

Analytical solutions of the Bohr Hamiltonian with the Morse potential

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Analytical solutions of the Bohr Hamiltonian are obtained in the γ -unstable case, as well as in an exactly separable rotational case with $\gamma \approx 0$, called the exactly separable Morse (ES-M) solution. Closed expressions for the energy eigenvalues are obtained through the asymptotic iteration method (AIM), the effectiveness of which is demonstrated by solving the relevant Bohr equations for the Davidson and Kratzer potentials. All medium mass and heavy nuclei with known β_1 and γ_1 bandheads have been fitted by using the two-parameter γ -unstable solution for transitional nuclei and the three-parameter ES-M for rotational ones. It is shown that bandheads and energy spacings within the bands are well reproduced for more than 50 nuclei in each case.

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I. INTRODUCTION

The recent introduction of the critical point symmetries E(5) [1] and X(5) [2], which describe shape phase transitions between vibrational and γ -unstable/prolate deformed rotational nuclei, respectively, has stirred much interest in special solutions of the Bohr Hamiltonian [3], describing collective nuclear properties in terms of the collective variables β and γ . Such solutions can describe nuclei in the whole region between different limiting symmetries, while critical point symmetries are appropriate for describing nuclei only at or near the critical point, in good agreement with experiment [4–8].

Shape phase transitions in nuclear structure have been first discovered [9] in the classical analog [10,11] of the interacting boson model [12], which describes collective nuclei in terms of collective bosons of angular momentum zero (s bosons) and two (d bosons) in the framework of a U(6) overall symmetry, possessing U(5) (vibrational), SU(3) (prolate deformed rotational), and O(6) (γ -unstable) limiting symmetries. To visualize these limiting symmetries and the transitions between them, it is useful to place them at the corners of a symmetry triangle [13]. A similar triangle for the collective model has been introduced [14]. In the IBM framework, it has been found that a first-order phase transition occurs between U(5) and SU(3), while a second-order phase transition occurs between U(5) and O(6) [9]. Within the collective model, X(5) corresponds to the first case and E(5) to the second.

It has been known for a long time [15] that simple special solutions of the Bohr Hamiltonian, resulting from exact separation of variables in the relevant Schrödinger equation, can be obtained in the γ -unstable case, in which the potential depends only on β , as well as in the case in which the potential can be written in the separable form

$$u(\beta, \gamma) = u(\beta) + \frac{u(\gamma)}{\beta^2}, \quad (1)$$

in the special cases of $\gamma \approx 0$ or $\gamma \approx \pi/6$ [16]. An approximate separation of variables has also been attempted for potentials of the form

$$u(\beta, \gamma) = u(\beta) + u(\gamma), \quad (2)$$

in the cases of $\gamma \approx 0$ [2] or $\gamma \approx \pi/6$ [17]. A brief summary of existing solutions is listed here.

- (i) The E(5) critical point symmetry [1] is a γ -unstable solution, using as $u(\beta)$ an infinite-well potential starting from $\beta = 0$. Displacing the well from $\beta = 0$ leads to the O(5)-confined beta soft [O(5)-CBS] model [18]. γ -unstable solutions have also been given for the Coulomb [19] and Kratzer [19] potentials, β^{2n} potentials ($n = 1, 2, 3, 4$) [20,21] (labeled as E(5)- β^{2n}), as well as for the Davidson potential [22–25]

$$u(\beta) = \beta^2 + \frac{\beta_0^4}{\beta^2}, \quad (3)$$

where β_0 is the position of the minimum of the potential. A solution using a well of finite depth has also been given [26].

- (ii) The X(5) critical point symmetry [2] is an approximate solution, using a potential of the form of Eq. (2) with $\gamma \approx 0$ [achieved by using as $u(\gamma)$ a harmonic oscillator potential with minimum at $\gamma = 0$] and an infinite-well potential starting from $\beta = 0$ as $u(\beta)$. Displacing the well from $\beta = 0$ leads to the confined beta soft (CBS) model [27]. Similar solutions have been obtained for β^{2n} potentials ($n = 1, 2, 3, 4$) [labeled as X(5)- β^{2n}] [28], as well as for the Davidson potential [25].
- (iii) Z(5) [17] is an approximate solution, using a potential of the form of Eq. (2) with $\gamma \approx \pi/6$.
- (iv) Exactly separable (ES) solutions using a potential of the form of Eq. (1) and $\gamma \approx 0$ have been obtained for the Coulomb [29] and Kratzer [29] potentials, the infinite-well potential [labeled as ES-X(5)] [30], the harmonic oscillator potential (labeled as ES-X(5)- β^2) [30], as well as for the Davidson potential [31] (labeled as ES-D).
- (v) Exactly separable solutions using a potential of the form of Eq. (1) and $\gamma \approx \pi/6$ have been obtained for the Coulomb [32,33], Kratzer [32,33], and Davidson [32,33] potentials.
- (vi) In addition to the special analytical solutions mentioned above, a powerful method for solving the Bohr Hamiltonian numerically has been developed recently

[34], evolving into an algebraic collective model [35]. The relations between the algebraic collective model and the different limiting symmetries of the interacting boson model [12] have been studied in Refs. [36,37]. Using this numerical method, the Bohr Hamiltonian has been solved [38] for the same potentials used in X(5), but avoiding the approximate separation of variables, resulting in evidence for strong β - γ mixing.

Chains of models mentioned above cover regions between different limiting symmetries. For example, the chain E(5)- β^{2n} ($n = 1, 2, 3, 4$), E(5), O(5)-CBS spans the region between the vibrational [U(5)] and γ -unstable [O(6)] limits, while the chain X(5)- β^{2n} , X(5), CBS spans the region between the vibrational and prolate deformed rotational [SU(3)] limits. Their predictions can therefore be tested against a large body of experimental data [39].

The potentials mentioned above (infinite well, harmonic oscillator, Coulomb, Kratzer, and Davidson) are known to be exactly soluble for all values of angular momentum L . In the present work, we introduce special solutions for the Morse potential [40],

$$u(\beta) = e^{-2a(\beta-\beta_e)} - 2e^{-a(\beta-\beta_e)}, \quad (4)$$

which is known [41,42] to be exactly soluble only for $L = 0$. The overall factor D of the Morse potential is set equal to unity, without affecting the method of solution, since it can be scaled out if ratios of energies are used, as in the present work. Analytical expressions for the spectra for any L are obtained by solving the relevant differential equation through the asymptotic iteration method (AIM) [43,44], after applying the Pekeris approximation [45]. Solutions for the γ -unstable case and the exactly separable rotational case with $\gamma \approx 0$ (to be called ES-M) are obtained. To demonstrate the effectiveness of AIM, we first apply it to the Davidson and Kratzer potentials in the same cases (γ -unstable, exactly separable rotational with $\gamma \approx 0$), recovering the above-mentioned solutions which can be obtained in terms of special functions.

A few advantages of the present approach are listed here.

- (i) In X(5) and related models, using potentials of the form of Eq. (2), the ground state and β bands depend only on the parameters of the β potential, while the γ bands depend also on an additional parameter introduced by the γ potential [usually the stiffness of the harmonic oscillator used as $u(\gamma)$]. When exactly separable potentials of the form of Eq. (1) are used, all bands (ground state, β , γ) depend on all parameters. Thus, all bands are treated on an equal footing, as in the case of the ES-D solution [31].
- (ii) A well-known problem of X(5) and related solutions is the overprediction of the energy spacings within the β band by almost a factor of 2 [6–8]. It is known that this problem can be avoided by replacing the infinite-well potential of X(5) by a potential with sloped walls [46]. The present solution avoids this problem, since the right branch of the Morse potential imitates the sloped wall.

To test the applicability of the Morse potential in the description of nuclear spectra, we have fitted all nuclei with mass $A \geq 100$ and $R_{4/2} = E(4)/E(2) < 2.6$ for which at least

the β_1 and γ_1 bandheads are known [47], using the γ -unstable solution of the Morse potential, which involves two free parameters (β_e, a). We have also fitted all nuclei with mass $A \geq 150$ and $R_{4/2} = E(4)/E(2) > 2.9$ for which at least the β_1 and γ_1 bandheads are known [47], using the exactly separable rotational solution of the Morse potential with $\gamma \approx 0$ (ES-M), which involves three free parameters (the Morse parameters β_e and a , as well as the stiffness c of the γ potential, for which a harmonic oscillator is used). A comparison of the latter to the fits provided by the Davidson potential in the exactly separable $\gamma \approx 0$ case [31] (ES-D), which contains two free parameters (β_0, c) instead of three, shows that the extra parameter extends the region of applicability of the model in the same nuclei to higher angular momenta, largely improving the quality of the fits.

In Sec. II of the present work, the asymptotic iteration method (AIM) is briefly reviewed. The method is then applied to the exactly separable rotational $\gamma \approx 0$ case for the Davidson, Kratzer, and Morse potentials in Sec. III, and to the γ -unstable case of the same potentials in Sec. IV, with the details of the calculations given in Appendixes A–F. Fits to experimental data are presented in Sec. V, while Sec. VI discusses the present results and plans for further work.

