_s)$ which through Eq. (2.7) generates the effective potential $\epsilon_l(R_s)$ in Eqs. (2.11) and (2.12) is, for large values of $R_s$, a double minimum potential. For $\lambda = \alpha$ (see Fig. 2 with $R_s = 6.27$ bohr, for example), the well at the left resembles the vibrational potential of the isolated diatomic reactant molecule BC and becomes equivalent to it as $R_s \rightarrow \infty$. The well at the right represents the products region of the potential energy surface; the coordinate $r_a$ is, however, not appropriate for the product vibration (which is described by $\lambda = \gamma$) so that the motion in that part of the potential along $r_a$ is not equivalent to that vibration. Only the solutions that are localized in the left well of the potential $V(r_a, R_s)$ evolve adiabatically from a particular vibrational state of the reactants and have the physical meaning of quasi-stationary vibrational states of the BC molecule perturbed by the presence of the atom A. The remaining solutions, localized in the right well of the double-well potential [Fig. 2(c)], do not correspond to stationary states of the products at large $R_s$ and are included in Eq. (2.10) only for the purpose of making the basis set given by Eq. (2.7) complete and hence, making Eq. (2.11) rigorously equivalent to the Schrödinger equation.

The effective adiabatic potential $\epsilon_l^a(R_s)$ for the motion along $R_s$ in Eq. (2.12) determined by the eigenvalues of the perturbed vibrational Hamiltonian in Eq. (2.7) can have a minimum at a finite value of $R_s$ (see Fig. 3). As a result, the eigenvalues $\epsilon_l^a(R_s)$, which appear in Eq. (2.12), will either support bound states or exhibit resonances at some energies. If the coupling of $\epsilon_l^a(R_s)$ to the other states in Eq. (2.11) is weak, the corresponding solution of Eq. (2.11) will also have a resonance at a nearby energy. The energy of the resonance or of the bound state of Eq. (2.12) would, in such a case, be a good approximation to the energy of the resonance obtained from an accurate solution of Eq. (2.11), i.e., to the resonance in the solution of the Schrödinger Eq. (2.3). The model resonance given by Eq. (2.12) may be shifted with respect to the actual resonance and it may have a different width. For narrow resonances, the size of the shift would be a good indication of the correctness of the model. The width of the actual resonance can typically be expected to be larger than the width of the model resonance as a result of the additional broadening due to the nonadiabatic coupling to other open states.

### III. RESULTS

In this section, we analyze the dynamics of the lowest energy resonance in the $F$ + $Y$ reaction, where $X$ and $Y$ are $H$ or $D$ atoms, in terms of the model of the previous section, which is based on the adiabaticity of the $XY$ vibration. We use the Muckerman V potential energy surface. Its parameters as well as the values of the physical constants used are the same as those of Ref. 18,43.

The potential energy surface for the reaction is shown in Fig. 1 for each of the four isotopic systems FHD, FH$_2$, FD$_2$, and FDH. The corresponding skew angles are 37.3°, 46.4°, 47.7°, and 56.7°, respectively, and the effective masses $\mu_e$ in units of the H atom mass are 1.31, 0.95, 1.82, and 1.31, respectively. The differences among these angles and masses are ultimately responsible for the differences among the characteristic parameters of the resonances in these four systems.

The eigenvalues $\epsilon_l^a(R_s)$ of Eq. (2.7) for the potential $V(r_a, R_s)$ at fixed values of $R_s$ were evaluated for each of the four isotopic systems as a function of the parameter $\tau_a$ using the finite element method of Malik et al. 19 For the $F$ + $H_2$ system, they are shown in Fig. 2 for a few values of $R_s$ together with plots of $V(r_a, R_s)$ versus $r_a$. The variation of $\epsilon_l^a(R_s)$ with $R_s$ for some $i$ is shown in Fig. 3 for all four isotopic reactions. The eigenvalue $\epsilon_l^a(R_s)$, which correlates with the ground vibrational state of the unperturbed $XY$ molecule at large $R_s$ and is used to construct the adiabatic model for the resonance, is shown as a bold curve on that figure.