II. OVERVIEW OF THE ASYMPTOTIC ITERATION METHOD

The asymptotic iteration method (AIM) has been proposed [43,44] and applied [48–53] to the solution of second-order differential equations of the form

$$y'' = \lambda_0(x)y' + s_0(x)y, \quad (5)$$

where $\lambda_0(x) \neq 0$ and the prime denotes the derivative with respect to x . The functions $s_0(x)$ and $\lambda_0(x)$ must be sufficiently differentiable. Equation (5) has the general solution [43]

$$y(x) = \exp\left(-\int^x \alpha(x_1)dx_1\right) \left[C_2 + C_1 \int^x \exp\left(\int^{x_1} [\lambda_0(x_2) + 2\alpha(x_2)]dx_2\right) dx_1 \right] \quad (6)$$

for sufficiently large $k, k > 0$, if

$$\frac{s_k(x)}{\lambda_k(x)} = \frac{s_{k-1}(x)}{\lambda_{k-1}(x)} = \alpha(x), \quad (7)$$

where

$$\lambda_k(x) = \lambda'_{k-1}(x) + s_{k-1}(x) + \lambda_0(x)\lambda_{k-1}(x), \quad (8)$$

$$s_k(x) = s'_{k-1}(x) + s_0(x)\lambda_{k-1}(x), \quad k = 1, 2, 3, \dots$$

For a given potential, the radial Schrödinger equation is converted to the form of Eq. (5). Then, $s_0(x)$ and $\lambda_0(x)$ are determined, and the functions $s_k(x)$ and $\lambda_k(x)$ are calculated by the recurrence relations of Eq. (8).

The termination condition of the method, given in Eq. (7), can be arranged as

$$\Delta_k(x) = \lambda_k(x)s_{k-1}(x) - \lambda_{k-1}(x)s_k(x) = 0, \quad (9)$$

$$k = 1, 2, 3, \dots$$

Then, the energy eigenvalues are obtained from the roots of Eq. (9) if the problem is exactly solvable. If not, for a specific principal quantum number n , we choose a suitable x_0 point, generally determined as the maximum value of the asymptotic wave function or the minimum value of the potential [43,50], and the approximate energy eigenvalues are obtained from the roots of this equation for sufficiently large values of k by iteration.

The corresponding eigenfunctions can be derived from the following wave function generator for exactly solvable potentials:

$$y_n(x) = C_2 \exp\left(-\int^x \frac{s_n(x_1)}{\lambda_n(x_1)} dx_1\right), \quad (10)$$

where n represents the radial quantum number.

Recently, Boztosun and Karakoc [54] further improved the method for the exactly solvable problems by rewriting the second-order differential equation of Eq. (5) in the form

$$y'' = -\frac{\tau(x)}{\sigma(x)} y' - \frac{\Omega_n}{\sigma(x)} y. \quad (11)$$

By comparison with Eq. (5), $\tau(x)$, $\sigma(x)$, and Ω_n can be found to be

$$-\frac{\tau(x)}{\sigma(x)} = \lambda_0(x), \quad -\frac{\Omega_n}{\sigma(x)} = s_0(x), \quad (12)$$

where Ω_n is a constant which comprises the eigenvalue. Then, the energy eigenvalues are obtained from

$$\Omega_n = -n\sigma'(x) - \frac{n(n-1)}{2}\sigma''(x). \quad (13)$$

The applicability of this new solution is demonstrated for the Morse potential in Appendixes C and F.

III. EXACTLY SEPARABLE SOLUTIONS FOR $\gamma \approx 0$

The original collective Bohr Hamiltonian [3] is

$$H = -\frac{\hbar^2}{2B} \left[\frac{1}{\beta^4} \frac{\partial}{\partial \beta} \beta^4 \frac{\partial}{\partial \beta} + \frac{1}{\beta^2 \sin 3\gamma} \frac{\partial}{\partial \gamma} \sin 3\gamma \frac{\partial}{\partial \gamma} - \frac{1}{4\beta^2} \sum_{k=1,2,3} \frac{Q_k^2}{\sin^2(\gamma - \frac{2\pi k}{3})} \right] + V(\beta, \gamma), \quad (14)$$

where β and γ are the usual collective coordinates which define the shape of the nuclear surface. Q_k ($k = 1, 2, 3$) represents the angular momentum components in the intrinsic frame, and B is the mass parameter. Reduced energies and reduced potentials are defined as $\epsilon = 2BE/\hbar^2$, $v = 2BV/\hbar^2$, respectively [1]. If the potential has a minimum around $\gamma = 0$, the angular momentum term in Eq. (14) can be written [2] as

$$\sum_{k=1}^3 \frac{Q_k^2}{\sin^2(\gamma - \frac{2\pi k}{3})} \approx \frac{4}{3} (Q_1^2 + Q_2^2 + Q_3^2) + Q_3^2 \left(\frac{1}{\sin^2 \gamma} - \frac{4}{3} \right). \quad (15)$$

Exact separation of variables can be achieved [15,31–33] for potentials of the form $u(\beta, \gamma) = u(\beta) + u(\gamma)/\beta^2$, given in

Eq. (1). We then assume wave functions of the form

$$\psi(\beta, \gamma, \theta_j) = \xi_L(\beta) \Gamma_K(\gamma) \mathcal{D}_{M,K}^L(\theta_j), \quad (16)$$

where θ_j ($j = 1, 2, 3$) are the Euler angles, $\mathcal{D}(\theta_j)$ represents Wigner functions of these angles, L stands for the eigenvalues of the angular momentum, while M and K are the eigenvalues of the projections of the angular momentum on the laboratory-fixed z axis and the body-fixed z' axis, respectively. The Schrödinger equation is thus separated, as in Refs. [15,31–33], into a “radial” part (depending on β) and a γ part.

A. Davidson potential

Solving through AIM, the β equation for the Davidson potential [22] of Eq. (3), $u(\beta) = \beta^2 + \beta_0^4/\beta^2$, where β_0 is the position of the minimum of the potential, we get the energy eigenvalues

$$\epsilon_{n,L} = 2n + 1 + \left[\frac{9}{4} + \frac{L(L+1)}{3} + \lambda + \beta_0^4 \right]^{1/2}, \quad (17)$$

where λ is a term coming from the exact separation of variables, determined from the γ equation.

The γ equation has been solved [2,31] for a potential

$$u(\gamma) = (3c)^2 \gamma^2, \quad (18)$$

leading to

$$\lambda = \epsilon_\gamma - \frac{K^2}{3}, \quad \epsilon_\gamma = (3C)(n_\gamma + 1), \quad (19)$$

where $C = 2c$. The final result for the energy eigenvalues coincides with the results of Ref. [31]. The details of the AIM calculation are given in Appendix A.

Special case 1: If we take $\beta_0 = 0$, the Davidson potential becomes the potential of a harmonic oscillator. Then, the expression for the energy eigenvalues coincides with the one obtained for the ES-X(5)- β^2 model of Ref. [30].

Special case 2: Taking $c = 0$ and $K = 0$ (i.e. $\lambda = 0$) in Eq. (17), we obtain for the ground state and β bands energy eigenvalues of the X(5)-Davidson solution of Ref. [25].

B. Kratzer potential

We again use an exactly separable potential of the form given in Eq. (1), with a Kratzer potential [19,29]

$$u(\beta) = -\frac{A}{\beta} + \frac{B}{\beta^2}, \quad A > 0, \quad (20)$$

in β ; while in γ , we use a harmonic oscillator potential, $u(\gamma) = c\gamma^2/2$, as in Ref. [29]. Solving the “radial” (β) equation by AIM, we obtain the energy eigenvalues

$$\epsilon_{n,L} = \frac{A^2/4}{\left(n + \frac{1}{2} + \sqrt{\frac{9}{4} + \lambda + B + \frac{L(L+1)}{3}}\right)^2}, \quad (21)$$

which, in comparison with the result of Ref. [29], contain the additional term λ , coming from the procedure of exact separation of variables, as shown in detail in Appendix B. The term λ is determined by solving the γ equation through

AIM, as exhibited in detail in Appendix B, the final result being

$$\lambda = (2c)^{1/2} (n_\gamma + 1) - K^2/3, \quad (22)$$

in agreement with the solution of the γ equation given in Ref. [29], as demonstrated in Appendix B.

Special case: Taking $B = 0$ in Eq. (20), the Coulomb potential is obtained. The energy eigenvalues are still given by Eq. (21) with $B = 0$. Again, the result differs from the one reported in Ref. [29] by the presence of the λ term, due to the exact separation of variables, as already remarked.

C. Morse potential

We use the exactly separable form of the potential given in Eq. (1) again. The Morse potential [40] is defined as

$$u(\beta) = e^{-2a(\beta-\beta_e)} - 2e^{-a(\beta-\beta_e)}. \quad (23)$$

Using the Pekeris approximation [45] and solving the β equation through AIM (the details are given in Appendix C), we obtain the energy eigenvalues

$$\epsilon_{n,L} = \frac{\mu c_0}{\beta_e^2} - \left[\frac{\gamma_1^2}{2\beta_e \gamma_2} - \left(n + \frac{1}{2} \right) \frac{\alpha}{\beta_e} \right]^2, \quad (24)$$

where

$$c_0 = 1 - \frac{3}{\alpha} + \frac{3}{\alpha^2}, \quad c_1 = \frac{4}{\alpha} - \frac{6}{\alpha^2}, \quad (25)$$

$$c_2 = -\frac{1}{\alpha} + \frac{3}{\alpha^2}, \quad \alpha = a\beta_e, \quad (26)$$

$$\gamma_1^2 = 2\beta_e^2 - \mu c_1, \quad \gamma_2^2 = \beta_e^2 + \mu c_2, \quad (27)$$

$$\mu = \frac{L(L+1)}{3} + 2 + \lambda. \quad (27)$$

λ in the last equation comes from the exact separation of variables and is determined from the γ equation. We use the same γ potential $u(\gamma) = (3c)^2 \gamma^2$ [Eq. (18)] as in the Davidson case (Sec. III A), leading to

$$\lambda = \epsilon_\gamma - \frac{K^2}{3}, \quad \epsilon_\gamma = (3C)(n_\gamma + 1), \quad (28)$$

where $C = 2c$, as given in Eq. (19).