Equation (2.12) with $i = 0$ was solved numerically using Gordon's method 44 to obtain the scattering phase shifts $\eta$ at large $R_s$. The dependence of $\eta$ on energy is shown in the upper part of Fig. 4 for the $F$ + $H_2$ reaction. The rapid increase in $\eta$ over a narrow energy range results in a sharp peak in the delay time $D = 2h(\partial\eta/\partial E)$ vs energy curve at the bottom of that figure. These features indicate the existence of a long-lived complex whose motion is trapped by the $\epsilon_l^a(R_s)$ well of Fig. 3(b). The right lower ordinate scale in Fig. 4 gives the delay time in units of the fundamental vibration period of an $X_a$-motion harmonic oscillator whose zero-point energy is equal to the model resonance energy of the $\epsilon_l^a(R_s)$ well, indicated by the solid horizontal line in Fig. 3(b). The left lower ordinate scale of Fig. 4 is in units of an $r_a$-motion harmonic oscillator fundamental vibration period whose zero-point energy is equal to that of the $V(R_s, r_a)$ potential, where $R_s$ is the value of $R_s$ corresponding to the bottom of the $\epsilon_l^a(R_s)$ well. For the $F$ + $HD$ and $F$ + $D_2$ systems, rapid increases of $\eta$ with energy over a narrow energy range are also observed. The model resonance energies of the solutions of Eq. (2.12) are determined by the position of the maximum of the peak of the delay time vs energy curve analogous to Fig. 4 for each of the isotopic systems $F$ + $HD$, $F$ + $H_2$, and $F$ + $D_2$.

The model resonance width for $F$ + $H_2$ and $F$ + $D_2$ are determined from the full width at half-maximum (FWHM) of the delay time vs energy curves. For the $F$ + $HD$ system, the thickness of the barrier in the effective potential $\epsilon_l^a(R_s)$ at the model resonance energy of 0.237 eV is considerably larger than that at the exact resonance energy of 0.245 eV, and the FWHM of the delay time vs energy peak would furnish an artificially narrow width. A better estimate of the adiabatic part of this width can be obtained semiclassically. There are accurate uniform semiclassical formulas 44 for the width of the resonances of one-dimensional potentials such as that appearing in Eq. (2.12) in which the resonance energy enters as a parameter, and therefore, that width can be calculated at the exact resonance en-
ergy rather than at the model one. For the present purposes, one of the simplest semiclassical expressions is sufficiently accurate and is used. It is given by

$$\Gamma = \hbar \nu_a^2 \ln(1 + e^{2Q^a})$$

(4.1)

where $\nu_a^2$ is the $R_a$ vibrational frequency, taken here as the fundamental vibration frequency of an $R_a$ motion harmonic oscillator whose zero-point energy is equal to the exact resonance energy measured with respect to the bottom of the $\epsilon_0(R_a)$ well and $Q^a$ is the tunneling phase.
FIG. 3. The vibrational eigenvalues $\epsilon_i^q(R_a)$ of Eq. (2.7) as a function of $R_a$ for all four isotopic variations. The zero of energy is the same as for Fig. 1. The $\epsilon_i^q(R_a)$ that correlate adiabatically to the XY vibrational eigenvalues at large $R_a$, i.e., which correspond to eigenfunctions that are localized in the reactant well at large $R_a$, are shown as full lines; the eigenvalues corresponding to eigenfunctions that are localized in the product well at large $R_a$ are shown as dashed curves. The bold curve marked $\epsilon_0(R_a)$, which correlates with the ground vibrational state of the reactants, is used in the adiabatic model of the lowest energy resonance. The horizontal line denotes the model resonance energy [Eq. (2.12)]. The horizontal dashed lines denote the exact resonance energy ($E^R$ of Table I). The horizontal arrows ending at the right ordinate scale indicate the ground state energies of the isolated hydrogenic molecules. The length in bohr, corresponding to 1 bohr in the abscissa of each of the four panels, has been scaled by the square root of the reduced mass of Eq. (2.3), i.e., in the ratio $1.14:0.97:1.35:1.14$, for the FHD, FH$_2$, FD$_2$, and FDH $R_a$ scales, respectively. With this scaling the $R_a$ motion is associated with a particle of the same mass for all four systems.
In Table I these model results are compared with the energies and the widths of the resonances obtained from the accurate coupled-channel calculations. As seen from the table, an excellent agreement for the resonance energies is obtained for \( F + H_2 \), \( F + HD \), and \( F + D_2 \). The differences between the exact resonance energies and the ones predicted by the model for \( F + H_2 \) and \( F + D_2 \) are actually smaller than the widths of the resonances, and such close agreement may be somewhat fortuitous. The resonance widths obtained from the present model reflect only the adiabatic contribution and may be viewed as an approximate lower bound to the actual resonance width.