IV. γ -UNSTABLE SOLUTIONS

In this case, the reduced potential is assumed to be γ independent, $v(\beta, \gamma) = u(\beta)$. Then the wave function is assumed to be of the form [15]

$$\psi(\beta, \gamma, \theta_j) = R(\beta)\Phi(\gamma, \theta_j). \quad (29)$$

The equation which includes the Euler angles and γ has been solved by Bès [55]. In this equation, the eigenvalues of the second-order Casimir operator of SO(5) occur, having the form $\Lambda = \tau(\tau + 3)$, where τ is the seniority quantum number, characterizing the irreducible representations of SO(5) and taking the values $\tau = 0, 1, 2, 3, \dots$ [56].

The values of the angular momentum L are given by the algorithm

$$\tau = 3\nu_\Delta + \lambda, \quad \nu_\Delta = 0, 1, 2, \dots, \quad (30)$$

$$L = \lambda, \lambda + 1, \dots, \quad 2\lambda - 2, 2\lambda \quad (31)$$

(with $2\lambda - 1$ missing), where ν_Δ is the missing quantum number in the reduction $SO(5) \supset SO(3)$ [56]. The ground state band levels are determined by $L = 2\tau$ and $n = 0$.

A. Davidson potential

The radial equation, when solved through the AIM, leads to the energy eigenvalues

$$\epsilon_{n,\tau} = 2n + 1 + \left[\frac{9}{4} + \tau(\tau + 3) + \beta_0^4 \right]^{1/2}, \quad (32)$$

the details of the calculation are given in Appendix D. This result coincides with those of Ref. [25].

Special case: Taking $\beta_0 = 0$ in Eq. (32), one gets the simplified expression

$$\epsilon_{n,\tau} = 2n + \tau + \frac{5}{2}, \quad (33)$$

which is the five-dimensional harmonic oscillator solution of Bohr [3].

B. Kratzer potential

Solving the radial equation by AIM (see Appendix E for details), we obtain the energy eigenvalues

$$\epsilon_{n,\tau} = \frac{A^2/4}{\left(n + \frac{1}{2} + \sqrt{\frac{9}{4} + \tau(\tau + 3) + B} \right)^2}, \quad (34)$$

which coincide with the results of Ref. [19].

Special case: For $B = 0$, the γ -unstable solution for the Coulomb potential is obtained, i.e.,

$$\epsilon_{n,\tau} = \frac{A^2/4}{(n + \tau + 2)^2}, \quad (35)$$

which coincides with the result of Ref. [19].

C. Morse potential

Using the Pekeris approximation [45] and AIM (see Appendix F for the details), we obtain the energy eigenvalues

$$\epsilon_{n,\tau} = \frac{\nu c_0}{\beta_e^2} - \left[\frac{\gamma_1^2}{2\beta_e \gamma_2} - \left(n + \frac{1}{2} \right) \frac{\alpha}{\beta_e} \right]^2, \quad (36)$$

where

$$\gamma_1^2 = 2\beta_e^2 - \nu c_1, \quad \gamma_2^2 = \beta_e^2 + \nu c_2, \quad (37)$$

$$\nu = \tau(\tau + 3) + 2, \quad (38)$$

with the rest of the quantities given again by Eq. (25).

V. NUMERICAL RESULTS

To test the applicability of the Morse potential in the description of nuclear spectra, we have fitted all nuclei with mass $A \geq 100$ and $R_{4/2} = E(4)/E(2) < 2.6$, for which at least the β_1 and γ_1 bandheads are known, using the γ -unstable solution of the Morse potential, which involves two free parameters (β_e, a), as described in Sec. IV C. Results for 54 nuclei are shown in Table I. The quality measure

$$\sigma = \sqrt{\frac{\sum_{i=1}^n (E_i(\text{exp}) - E_i(\text{th}))^2}{(n-1)E(2_1^+)}} \quad (39)$$

used in the rms fits, remains below 1 in most cases.

TABLE I. Comparison of theoretical predictions of the γ -unstable Bohr Hamiltonian with Morse potential (described in Sec. IV C) and experimental data [47] of nuclei with $A \geq 100$, $R_{4/2} \leq 2.6$, and known 0_2^+ and 2_2^+ states. The $R_{4/2} = E(4_1^+)/E(2_1^+)$ ratios, as well as the β and γ bandheads normalized to the 2_1^+ state and labeled as $R_{0/2} = E(0_2^+)/E(2_1^+)$ and $R_{2/2} = E(2_2^+)/E(2_1^+)$, respectively, are shown. The angular momenta of the highest levels of the ground state, β , and γ bands included in the rms fit are labeled as L_g , L_β , and L_γ , respectively; n indicates the total number of levels involved in the fit and σ is the quality measure of Eq. (39). See Sec. V for further discussion.

Nucleus	$R_{4/2}$ exp	$R_{4/2}$ th	$R_{0/2}$ exp	$R_{0/2}$ th	$R_{2/2}$ exp	$R_{2/2}$ th	β_0	a	L_g	L_β	L_γ	n	σ
⁹⁸ Ru	2.14	2.25	2.0	2.4	2.2	2.3	3.84	0.44	18	0	4	12	0.659
¹⁰⁰ Ru	2.27	2.29	2.1	2.8	2.5	2.3	4.43	0.36	28	0	4	17	0.291
¹⁰² Ru	2.33	2.25	2.0	2.3	2.3	2.2	3.78	0.42	16	0	5	12	0.341
¹⁰⁴ Ru	2.48	2.33	2.8	2.9	2.5	2.3	7.57	0.10	8	2	8	12	0.433
¹⁰² Pd	2.29	2.28	2.9	2.6	2.8	2.3	4.30	0.34	26	4	4	18	0.219
¹⁰⁴ Pd	2.38	2.28	2.4	2.6	2.4	2.3	4.15	0.41	18	2	4	13	0.300
¹⁰⁶ Pd	2.40	2.26	2.2	2.4	2.2	2.3	3.93	0.43	16	4	5	14	0.343
¹⁰⁸ Pd	2.42	2.28	2.4	2.5	2.1	2.3	4.36	0.30	14	4	4	12	0.313
¹¹⁰ Pd	2.46	2.29	2.5	2.0	2.2	2.3	4.01	0.26	12	10	4	14	0.338
¹¹² Pd	2.53	2.33	2.6	2.5	2.1	2.3	4.11	0.60	6	0	3	5	0.485
¹¹⁴ Pd	2.56	2.31	2.6	2.9	2.1	2.3	5.12	0.24	16	0	11	18	0.727
¹¹⁶ Pd	2.58	2.34	3.3	3.5	2.2	2.3	7.44	0.13	16	0	9	16	0.626
¹⁰⁶ Cd	2.36	2.37	2.8	2.9	2.7	2.4	4.45	0.62	12	0	2	7	0.196
¹⁰⁸ Cd	2.38	2.26	2.7	2.5	2.5	2.3	3.97	0.43	18	0	5	13	0.688
¹¹⁰ Cd	2.35	2.24	2.2	2.2	2.2	2.2	3.66	0.47	16	6	5	15	0.269
¹¹² Cd	2.29	2.23	2.0	2.1	2.1	2.2	3.55	0.50	12	8	11	20	0.542
¹¹⁴ Cd	2.30	2.21	2.0	2.0	2.2	2.2	3.43	0.51	14	4	3	11	0.359
¹¹⁶ Cd	2.38	2.29	2.5	2.7	2.4	2.3	4.10	0.47	14	2	3	10	0.408
¹¹⁸ Cd	2.39	2.29	2.6	2.7	2.6	2.3	4.11	0.47	14	0	3	9	0.313
¹²⁰ Cd	2.38	2.28	2.7	2.6	2.6	2.3	4.09	0.44	16	0	2	9	0.379
¹¹⁸ Xe	2.40	2.31	2.5	2.7	2.8	2.3	5.40	0.19	16	4	10	19	0.343
¹²⁰ Xe	2.47	2.35	2.8	3.8	2.7	2.4	7.05	0.16	26	4	9	23	0.652
¹²² Xe	2.50	2.43	3.5	3.5	2.5	2.4	5.03	0.66	10	0	9	13	0.501
¹²⁴ Xe	2.48	2.35	3.6	3.8	2.4	2.4	6.88	0.17	20	2	11	21	0.562
¹²⁶ Xe	2.42	2.33	3.4	3.2	2.3	2.3	6.12	0.18	12	4	9	16	0.576
¹²⁸ Xe	2.33	2.33	3.6	3.3	2.2	2.3	4.74	0.39	10	2	7	12	0.522
¹³⁰ Xe	2.25	2.27	3.3	2.6	2.1	2.3	4.05	0.44	14	0	5	11	0.476
¹³² Xe	2.16	2.17	2.8	1.4	1.9	2.2	3.16	0.66	6	0	5	7	0.731
¹³⁴ Xe	2.04	2.11	1.9	1.0	1.9	2.1	2.91	0.74	6	0	5	7	0.753
¹³⁰ Ba	2.52	2.46	3.3	3.3	2.5	2.5	5.58	0.77	12	0	6	11	0.416
¹³² Ba	2.43	2.29	3.2	2.7	2.2	2.3	4.63	0.29	14	0	8	14	0.609
¹³⁴ Ba	2.32	2.26	2.9	2.4	1.9	2.3	3.82	0.50	8	0	4	7	0.483
¹³⁶ Ba	2.28	2.18	1.9	1.7	1.9	2.2	3.24	0.60	6	0	2	4	0.454
¹⁴² Ba	2.32	2.44	4.3	4.3	4.0	2.4	5.45	0.60	14	0	2	8	0.605
¹³⁴ Ce	2.56	2.37	3.7	4.1	2.4	2.4	6.00	0.27	34	2	8	25	0.502
¹³⁶ Ce	2.38	2.24	1.9	2.2	2.0	2.2	3.66	0.47	16	0	3	10	0.618
¹³⁸ Ce	2.32	2.23	1.9	2.1	1.9	2.2	3.55	0.50	14	0	2	8	1.308
¹⁴⁰ Nd	2.33	2.19	1.8	1.7	1.9	2.2	3.27	0.60	6	0	2	4	0.265
¹⁴⁸ Nd	2.49	2.32	3.0	2.9	4.1	2.3	6.40	0.14	12	8	4	13	0.810
¹⁴⁰ Sm	2.35	2.38	1.9	1.9	2.7	2.4	4.20	0.77	8	0	2	5	0.153
¹⁴² Sm	2.33	2.20	1.9	1.7	2.2	2.2	3.33	0.61	8	0	2	5	0.173
¹⁴² Gd	2.35	2.28	2.7	2.7	1.9	2.3	4.17	0.42	16	0	2	9	0.188
¹⁴⁴ Gd	2.35	2.36	2.5	2.5	2.5	2.4	4.24	0.65	6	0	2	4	0.102
¹⁵² Gd	2.19	2.26	1.8	2.4	3.2	2.3	3.93	0.40	16	10	7	19	0.436
¹⁵⁴ Dy	2.23	2.28	2.0	2.7	3.1	2.3	4.22	0.38	26	10	7	24	0.371
¹⁵⁶ Er	2.32	2.31	2.7	3.1	2.7	2.3	4.75	0.34	20	4	5	16	0.374
¹⁸⁶ Pt	2.56	2.34	2.5	1.7	3.2	2.3	6.18	0.07	26	6	10	25	1.070
¹⁸⁸ Pt	2.53	2.45	3.0	3.3	2.3	2.5	5.45	0.75	16	2	4	12	0.356
¹⁹⁰ Pt	2.49	2.34	3.1	3.6	2.0	2.3	5.08	0.35	18	2	6	15	0.566
¹⁹² Pt	2.48	2.35	3.8	3.7	1.9	2.3	6.42	0.19	10	0	8	12	0.681
¹⁹⁴ Pt	2.47	2.35	3.9	3.6	1.9	2.3	7.28	0.14	10	4	5	11	0.657
¹⁹⁶ Pt	2.47	2.32	3.2	2.9	1.9	2.3	6.26	0.15	10	2	6	11	0.627
¹⁹⁸ Pt	2.42	2.25	2.2	2.3	1.9	2.3	3.87	0.39	6	2	4	7	0.374
²⁰⁰ Pt	2.35	2.20	2.4	1.7	1.8	2.2	3.31	0.59	4	0	4	5	0.676