The accurate reaction probability vs energy curve for each of the four systems is given in Fig. 5. It can be seen that the curve for the \( F + DH - FD + H \) system does not display a sharp structure analogous to that of the other three curves. However, a recent collision lifetime analysis shows that a very short-lived (0.06 ps) lifetime does occur for this system at a total energy of 0.253 eV and an initial translational energy of 0.020 eV, significantly below the position of the probability peak at 0.044 eV translational energy. The present model does not predict this very weak resonant feature.

IV. DISCUSSION

The resonance positions and widths predicted by the present model agree reasonably well with those found in the accurate coupled-channel results for all but the very weak \( F + DH - FD + H \) resonance. Such agreement indicates that the model reproduces the essential dynamical features of the quasibound state which is responsible for the lowest energy resonances in the \( F + HD \), \( F + H_2 \), and \( F + D_2 \) reactions. As a result, the model can be used to identify the important features of
these systems which lead to the existence, the energy, and the width of these resonances.

The quasibound states responsible for the resonances being considered are formed adiabatically from the separated F + XY reactants. As the F to XY distance decreases, the XY vibration, although somewhat distorted, retains its identity and quantum number, as well as its \( r_a \) direction of motion in the two-dimensional configuration space and becomes a distinct mode of motion of the complex. As a result, the effective potential \( \epsilon_{\nu}(R_a) \), which appears in Eq. (2.12), has a shape which depends largely on the shape of the potential energy function \( V(R_\alpha, r_\alpha) \) along the \( R_\alpha \) direction. In particular, the height of the barrier and the depth of the well in \( \epsilon_{\nu}(R_a) \), which play an important role in determining the characteristics of the resonance, correlate well with the features of \( V(R_\alpha, r_\alpha) \) along \( R_\alpha \) in the vicinity and to the left of the saddle point region. Very approximately, the values of \( \epsilon_{\nu}(R_a) \) are shifted upwards from those of \( V(R_\alpha, r_\alpha) \) in the neighborhood of the minimum energy path, by the zero-point energy of the isolated XY hydrogenic molecule. This overall trend suggests that, in going from one isotopic system to another, the shift in the resonance energy should correlate approximately with the shift in the corresponding XY zero-point level. This is indeed the case, for both the model and the accurate results, as can be seen from a comparison of the second column of Table I with the next three. Thus, the qualitative nature of the variation of the position of the resonance with the isotopic system being considered is easily explained. A more quantitative explanation, encompassed in the model, includes the effects on the \( \epsilon_{\nu}(R_a) \) vs \( R_\alpha \) curves of the changes produced in the \( r_a \) motion zero-point energy by the approaching F atom.

The trend in the widths of the resonances, as we compare the several isotopic systems, is of a more subtle nature. Couplings of the entrance channel quasibound complex to both reagent and product states are involved, since both these couplings lead to its decay. The relatively narrow widths of those resonances for the FHID, FHI, and FD systems, as obtained from the coupled-channel calculations, indicates that the couplings of the multiplicative wave function described by Eq. (2.9) to other states is relatively weak. This weakness is, to some extent, due to the significant separation of the time scales of the motion along \( R_\alpha \) and \( r_\alpha \) in the complex, which separation in turn depends largely on the ratio of the effective masses for the motions along \( R_\alpha \) and \( r_\alpha \). This mass ratio \( M_\alpha/m_\alpha \) is about 4 for all four isotopic systems. Another factor is that the quasibound complex is in the ground vibrational state and its \( r_\alpha \) motion is mainly confined to a strip of the potential energy surface along the \( R_\alpha \) path. In a considerable portion of the region of the surface in which the complex is localized, the potential energy function is separable within that strip, leading to the validity of Eq. (2.9) and consequently, to the weakness of the couplings of that state to other states which may lead to its decay.

Let us now analyze the nature of the couplings of the quasibound state to the reactant and product states. In the present systems, the decay of the resonance state into reactants occurs adiabatically into the ground vibrational state of the latter, and is due to tunneling through or motion over the effective dynamic potential barrier of the \( \epsilon_{\nu}(R_a) \) curve of Fig. 3 along the outward \( R_\alpha \) direction. This \( R_\alpha \) mode of motion of the complex transforms thereby into reactant translation, while the \( r_\alpha \) mode remains largely unchanged and becomes the reactant vibration. The part of the resonance width that is due to this decay mechanism is controlled by the barrier traversal amplitude, i.e., by the likelihood of the system either tunneling through the barrier or moving over it, and is determined from the dependence of the phase shift resulting from Eq. (2.12) on energy for the F + H and F + D systems or from the tunneling coefficient at the exact resonance energy for the F + HD system. It is the only part of the width that can be readily estimated from the present model.

The complex can, in addition, decay into (or be formed from) the scattering states of the products. This decay pathway requires a more severe restructuring of the complex and involves vibrationally nonadiabatic processes. The enhancement of all of the state-to-state reactive transition probabilities in the reso-
nance energy region, seen in Fig. 5 and in the other exact reaction probability vs energy curves of Refs. 18 and 29, indicates that all the energetically accessible product states interact with the complex, although the present model furnishes an estimate of the strength of the interaction of the complex with the reactants, it does not lead to an estimation of its coupling with the products. The reason for this is that the coordinates that are appropriate for describing product states are different from those that are appropriate for describing the complex. The strength of the nonadiabatic coupling between two states is usually, to some extent at least, inversely proportional to the energy gap between them. One can therefore expect the energetically closest product state, in this case the highest accessible one, to interact the most strongly with the complex. An examination of the transition probabilities to different vibrational states of the product (Fig. 5 and Ref. 18) does indeed indicate the existence of such a trend.

The partial width of the resonance due to the coupling of the complex to the product states should roughly correspond to the difference between the exact width and the adiabatic partial width due to coupling with the reactants, determined in Sec. III, and presented in Table I. This difference, obtained from the last two columns of that table, suggests that the coupling to the product states is of the same order of magnitude as the coupling to the reactant states.

With these considerations in mind, let us attempt to interpret the trend in the resonance widths calculated for the FHD, FH₂, and FD₂ series by the model and by the coupled-channel method and displayed in the last two columns of Table I. A noticeable feature in that series is the broadening of the resonances in the order given. The potential energy surface for all three systems is the same function of the internuclear coordinates. However, when expressed in the (unscaled) Jacobi Cartesian coordinates Rₐ, Rₐ', some differences appear. The surfaces for FH₂ and FD₂ continue to be equal to each other in these coordinates, but that for FHD is slightly different, due to the shift of the center of mass of HD away from its midpoint and to the change of the (unscaled coordinate) skew angle from tan⁻¹ 63. 4° to tan⁻¹ 3 = 71. 6°. In the following qualitative discussion, this small difference will be disregarded. In addition, although the mass mₐ changes from ½ to 3 atomic hydrogen mass units in going from F + HD to F + HD₂, the effect of this change on the ε₂(R₂ₐ) (rather than Rₐ) curve is also small. The reason for this smallness is that ε₂(R₂ₐ) depends largely on the characteristics of V(Rₐ; R₂ₐ) in the general vicinity of its saddle point, and only to a minor extent on mₐ, as is also the case for ε₂(R₂ₐ). As a result, ε₂(R₂ₐ) is approximately the same for both these two systems. The most important difference between them is due to Mₐ, which is 1. 81 and 2. 59 atomic hydrogen mass units for F + HD and F + HD₂, respectively. The effect of this mass change is to lower the energy of the quasibound state in Eq. (2.12) [with respect to the bottom of the ε₂(R₂ₐ) well] by approximately the square root of the ratio of these masses. An equivalent way of arriving at the same conclusion is to replace the variable Rₐ in that equation by the variable ρ₂ₐ = μ₁/²Rₐ = Mₐ₁/²R₂ₐ (as done in Fig. 3). This makes the well in the ε₂(ρ₂ₐ) vs ρ₂ₐ curve for F + HD be somewhat broader than for the F + HD₂ one, but the new effective mass in the transformed equation is now the same for both systems, resulting in the lowering of the quasibound level. This lowering results in the increase of the thickness of the barrier through which tunneling must occur for the system to move from this well region to the reactant region. Since the lifetime associated with the tunneling increases exponentially with this thickness, the resonance lifetime for FH₁₂ is higher than that of FH₂. In going from FHD to FD₂, the ε₂(ρ₂ₐ) well width increases further, which would tend to make the lifetime of the FD₂ resonance longer than that of the FHD one. However, as can be seen from Fig. 3(c), an avoided crossing between the ε₂(ρ₂ₐ) curve and another one [which localizes the corresponding eigenfunction φ₂(r₂ₐ; R₂ₐ) in the product region] reduces the depth of the well in this curve and pushes the corresponding quasibound state slightly higher than the top of the barrier. This effect is dominant and results in an FD₂ resonance broader than those for the FHD and FH₂ systems. It should be noted that this analysis refers to the adiabatic part of the resonance width. The nonadiabatic part associated with the interaction of the complex with the products should further depend on the Delves coordinate skew angle, and should increase as this angle increases, in the direction FHD, FH₂, FD₂ as is observed in the accurate calculations.