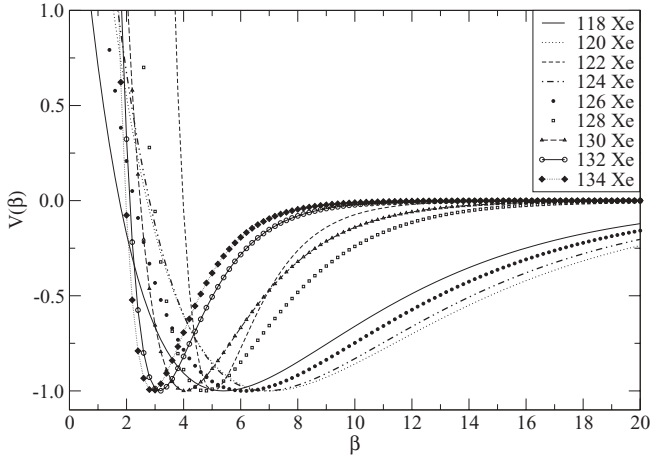


FIG. 1. Evolution of Morse potential shapes for the $_{54}\text{Xe}$ isotopes, with the parameters given in Table I. See Sec. V for further discussion.

The Morse potentials obtained for the $_{54}\text{Xe}$ isotopes are shown in Fig. 1. The evolution of the parameters and the shapes of the potentials are clear. As one moves from $^{134}\text{Xe}_{80}$, which is just below the $N = 82$ magic number, to the midshell nucleus $^{120}\text{Xe}_{66}$, the β_e parameter (which is the position of the minimum of the potential) increases, while the parameter a , which corresponds to the steepness of the potential, decreases. As a result, one gradually obtains less steep potentials with a minimum farther away from the origin. The trends start to be reversed at $^{118}\text{Xe}_{64}$, which is just below midshell.

We have also fitted all nuclei with mass $A \geq 150$ and $R_{4/2} = E(4)/E(2) > 2.9$ for which at least the β_1 and γ_1 bandheads are known, using the exactly separable rotational solution of the Morse potential with $\gamma \approx 0$ (ES-M), which involves three free parameters (the Morse parameters β_e and a , as well as the stiffness C of the γ potential), as described in Sec. III C. All bands are treated on an equal footing, depending on all three parameters. Results for 45 rare earths and 13 actinides are shown in Table II. The quality measure σ of Eq. (39), used in the rms fits, remains below 1 in most cases.

The Morse potentials obtained for the $_{70}\text{Yb}$ isotopes are shown in Fig. 2. The evolution of the parameters and the shapes of the potentials are again clear. As one moves from $^{164}\text{Yb}_{94}$ to the midshell nucleus $^{174}\text{Yb}_{104}$, the β_e parameter (which is the position of the minimum of the β potential) again increases, while the parameter a , which corresponds to the steepness of the β potential, again decreases. The C parameter, which is related to the stiffness of the γ potential, increases. As a result, one gradually obtains less steep β potentials with a minimum farther away from the origin, while the γ potentials get stiffer at the same time.

A notable exception occurs in the $N = 90$ isotones ^{150}Nd , ^{152}Sm , ^{154}Gd , which are known [6,7,58] to be good examples of the X(5) critical point symmetry, along with ^{178}Os [59]. The relative failure of the Morse potential to describe critical nuclei is expected. The potential at the critical point is expected to be flat, as the infinite-well potential used in X(5), or to have a little bump in the middle [8,39]. Microscopic relativistic mean field calculations [60–62] of potential energy surfaces support these assumptions. Since the Morse potential cannot imitate

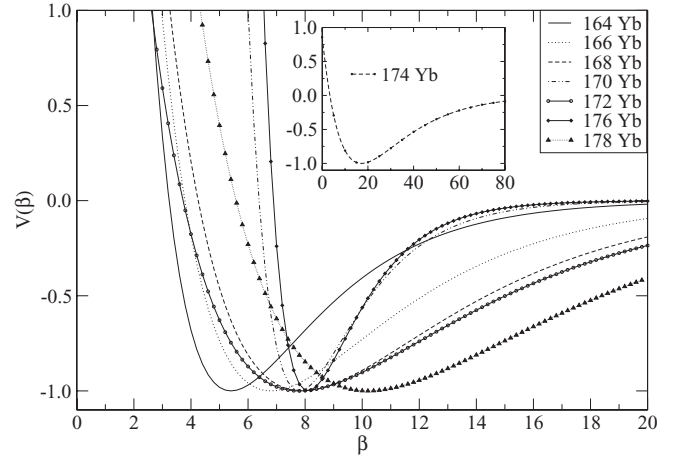


FIG. 2. Evolution of Morse potential shapes for the $_{70}\text{Yb}$ isotopes, with the parameters given in Table II. See Sec. V for further discussion.

a flat potential, with or without a bump in the middle, it is expected that it cannot describe these nuclei satisfactorily.

A comparison of the fits of Table II to the results provided by the Davidson potential in the exactly separable $\gamma \approx 0$ case [31] (ES-D), which contains two free parameters (β_0 , c) instead of three (see Table I of Ref. [31]), shows that the extra parameter extends the region of applicability of the model in most nuclei to higher angular momenta, largely improving the quality of the fits.

As an example, the spectra of ^{154}Dy (γ -unstable case) and ^{232}Th (exactly separable rotational case with $\gamma \approx 0$) are shown in Table III. The overall agreement between theory and experiment is very good in both cases. In the theoretical predictions for the γ band of ^{154}Dy , the O(5) degeneracies are present, limiting the flexibility of the model to agree to experiment. Spacings within all bands of ^{232}Th , including the β band (in which spacings in X(5) are overpredicted by almost a factor of 2 [6–8]), are reproduced very accurately.

For the construction of complete level schemes, the calculation of $B(E2)$ transition rates is required, for which the wave functions are needed. Work in this direction is in progress.

VI. DISCUSSION

The Bohr Hamiltonian has been solved with the Morse potential for any angular momentum, both in the γ -unstable case and in the exactly separable rotational case with $\gamma \approx 0$ (in which a harmonic oscillator is used for the γ potential), labeled as ES-M. The solution has been achieved through the asymptotic iteration method (AIM) and has involved the Pekeris approximation. The effectiveness of AIM has been demonstrated by applying it to the γ -unstable case and to the exactly separable rotational case with $\gamma \approx 0$ for the Davidson and Kratzer potentials.

Numerical results have been presented for both solutions, including all relevant medium mass and heavy nuclei for which at least the β_1 and γ_1 bandheads are known. The success of the present solutions in reproducing quite well both the

TABLE II. Comparison of theoretical predictions of the exactly separable Morse model (ES-M) (described in Sec. III C) and experimental data [47] of rare earth and actinides with $A \geq 150$, $R_{4/2} > 2.9$, and known 0_2^+ and 2_7^+ states. Data for ^{228}Ra come from Ref. [57]. The labels are the same as in Table I.