Another consequence of the analysis above is that the resonance characteristics are very sensitive to the shape and size of the barrier in the effective potential ε₂(R₂ₐ) vs R₂ₐ curve and therefore, on the shape and size of the barrier in the V(Rₐ; R₂ₐ) surface along the Rₐ direction. As a result, small changes in the details of the saddle point region of the surface (smaller in magnitude than 1 kcal/mole) can be expected to introduce radical changes in the resonance characteristics, including the possibility of their complete disappearance. Consequently, such resonances will constitute a very sensitive probe for the experimental characterization of that region of the FHD potential energy surface.

As observed in Fig. 5, the off-resonance transition probabilities for the F + H₂ and F + D₂ reactions are much larger than for the F + HD one. This behavior can be understood by considering that the pronounced narrowness of the FHD resonance is a manifestation of the weakness of the coupling of the reactant states to the product states through the strong interaction (complex) region, with a resulting small reaction probability away from resonance when compared with the other two systems.

The present model for the resonance in the F + H₂, F + HD, and F + D₂ reactions is similar to the one used by Chapman and Hayes, 3,41 to study the resonances in H + H₂ reactive collisions and in H⁺ + H₂ inelastic collisions. In their model, a zero-order approximation for the wave function of the complex is obtained in the form ψ = G(R₂ₐ; ρ₂ₐ'), where G(ρ₂ₐ') is the isolated reagent diatomic molecule wave function and G(R₂ₐ) is the eigenfunction associated with V(R₂ₐ; ρ₂ₐ'), i.e., calculated.
along the potential energy surface with $r_d^*$ fixed at its isolated reagent equilibrium position. The distortion of the diatomic molecule during the collision is accounted for by a first-order perturbation correction to the asymptotic eigenvalues. The nature of the resonances in the present systems is, however, markedly different from the ones in He + H$_2$ and I + H$_2$. The latter are supported by actual two-dimensional potential energy wells in the corresponding surfaces. No such wells exist in the present case, and the origin of the stability of the complex is of a purely dynamical nature.

The quantum dynamics of the resonances in one of the reactions studied here, namely F + H$_2$, has been a subject of a detailed study by Latham et al. by means of a vibrationally adiabatic analysis, a vibrational entropy analysis, and a scattering wave function probability density analysis. Their vibrationally adiabatic analysis in a system of curved reaction coordinates shows that no single channel model in those coordinates can account for the resonances, whereas in our model it does. The sharp increase in the scattering wave function probability density in a small region of the surface at the resonance energy indicates clearly the location on the surface where the complex is formed. The location of the complex predicted by the present model appears to be in reasonable agreement with that indicated by the scattering probability density maps in Ref. 20.

Another study of the resonances on the F + H$_2$ reaction by Hayes and Walker came to our attention during the preparation of this manuscript. One of the aspects of their work is the study of the sensitivity of the resonances to the shape of the potential in the entrance channel. They have shown that if the entrance channel barrier is removed from the surface, the scattering results on the new surface do not show a resonance. The disappearance of the resonance as a result of this removal is in agreement with the conclusions drawn above from the validity of the present model.