Nucleus	$R_{4/2}$ exp	$R_{4/2}$ th	$R_{0/2}$ exp	$R_{0/2}$ th	$R_{2/2}$ exp	$R_{2/2}$ th	β_0	C	a	L_g	L_β	L_γ	n	σ
^{150}Nd	2.93	3.21	5.2	6.2	8.2	8.2	5.2	6.0	0.41	14	6	4	13	1.129
^{152}Sm	3.01	3.22	5.6	6.9	8.9	9.5	5.3	7.0	0.34	16	14	9	23	1.007
^{154}Sm	3.25	3.28	13.4	13.5	17.6	18.6	8.4	13.7	0.20	16	6	7	17	0.484
^{154}Gd	3.02	3.20	5.5	7.4	8.1	5.1	5.4	3.4	0.31	26	26	7	32	1.849
^{156}Gd	3.24	3.27	11.8	11.6	13.0	14.4	7.5	10.5	0.22	26	12	16	34	0.605
^{158}Gd	3.29	3.30	15.0	15.4	14.9	15.1	8.3	10.6	0.36	12	6	6	14	0.224
^{160}Gd	3.30	3.31	17.6	17.8	13.1	13.1	8.6	8.8	0.44	16	4	8	17	0.169
^{162}Gd	3.29	3.31	19.8	19.8	12.0	12.0	9.5	8.1	0.30	14	0	4	10	0.082
^{158}Dy	3.21	3.25	10.0	10.1	9.6	10.0	6.7	7.1	0.25	28	8	8	25	0.495
^{160}Dy	3.27	3.28	14.7	14.5	11.1	11.8	9.2	8.2	0.18	28	4	23	38	0.522
^{162}Dy	3.29	3.31	17.3	17.1	11.0	10.8	8.3	7.2	0.44	18	14	14	29	0.312
^{164}Dy	3.30	3.31	22.6	22.2	10.4	10.2	13.1	6.8	0.14	20	0	10	19	0.188
^{166}Dy	3.31	3.31	15.0	15.1	11.2	11.2	7.9	7.5	0.52	6	2	5	8	0.037
^{160}Er	3.10	3.20	7.1	7.9	6.8	3.9	5.6	2.5	0.30	26	2	5	18	1.790
^{162}Er	3.23	3.26	10.7	11.1	8.8	9.9	7.1	7.0	0.26	20	4	12	23	0.588
^{164}Er	3.28	3.27	13.6	12.9	9.4	9.6	9.3	6.6	0.14	22	10	18	33	0.827
^{166}Er	3.29	3.30	18.1	17.8	9.8	9.6	9.7	6.5	0.23	16	10	14	26	0.306
^{168}Er	3.31	3.32	15.3	15.5	10.3	10.2	8.1	6.7	0.59	18	6	8	19	0.176
^{170}Er	3.31	3.29	11.3	9.9	11.9	13.1	6.5	9.0	0.17	24	10	19	35	0.864
^{164}Yb	3.13	3.21	7.9	7.4	7.0	7.3	5.4	5.2	0.32	18	0	5	13	0.471
^{166}Yb	3.23	3.25	10.2	10.1	9.1	9.5	6.8	6.7	0.23	24	10	13	29	0.688
^{168}Yb	3.27	3.27	13.2	11.9	11.2	11.4	7.9	8.0	0.19	34	4	7	25	0.768
^{170}Yb	3.29	3.31	12.7	13.8	13.6	13.5	7.8	9.2	0.49	20	10	17	31	0.509
^{172}Yb	3.31	3.29	13.2	12.4	18.6	19.1	7.8	13.7	0.17	16	12	5	18	0.851
^{174}Yb	3.31	3.31	19.4	19.3	21.4	21.5	17.2	15.1	0.05	20	4	5	16	0.170
^{176}Yb	3.31	3.32	13.9	14.0	15.4	15.3	8.1	10.4	0.57	20	2	5	15	0.254
^{178}Yb	3.31	3.29	15.7	15.6	14.5	14.5	10.2	10.3	0.15	6	4	2	6	0.045
^{168}Hf	3.11	3.22	7.6	7.5	7.1	7.4	5.4	5.3	0.32	22	4	4	16	0.438
^{170}Hf	3.19	3.23	8.7	8.6	9.5	8.1	6.0	5.7	0.27	34	4	4	22	0.964
^{172}Hf	3.25	3.26	9.2	10.1	11.3	11.7	6.7	8.5	0.24	38	4	6	26	0.444
^{174}Hf	3.27	3.26	9.1	9.0	13.5	14.1	6.1	10.4	0.26	26	26	5	30	0.484
^{176}Hf	3.28	3.28	13.0	12.3	15.2	16.1	7.9	11.7	0.18	18	10	8	21	0.622
^{178}Hf	3.29	3.28	12.9	12.7	12.6	12.9	8.5	9.2	0.16	18	6	6	17	0.298
^{180}Hf	3.31	3.31	11.8	12.2	12.9	12.9	7.7	8.7	0.61	12	4	5	12	0.199
^{176}W	3.22	3.24	7.8	8.2	9.6	10.2	5.8	7.4	0.29	22	12	5	21	0.578
^{178}W	3.24	3.25	9.4	9.5	10.5	10.5	6.4	7.6	0.26	18	10	2	15	0.177
^{180}W	3.26	3.27	14.6	12.7	10.8	11.4	7.9	8.0	0.24	24	0	7	18	0.838
^{182}W	3.29	3.28	11.3	11.6	12.2	12.4	8.3	8.6	0.13	18	4	6	16	0.282
^{184}W	3.27	3.27	9.0	8.9	8.1	8.1	6.6	5.5	0.15	10	4	6	12	0.094
^{186}W	3.23	3.24	7.2	7.2	6.0	6.2	5.3	4.2	0.22	14	4	6	14	0.133
^{178}Os	3.02	3.19	4.9	5.8	6.6	7.1	4.8	5.1	0.39	16	6	5	15	0.724
^{180}Os	3.09	3.23	5.6	7.1	6.6	7.0	5.1	4.9	0.27	10	6	7	14	1.122
^{184}Os	3.20	3.24	8.7	9.4	7.9	8.5	6.3	6.0	0.28	22	0	6	16	0.600
^{186}Os	3.17	3.22	7.7	7.7	5.6	6.1	5.6	4.2	0.27	14	10	13	24	0.200
^{188}Os	3.08	3.20	7.0	7.3	4.1	4.3	5.3	2.8	0.34	12	2	7	13	0.213
^{228}Ra	3.21	3.27	11.3	11.3	13.3	13.0	7.3	9.5	0.24	22	4	3	15	0.387
^{228}Th	3.24	3.28	14.4	13.9	16.8	17.0	8.4	12.4	0.25	18	2	5	14	0.514
^{230}Th	3.27	3.28	11.9	11.7	14.7	14.8	7.6	10.7	0.19	24	4	4	17	0.276
^{232}Th	3.28	3.29	14.8	14.7	15.9	16.5	9.6	11.8	0.15	30	20	12	36	0.321
^{232}U	3.29	3.29	14.5	14.4	18.2	18.3	9.5	13.1	0.14	20	10	4	18	0.158
^{234}U	3.30	3.30	18.6	18.7	21.3	21.7	12.1	15.5	0.12	28	8	7	24	0.219
^{236}U	3.30	3.30	20.3	20.4	21.2	21.2	13.8	15.0	0.10	30	4	5	21	0.266
^{238}U	3.30	3.31	20.6	20.9	23.6	24.7	13.9	17.6	0.10	30	4	27	43	0.716
^{238}Pu	3.31	3.32	21.4	21.7	23.3	23.4	9.8	15.9	0.50	26	2	4	17	0.857
^{240}Pu	3.31	3.31	20.1	19.1	26.6	27.1	13.0	19.4	0.09	26	4	4	18	0.539
^{242}Pu	3.31	3.32	21.5	21.6	24.7	24.8	9.9	17.0	0.46	26	2	2	15	0.926
^{248}Cm	3.31	3.31	25.0	25.1	24.2	24.2	16.4	17.0	0.09	28	4	2	17	0.105
^{250}Cf	3.32	3.32	27.0	27.0	24.2	24.2	12.8	16.7	0.20	8	2	4	8	0.018

TABLE III. Theoretical predictions of the γ -unstable Morse potential (described in Sec. IV C) with $\beta_0 = 4.22$ and $a = 0.38$ compared with experimental data for ^{154}Dy [47], and theoretical predictions of the exactly separable Morse (ES-M) (described in Sec. III C) with $\beta_0 = 9.6$, $C = 11.8$, $a = 0.15$ compared with experimental data for ^{232}Th [47]. All states are normalized to the 2_1^+ state. See Sec. V for further discussion.

L	^{154}Dy		^{232}Th		^{154}Dy		^{232}Th		L	^{154}Dy		^{232}Th	
	gsb	gsb	gsb	gsb	β_1	β_1	β_1	β_1		γ_1	γ_1	γ_1	γ_1
	exp	th	exp	th	exp	th	exp	th		exp	th	exp	th
0	0.00	0.00	0.00	0.00	1.98	2.66	14.79	14.72	2	3.07	2.28	15.91	16.48
2	1.00	1.00	1.00	1.00	2.71	3.35	15.68	15.63	3	3.99	3.71	16.80	17.32
4	2.23	2.28	3.28	3.29	3.74	4.24	17.68	17.72	4	4.31	3.71	18.03	18.42
6	3.66	3.71	6.75	6.76	4.96	5.25	20.72	20.90	5	5.20	5.25	19.45	19.79
8	5.22	5.25	11.28	11.29	6.47	6.37	24.75	25.06	6	5.64	5.25	21.27	21.40
10	6.89	6.86	16.75	16.73	8.24	7.58	29.76	30.08	7	6.53	6.86	23.21	23.26
12	8.65	8.56	23.03	22.97			35.55	35.86	8			25.50	25.35
14	10.49	10.37	30.04	29.89			42.14	42.32	9			27.75	27.65
16	12.47	12.30	37.65	37.43			49.44	49.38	10			30.62	30.17
18	14.55	14.36	45.84	45.52			57.36	57.03	11			33.22	32.88
20	16.71	16.58	54.52	54.16			65.81	65.23	12			36.48	35.78
22	18.98	18.95	63.69	63.31									
24	21.40	21.48	73.32	72.99									
26	23.99	24.19	83.38	83.20									
28			93.82	93.94									
30			104.56	105.24									

bandheads of and the spacings within the ground, β_1 , and γ_1 bands indicates that a detailed study of γ_2 and β_2 bands within this framework might be fruitful, although the difficulties in singling out the experimental β_2 band [63] should be kept in mind. The influence of the finite depth of the potential is also worth considering in more detail. From the findings of Ref. [26], where the E(5) case was solved for a finite well, the influence of the finite depth of the potential is expected to show up more clearly in the higher excited states. Work on the calculation of wave functions and $B(E2)$ transition rates is in progress.