Although the present study was done using only one semiempirical surface, most of the collinear F + H$_2$ surfaces used so far have similar basic features, and we hope that our model will also be useful for predicting the absence or presence and the characteristics of this type of resonance on other more accurate surfaces for the F + H$_2$ reaction.

Resonances of a similar nature can be expected to be seen in other exothermic reactions with a small barrier early in the entrance channel. The present type of analysis can also be expected to be useful in modeling resonances in low barrier asymmetric reactions of the heavy–light–heavy type. On the other hand, this approach will be less useful for reactions with light–heavy–light (LHL) mass combinations since the reagent vibrational motion is in such cases almost parallel to the product translational motion (in configuration space) making it unlikely that a complex of the present type would exist. In the limit of infinite central atom mass, even Eq. (2.11) becomes invalid. The resonances in thermonuclear or nearly thermonuclear reactions of the LHL type usually have a different physical origin and can be successfully modeled using different techniques.

Removing the collinear restriction from the motion of the triatomic complex would introduce changes in its dynamics which are, as shown previously for the case of the H + H$_2$ and F + H$_2$ systems, to a certain extent, predictable. The equilibrium configuration of the complex would, based on the structure of the surface, remain collinear. There would be two additional degrees of freedom of the complex, described either as a doubly degenerate hindered rotation or a doubly degenerate bending vibration. For the resonances studied here, they are likely to be most separable if treated as hindered rotations of the H$_2$ molecule around its center of mass. The existence of these two additional modes of motion would shift the energy of the complex for collisions with zero total angular momentum upwards by the sum of their zero-point energies.

The most serious complication introduced by removing the collinear restriction for the F + H$_2$ reaction is the large number of possible rotational states of the complex that spread over a wide energy range. As a result, the long-lived complex can be formed, as shown by approximate coupled-channel calculations, over a wide range of collisional energies and does not produce strong structure in the integral reaction cross section vs energy curve. However, angular distributions of reaction products are expected to be quite different at resonance energies as compared with the nonresonance ones. As a result, it is important to develop models that predict the shape of the differential reaction cross section vs scattering angle curves at resonance. Such models could permit the establishment of relations between the characteristics of the shape of the potential energy surface in the strong interaction region and characteristics of those angular distributions. These relationships would be very useful in extracting information about the details of potential energy surfaces from experimental observations of reactive scattering resonances.

V. SUMMARY

The results and conclusions given in the present paper can be summarized as follows:

(1) The lowest resonance in the collinear reactions $F + H_2(v = 0) \rightarrow FH + H$, $F + HD(v = 0) \rightarrow FH + D$, and $F + DH(v = 0) \rightarrow FD + D$ is due to the formation of a quasi-bound complex which is localized in the entrance channel region of the corresponding potential energy surface, slightly to the left of its saddle point.

(2) The ordering of the positions of these resonances is the same as that of the zero-point energies of the reactant molecules. The reason for this interrelation is that the shape of the effective potential in the $R_a$ direction, which correlates adiabatically with the ground state of the reactant, is determined largely by the shape of the potential energy surface $V(R_a, r_d)$ along $R_a$.

(3) The nonadiabatic coupling of the complex to the reaction products is of the same order of magnitude as its adiabatic coupling to the reagents.

(4) The adiabatic part of the resonance width increases in the order FHD, FH$_2$, FD$_2$. The increase from
FHD to FH₂ is due to a corresponding lowering of the effective adiabatic barrier height. The increase from FHD to FD₄ is due to a decrease of the depth of the adiabatic well.

(5) The position and width of the resonance is a very sensitive function of the shape of $V(r_1, r_2)$ in the saddle point region and may change or the resonances may even disappear if that region of the potential is modified.

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49. There are small differences in the resonance energies found in different coupled-channel calculations on the Mukeraman V potential energy surface in the literature due to slight variations in the equilibrium parameters and physical constants used as well as to the use of different analytic fits to the surface.
49. However, when the resonance energy found by the model is significantly higher than the exact one (which is not the present case), the width of the model resonance can be larger than that of the exact results due to a possible overestimate of the likelihood of adiabatic decay.