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APPENDIX A: EXACTLY SEPARABLE $\gamma \approx 0$ SOLUTION FOR THE DAVIDSON POTENTIAL

Assuming the reduced potential to be of the form of Eq. (1), $u(\beta, \gamma) = u(\beta) + u(\gamma)/\beta^2$, and plugging it into the Bohr Hamiltonian of Eq. (14), we obtain the “radial” and γ equations [31]

$$\left[-\frac{1}{\beta^4} \frac{\partial}{\partial \beta} \beta^4 \frac{\partial}{\partial \beta} + \frac{L(L+1)}{3\beta^2} + \frac{\lambda}{\beta^2} + u(\beta) \right] \times \xi_L(\beta) = \epsilon \xi_L(\beta), \quad (\text{A1})$$

$$\left[-\frac{1}{\sin 3\gamma} \frac{\partial}{\partial \gamma} \sin 3\gamma \frac{\partial}{\partial \gamma} + \frac{K^2}{4} \left(\frac{1}{\sin^2(\gamma)} - \frac{4}{3} \right) + u(\gamma) \right] \times \Gamma_K(\gamma) = \lambda \Gamma_K(\gamma). \quad (\text{A2})$$

Assuming $u(\gamma) = (3c)^2 \gamma^2$ and expanding Eq. (A2) in powers of γ for $\gamma \simeq 0$, we get [31]

$$\left[-\frac{1}{\gamma} \frac{\partial}{\partial \gamma} \gamma \frac{\partial}{\partial \gamma} + \frac{K^2}{4\gamma^2} + (3c^2)\gamma^2 \right] \Gamma_K(\gamma) = \epsilon_\gamma \Gamma_K(\gamma), \quad (\text{A3})$$

where $\epsilon_\gamma = \lambda + K^2/3$. The solution of this equation is given [31] in terms of Laguerre polynomials with

$$\epsilon_\gamma = (3C)(n_\gamma + 1), \quad C = 2c, \quad n_\gamma = 0, 1, 2, 3, \dots \quad (\text{A4})$$

We now solve the radial equation by using AIM. Plugging the Davidson potential of Eq. (3), $u(\beta) = \beta^2 + \beta_0^4/\beta^2$, into Eq. (A1), we get

$$\left[-\frac{1}{\beta^4} \frac{\partial}{\partial \beta} \beta^4 \frac{\partial}{\partial \beta} + \frac{\frac{L(L+1)}{3} + \lambda + \beta_0^4}{\beta^2} + \beta^2 \right] \xi_L(\beta) = \epsilon \xi_L(\beta). \quad (\text{A5})$$

Transforming ξ_L into χ_L by the relation

$$\xi_L(\beta) = \beta^{-2} \chi_L(\beta), \quad (\text{A6})$$

and plugging it into Eq. (A5), we obtain

$$\chi_L''(\beta) + \left[\epsilon - \frac{\frac{L(L+1)}{3} + \beta_0^4 + \lambda + 2}{\beta^2} - \beta^2 \right] \chi_L(\beta) = 0. \quad (\text{A7})$$

To simplify this equation, we define

$$\frac{L(L+1)}{3} + \lambda + \beta_0^4 + 2 = \mu(\mu+1), \quad (\text{A8})$$

obtaining

$$\chi_L''(\beta) + \left[\epsilon - \frac{\mu(\mu+1)}{\beta^2} - \beta^2 \right] \chi_L(\beta) = 0. \quad (\text{A9})$$

This second-order differential equation must have a solution of the form

$$\chi_L(\beta) = \beta^{\mu+1} e^{-\frac{\beta^2}{2}} f_{n,L}(\beta). \quad (\text{A10})$$

Using this function in Eq. (A9), one can get

$$f_{n,L}''(\beta) = \left(2\beta - \frac{2\mu+2}{\beta} \right) f_{n,L}'(\beta) + (2\mu+3-\epsilon) f_{n,L}(\beta). \quad (\text{A11})$$

From Eq. (5), one can define $\lambda_0(\beta)$ and $s_0(\beta)$ as

$$\lambda_0(\beta) = 2\beta - \frac{2\mu+2}{\beta}, \quad s_0(\beta) = 2\mu+3-\epsilon. \quad (\text{A12})$$

Then, $s_k(x)$ and $\lambda_k(x)$ are calculated by the recurrence relations of Eq. (8) as

$$\lambda_1(\beta) = \frac{4\beta^4 - (6\mu + \epsilon + 3)\beta^2 + 4\mu^2 + 10\mu + 6}{\beta^2}, \quad (\text{A13})$$

$$s_1(\beta) = \frac{2(2\mu + 3 - \epsilon)(\beta^2 - \mu - 1)}{\beta}, \quad (\text{A14})$$

$$\lambda_2(\beta) = \frac{-4[(6 + 13\mu + 9\mu^2 + 2\mu^3) - \beta^2(3 + \epsilon + 7\mu + \mu\epsilon + 4\mu^2) + \beta^4(4\mu + \epsilon) - 2\beta^6]}{\beta^3}, \quad (\text{A15})$$

$$s_2(\beta) = \frac{(24 + 52\mu - 8\epsilon + 36\mu^2 - 12\epsilon\mu - 4\epsilon\mu^2 + 8\mu^3)}{\beta^2} + \beta^2(12 + 8\mu - 4\epsilon) - (3 + 2\epsilon + 20\mu - 4\mu\epsilon + 12\mu^2 - \epsilon^2), \quad (\text{A16})$$

⋮

By applying the termination condition of AIM, Eq. (9), we obtain the energy eigenvalues from the roots of Δ_k as

$$\epsilon_0 = \mu + \frac{3}{2}, \quad \epsilon_1 = \mu + \frac{7}{2}, \quad \epsilon_2 = \mu + \frac{11}{2}, \quad \dots, \quad (\text{A17})$$

while the general expression is

$$\epsilon_{n,\mu} = \mu + \frac{3}{2} + 2n. \quad (\text{A18})$$

Then, substituting the value of μ from Eq. (A8) yields the energy eigenvalues

$$\epsilon_{n,L} = 2n + 1 + \left[\frac{9}{4} + \frac{L(L+1) - K^2}{3} + 3C(n_\gamma + 1) + \beta_0^4 \right]^{1/2}, \quad (\text{A19})$$

which are identical to the ones found in Ref. [31].

APPENDIX B: EXACTLY SEPARABLE $\gamma \approx 0$ SOLUTION FOR THE KRATZER POTENTIAL

Separation of variables proceeds as in Appendix A. The radial equation is

$$\chi_L''(\beta) + \left[\epsilon - \frac{\frac{L(L+1)}{3} + 2 + \lambda}{\beta^2} + \frac{A}{\beta} - \frac{B}{\beta^2} \right] \chi_L(\beta) = 0, \quad (\text{B1})$$

where we have used the wave function of Eq. (16) and the transformation $\xi_L(\beta) = \beta^{-2} \chi_L(\beta)$. This equation differs from the corresponding one of Ref. [29] by the term λ , which comes

from the exact separation of variables and will be determined below from the γ equation. Defining a new parameter set

$$\frac{L(L+1)}{3} + 2 + \lambda + B = p(p+1), \quad (\text{B2})$$

$$\epsilon = -\epsilon, \quad 2\beta\sqrt{\epsilon} = x, \quad \frac{A}{2\sqrt{\epsilon}} = k, \quad (\text{B3})$$

Eq. (B1) becomes

$$\chi_L''(x) + \left[-\frac{1}{4} - \frac{p(p+1)}{x^2} + \frac{k}{x} \right] \chi_L(x) = 0. \quad (\text{B4})$$

This second-order differential equation must have a solution of the form

$$\chi_L(x) = x^{p+1} e^{-\frac{x^2}{2}} R_{n,L}(x), \quad (\text{B5})$$

which leads to

$$R_{n,L}''(x) = \left(\frac{x-2p-2}{x} \right) R_{n,L}'(x) + \left(\frac{p+1-k}{x} \right) R_{n,L}(x). \quad (\text{B6})$$

According to Eq. (5) of AIM, one then has

$$\lambda_0(x) = \frac{x-2p-2}{x}, \quad s_0(x) = \frac{p+1-k}{x}. \quad (\text{B7})$$

Using the recurrence relations of Eq. (8), one can then determine $\lambda_k(x)$ and $s_k(x)$ as

$$\lambda_1(x) = \frac{10p+6-3xp-3x-kx+x^2+4p^2}{x^2}, \quad (\text{B8})$$

$$s_1(x) = \frac{-5p - 3 + 3k + xp - 2p^2 + x - kx + 2kp}{x^2}, \quad (\text{B9})$$

$$\lambda_2(x) = \frac{(-24 - 52p + 12x - 4x^2 - 36p^2 + 6kx + 20xp + x^3 + 8p^2x - 4px^2 + 4pkx - 8p^3 - 2kx^2)/x^3,}{(\text{B10})}$$

$$s_2(x) = \frac{[(1 + p - k)x^2 + (3k - 4 - 7p - 3p^2 + k^2 + 2pk)x + 2(6 + 13p - 6k + 9p^2 - 7kp + 2p^3 - 2kp^2)]/x^3,}{\vdots} \quad (\text{B11})$$

After determining $\Delta_k(x)$ by using the termination condition in Eq. (9), the energy eigenvalues are found from the roots of $\Delta_k(x)$ as

$$k_0 = p + 1, \quad k_1 = p + 2, \quad k_2 = p + 3, \dots, \quad (\text{B12})$$

generalized into

$$k_n = p + 1 + n. \quad (\text{B13})$$

From Eqs. (B2) and (B3), one then obtains

$$\varepsilon_{n,L} = \frac{A^2/4}{\left(n + \frac{1}{2} + \sqrt{\frac{9}{4} + \lambda + B + \frac{L(L+1)}{3}}\right)^2}. \quad (\text{B14})$$

To find λ , we have to solve the γ equation of Eq. (A2), using [29] $u(\gamma) = c\gamma^2/2$. Expanding around $\gamma = 0$ and taking $\varepsilon_\gamma = \lambda + K^2/3$, one obtains

$$\left[-\frac{1}{\gamma} \frac{\partial}{\partial \gamma} \gamma \frac{\partial}{\partial \gamma} + \frac{K^2}{4\gamma^2} + \frac{c\gamma^2}{2}\right] \Gamma_K(\gamma) = \varepsilon_\gamma \Gamma_K(\gamma), \quad (\text{B15})$$

which, through the transformation $\Gamma_K(\gamma) = \gamma^{-1/2} \xi_K(\gamma)$, leads to

$$\xi_K''(\gamma) + \left[\varepsilon_\gamma + \frac{(1 - K^2)/4}{\gamma^2} - \frac{c\gamma^2}{2}\right] \xi_K(\gamma) = 0. \quad (\text{B16})$$

Defining

$$\left(\frac{c}{2}\right)^{1/4} \gamma = y, \quad \mu(\mu + 1) = (K^2 - 1)/4, \quad (\text{B17})$$

$$\varepsilon_\gamma = \left(\frac{c}{2}\right)^{-1/2} \varepsilon_\gamma,$$

this equation is brought into the form

$$\xi_K''(y) + \left[\varepsilon_\gamma - \frac{\mu(\mu + 1)}{y^2} - y^2\right] \xi_K(y) = 0, \quad (\text{B18})$$

which is suitable for solving through AIM, by considering a solution of the form

$$\xi_K(y) = y^{\mu+1} e^{-\frac{y^2}{2}} G_{m,K}(y). \quad (\text{B19})$$

Following the same procedure as above, we obtain the differential equation in the form

$$G_{m,K}''(y) = \left(\frac{2y^2 - 2\mu - 2}{y}\right) G_{m,K}'(y) + \left(\frac{2y\mu + 3y - y\varepsilon_\gamma}{y}\right) G_{m,K}(y). \quad (\text{B20})$$

Comparison with Eq. (5) leads to

$$\lambda_0(y) = \frac{2y^2 - 2\mu - 2}{y}, \quad s_0(y) = 2\mu + 3 - \varepsilon_\gamma. \quad (\text{B21})$$

By using the recurrence relations of Eq. (8) and the termination condition of Eq. (9), one can find the energy eigenvalues to be

$$(\varepsilon_\gamma)_0 = 2\mu + 3, \quad (\varepsilon_\gamma)_1 = 2\mu + 7, \quad (\varepsilon_\gamma)_2 = 2\mu + 11, \dots, \quad (\text{B22})$$

and the generalized form is

$$(\varepsilon_\gamma)_{\mu,m} = 2\mu + 3 + 4m. \quad (\text{B23})$$

Using μ from Eq. (B17), this becomes

$$(\varepsilon_\gamma)_{K,m} = \left(\frac{c}{2}\right)^{1/2} [4m + 2 + K], \quad (\text{B24})$$

leading to

$$\lambda = \left(\frac{c}{2}\right)^{1/2} [4m + 2 + K] - K^2/3. \quad (\text{B25})$$

This result agrees with Ref. [29], with $K = 2n_\gamma - 4m$, $m = 0, 1, \dots, n_\gamma$, as stated there. It is also in full agreement with the results of Ref. [31], taking into account the different coefficients of γ^2 in $u(\gamma)$.

APPENDIX C: EXACTLY SEPARABLE $\gamma \approx 0$ SOLUTION FOR THE MORSE POTENTIAL

Separation of variables again proceeds as in Appendix A. Using the transformation $\xi_L(\beta) = \beta^{-2} \chi_L(\beta)$, the radial equation becomes

$$\chi_L''(\beta) + \left[\varepsilon - \frac{\frac{L(L+1)}{3} + \lambda + 2}{\beta^2} - e^{-2a(\beta-\beta_e)} + 2e^{-a(\beta-\beta_e)}\right] \chi_L(\beta) = 0. \quad (\text{C1})$$

Defining

$$x = \frac{\beta - \beta_e}{\beta_e}, \quad \alpha = a\beta_e, \quad (\text{C2})$$

$$\beta_e^2 \varepsilon = \varepsilon, \quad \frac{L(L+1)}{3} + 2 + \lambda = \mu,$$

the radial equation becomes

$$\chi_L''(x) + \left[\varepsilon - \frac{\mu}{(1+x)^2} - \beta_e^2 e^{-2\alpha x} + 2\beta_e^2 e^{-\alpha x}\right] \chi_L(x) = 0. \quad (\text{C3})$$

We now apply the Pekeris approximation [45]. Renaming $\frac{\mu}{(1+x)^2}$ as $u_L(x)$ and expanding in a series around $x = 0$, we obtain

$$u_L(x) = \mu(1 - 2x + 3x^2 - 4x^3 + \dots). \quad (\text{C4})$$

In the exponential form, $u_L(x)$ can be written as

$$\tilde{u}_L(x) = \mu(c_0 + c_1 e^{-\alpha x} + c_2 e^{-2\alpha x} + \dots). \quad (\text{C5})$$

Expanding also this equation in a series around $x = 0$ produces

$$\tilde{u}_L(x) = \mu \left(c_0 + c_1 + c_2 - [c_1 + 2c_2]\alpha x + \left[\frac{c_1}{2} + 2c_2 \right] \alpha^2 x^2 + \dots \right). \quad (\text{C6})$$

Comparing Eqs. (C4) and (C6), one can now determine the c_i coefficients

$$c_0 = 1 - \frac{3}{\alpha} + \frac{3}{\alpha^2}, \quad c_1 = \frac{4}{\alpha} - \frac{6}{\alpha^2}, \quad c_2 = -\frac{1}{\alpha} + \frac{3}{\alpha^2}. \quad (\text{C7})$$

Returning to the radial equation,

$$\chi_L''(x) + [\varepsilon - \mu(c_0 + c_1 e^{-\alpha x} + c_2 e^{-2\alpha x}) - \beta_e^2 e^{-2\alpha x} + 2\beta_e^2 e^{-\alpha x}] \chi_L(x) = 0, \quad (\text{C8})$$

and by using the ansatz

$$\varepsilon - \mu c_0 = -\rho^2, \quad 2\beta_e^2 - \mu c_1 = \gamma_1^2, \quad \beta_e^2 + \mu c_2 = \gamma_2^2, \quad (\text{C9})$$

we get

$$\chi_L''(x) + [-\rho^2 + \gamma_1^2 e^{-\alpha x} - \gamma_2^2 e^{-2\alpha x}] \chi_L(x) = 0. \quad (\text{C10})$$

Rewriting this equation by using the new variable $y = e^{-\alpha x}$, we obtain

$$\chi_L''(y) + \frac{1}{y} \chi_L'(y) + \left[-\frac{\rho^2}{\alpha^2 y^2} + \frac{\gamma_1^2}{\alpha^2 y} - \frac{\gamma_2^2}{\alpha^2} \right] \chi_L(y) = 0. \quad (\text{C11})$$

Inserting a wave function of the form

$$\chi_L(y) = y^{\frac{\rho}{\alpha}} e^{-\frac{\gamma_2}{\alpha} y} R_{n,L}(y), \quad (\text{C12})$$

the second-order differential equation becomes

$$R_{n,L}''(y) = \left(\frac{2\gamma_2 \alpha y - 2\alpha \rho - \alpha^2}{\alpha^2 y} \right) R_{n,L}'(y) + \left(\frac{2\rho \gamma_2 + \alpha \gamma_2 - \gamma_1^2}{\alpha^2 y} \right) R_{n,L}(y). \quad (\text{C13})$$

Comparison with Eq. (11) leads to the identifications

$$\begin{aligned} \tau(y) &= 2\gamma_2 \alpha y - 2\rho - \alpha, \\ \sigma(y) &= \alpha^2 y, \\ \Omega_n &= 2\gamma_2 \rho + \alpha \gamma_2 - \gamma_1^2. \end{aligned} \quad (\text{C14})$$

From Eq. (13), we get

$$2\gamma_2 \rho + \alpha \gamma_2 - \gamma_1^2 = -n(2\gamma_2 \alpha), \quad (\text{C15})$$

while the generalized form is written as

$$\rho_{n,L} = \frac{\gamma_1^2}{2\gamma_2} - \left(n + \frac{1}{2} \right) \alpha. \quad (\text{C16})$$

Using Eq. (C9), one then obtains

$$\epsilon_{n,L} = \frac{\mu c_0}{\beta_e^2} - \left[\frac{\gamma_1^2}{2\beta_e \gamma_2} - \left(n + \frac{1}{2} \right) \frac{\alpha}{\beta_e} \right]^2. \quad (\text{C17})$$

APPENDIX D: γ -UNSTABLE SOLUTION FOR THE DAVIDSON POTENTIAL

In this case, the reduced potential depends only on β . Using the wave function of Eq. (29), the relevant radial equation becomes

$$\chi_\tau''(\beta) + \left[\epsilon - \beta^2 - \frac{p(p+1)}{\beta^2} \right] \chi_\tau(\beta) = 0, \quad (\text{D1})$$

where $\chi_\tau(\beta) = \beta^{-2} R_\tau(\beta)$ and

$$p(p+1) = \tau(\tau+3) + \beta_0^4 + 2. \quad (\text{D2})$$

For this differential equation, one looks for a solution of the form

$$\chi_\tau(\beta) = \beta^p e^{-\frac{\beta^2}{2}} f_{n,\tau}(\beta). \quad (\text{D3})$$

Using this function in Eq. (D1), we obtain a differential equation that is similar to Eq. (5), i.e.,

$$\begin{aligned} f_{n,\tau}''(\beta) &= \left(\frac{2\beta^3 - 2p\beta}{\beta^2} \right) f_{n,\tau}'(\beta) \\ &+ \left(\frac{\beta^2 + 2p + 2p\beta^2 - \epsilon\beta^2}{\beta^2} \right) f_{n,\tau}(\beta). \end{aligned} \quad (\text{D4})$$

Comparing to Eq. (5), we identify

$$\lambda_0(\beta) = \frac{2\beta^3 - 2p\beta}{\beta^2}, \quad s_0(\beta) = \frac{\beta^2 + 2p + 2p\beta^2 - \epsilon\beta^2}{\beta^2}. \quad (\text{D5})$$

Then the recurrence relations of Eq. (8) give $\lambda_k(\beta)$ and $s_k(\beta)$, while by using the termination relations of Eq. (9), we can obtain the energy eigenvalues

$$\epsilon_0 = p + \frac{3}{2}, \quad \epsilon_1 = p + \frac{7}{2}, \quad \epsilon_2 = p + \frac{11}{2}, \dots, \quad (\text{D6})$$

which give the generalized form

$$\epsilon_{n,p} = 2n + p + \frac{3}{2}. \quad (\text{D7})$$

Substituting in this expression the value of p from Eq. (D2), one gets

$$\epsilon_{n,\tau} = 2n + 1 + \left[\frac{9}{4} + \tau(\tau+3) + \beta_0^4 \right]^{1/2}, \quad (\text{D8})$$

in agreement with the result obtained in Ref. [25].

APPENDIX E: γ -UNSTABLE SOLUTION FOR THE KRATZER POTENTIAL

Using wave functions of the form $\Psi(\beta, \gamma, \theta_i) = \chi(\beta) \Phi(\gamma, \theta_i)$, the radial equation becomes

$$\left[-\frac{1}{\beta^4} \frac{\partial}{\partial \beta} \beta^4 \frac{\partial}{\partial \beta} + \frac{\tau(\tau+3)}{\beta^2} - \frac{A}{\beta} + \frac{B}{\beta^2} \right] \chi(\beta) = \epsilon \chi(\beta), \quad (\text{E1})$$

in agreement with Ref. [19]. Substituting $\chi(\beta) = \beta^{-2} \xi(\beta)$, this becomes

$$\xi''(\beta) + \left[\epsilon - \frac{\tau(\tau+3) + B + 2}{\beta^2} + \frac{A}{\beta} \right] \xi(\beta) = 0. \quad (\text{E2})$$

Using the parameter set

$$\begin{aligned} \epsilon &= -\varepsilon, \quad 2\beta\sqrt{\varepsilon} = y, \\ \frac{A}{2\sqrt{\varepsilon}} &= k, \quad \tau(\tau + 3) + B + 2 = \nu(\nu + 1), \end{aligned} \quad (\text{E3})$$

the differential equation becomes

$$\xi''_{\tau}(y) + \left[-\frac{1}{4} + \frac{k}{y} - \frac{\nu(\nu + 1)}{y^2} \right] \xi_{\tau}(y) = 0. \quad (\text{E4})$$

Assuming that this equation has a solution of the form

$$\xi_{\tau}(y) = y^{\nu+1} e^{-\frac{y}{2}} R_{n,\tau}(y), \quad (\text{E5})$$

we bring it into the form

$$R''_{n,\tau}(y) = \left(\frac{y - 2\nu - 2}{y} \right) R'_{n,\tau}(y) + \left(\frac{\nu + 1 - k}{y} \right) R_{n,\tau}(y). \quad (\text{E6})$$

Comparison with Eq. (5) then provides

$$\lambda_0(y) = \frac{y - 2\nu - 2}{y}, \quad s_0(y) = \frac{\nu + 1 - k}{y}. \quad (\text{E7})$$

By using the recurrence relations of Eq. (8) and the termination conditions of Eq. (9), one then obtains the following energy eigenvalues from the roots of $\Delta_i(y)$:

$$k_0 = \nu + 1, \quad k_1 = \nu + 2, \quad k_2 = \nu + 3, \dots \quad (\text{E8})$$

These are generalized into

$$k_n = \nu + 1 + n. \quad (\text{E9})$$

From Eq. (E3), one then obtains the energy eigenvalues

$$\varepsilon_{n,\tau} = \frac{A^2/4}{\left(n + \frac{1}{2} + \sqrt{\frac{9}{4} + \tau(\tau + 3) + B} \right)^2}, \quad (\text{E10})$$

in agreement with Ref. [19].

APPENDIX F: γ -UNSTABLE SOLUTION FOR THE MORSE POTENTIAL

Using a wave function of the form of Eq. (29), the radial equation is

$$\left[-\frac{1}{\beta^4} \frac{\partial}{\partial \beta} \beta^4 \frac{\partial}{\partial \beta} + \frac{\tau(\tau + 3)}{\beta^2} + u(\beta) \right] \xi(\beta) = \epsilon \xi(\beta). \quad (\text{F1})$$

Taking $\xi(\beta) = \beta^{-2} \chi(\beta)$ and

$$\tau(\tau + 3) + 2 = \nu, \quad (\text{F2})$$

this equation becomes

$$\chi''(\beta) + \left[\epsilon - \frac{\nu}{\beta^2} - u(\beta) \right] \chi(\beta) = 0. \quad (\text{F3})$$

Using the parametrization

$$x = \frac{\beta - \beta_e}{\beta_e}, \quad \alpha = a\beta_e, \quad \varepsilon = \beta_e^2 \epsilon, \quad (\text{F4})$$

one obtains

$$\chi''(x) + \left[\varepsilon - \frac{\nu}{(1+x)^2} - \beta_e^2 e^{-2\alpha x} + 2\beta_e^2 e^{-\alpha x} \right] \chi(x) = 0. \quad (\text{F5})$$

Applying now the Pekeris approximation [45] as in Appendix C, we replace $1/(1+x)^2$ by its approximate expression, obtaining

$$\chi''(x) + \left[\varepsilon - \nu(c_0 + c_1 e^{-\alpha x} + c_2 e^{-2\alpha x}) - \beta_e^2 e^{-\alpha x} + 2\beta_e^2 e^{-2\alpha x} \right] \chi(x) = 0. \quad (\text{F6})$$

Using the parametrization

$$\varepsilon - \nu c_0 = -K^2, \quad 2\beta_e^2 - \nu c_1 = \gamma_1^2, \quad \beta_e^2 + \nu c_2 = \gamma_2^2, \quad (\text{F7})$$

the differential equation is brought into the form

$$\chi''(x) + \left[-K^2 + \gamma_1^2 e^{-\alpha x} - \gamma_2^2 e^{-2\alpha x} \right] \chi(x) = 0. \quad (\text{F8})$$

Introducing a new variable $y = e^{-\alpha x}$, one has

$$\chi''_{\tau}(y) + \frac{1}{y} \chi'_{\tau}(y) + \left[-\frac{K^2}{\alpha^2 y^2} + \frac{\gamma_1^2}{\alpha^2 y} - \frac{\gamma_2^2}{\alpha^2} \right] \chi_{\tau}(y) = 0. \quad (\text{F9})$$

Inserting a wave function of the form

$$\chi_{\tau}(y) = y^{\frac{K}{\alpha}} e^{-\frac{\gamma_2}{\alpha} y} f_{n,\tau}(y), \quad (\text{F10})$$

the differential equation becomes

$$\begin{aligned} f''_{n,\tau}(y) &= \left(\frac{2\gamma_2 \alpha y - 2\alpha K - \alpha^2}{\alpha^2 y} \right) f'_{n,\tau}(y) \\ &+ \left(\frac{2K\gamma_2 + \alpha\gamma_2 - \gamma_1^2}{\alpha^2 y} \right) f_{n,\tau}(y). \end{aligned} \quad (\text{F11})$$

Comparison with Eq. (11) leads to

$$\tau(y) = 2\gamma_2 y - 2K_n - \alpha, \quad \sigma(y) = \alpha^2 y, \quad (\text{F12})$$

$$\Omega_n = 2\gamma_2 K_n + \alpha\gamma_2 - \gamma_1^2.$$

Using Eq. (13), one then has

$$K_{n,\tau} = \frac{\gamma_1^2}{2\gamma_2} - \left(n + \frac{1}{2} \right) \alpha. \quad (\text{F13})$$

From Eq. (F7), we obtain the energy eigenvalues

$$\varepsilon_{n,\tau} = \frac{\nu c_0}{\beta_e^2} - \left[\frac{\gamma_1^2}{2\beta_e \gamma_2} - \left(n + \frac{1}{2} \right) \frac{\alpha}{\beta_e} \right]^2. \quad (\text{F14})$$

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