DEFORMED ENERGY LEVELS OF A PSEUDOHARMONIC POTENTIAL: NONRELATIVISTIC QUANTUM MECHANICS

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ABSTRACT

The paper described the exact solutions of Schrödinger equation for diatomic molecules with pseudoharmonic potential $V(r) = D\left(\frac{r}{r_e} - \frac{r_e}{r}\right)^2$, it was solved by applying the Boopp's shift method to first order in the non-commutativity parameter θ and standard perturbation theory, the bound state energy (eigen values) found to be degenerate. It's shown that the noncommutative Hamiltonian (NC) described automatically a particle with spin (1/2) interacted with an external magnetic field, including the usual Zeeman effect in commutative space. Furthermore, I observed the NC Hamiltonian can be described automatically the spin-orbit interaction, I justified my results by calculate the absolute incertitude.

Keywords: *Star product, Boopp's shift method, diatomic molecules, pseudoharmonic potential.*

1. INTRODUCTION

Recently, the three fundamental equations: Schrödinger, Klein-Gordon and Dirac were satisfied a beautiful Physics success at low and high energy, respectively. Furthermore gauge theory and standard model was the second revolution in modern Physics after general relativity. The Schrödinger equation played a crucial for describing many Physics phenomena, the exact solutions for a Colombian and for a harmonic oscillator as well as in an arbitrary number of spatial dimensions represents typical models in Quantum mechanics [1-18]. Although a new concepts of space-time, known by of noncommutative spaces, represent a hop to obtained a new and profound interpretations at microscopic scales, the noncommutativity

is introduced by many ways, the simple approach, it consider the position and momentum operators obeys to the Heisenberg commutation relation, that is similarly to quantize space-time coordinates, when the commutators $[\hat{x}_i, \hat{x}_j] \neq 0$ [22], the formalism of star product, Boopp's shift method and the Seiberg-Witten map are plays a fundamental roles in this new theory. The rich mathematical structure of the noncommutative theory gives a rise to the hop to get a better understanding of Physics phenomena at smallness distances and hop to solve above mentioned problems. The Physics idea of a noncommutative space satisfied by on based, in addition, to usual Quantum commutators, the following special commutators [18-36]:

$$\left[\hat{x}_i^*, \hat{x}_j \right] = i\theta_{ij} \quad (1)$$

The parameters θ^{ij} are an antisymmetric real matrix of dimension square length in the noncommutative canonical-type space, throughout this paper the natural unites ($c = \hbar = 1$) and $(\mu = s)$ are employed. It's important to notice that, the fundamental commutator which represented by eq. (1), it has been deduced from the star product between two arbitrary functions $f(x)$ and $g(x)$ in the first order of θ [19,20,35,36]:

$$f(x)*g(x) = f(x)g(x) - \frac{i}{2}\theta^{\alpha\beta}\partial_\alpha f(x)\partial_\beta g(x) \quad (2)$$

Thus, the usual product $f(x)g(x)$ replace by new star product $f(x)*g(x)$ which represented by eq (2). Actually there are many attempts to study noncommutative space time in three dimensional spaces, but limited Physics phenomena are studied in two dimensional spaces for examples (the isotropic Harmonic oscillator plus inverse quadratic, the sextic, the singular even-power, the mixed, the Inverse-Power) the study of bound-states of the Schrödinger equation for the previously potentials is fundamental to understanding the molecular spectrum of a diatomic molecule, as a good example, of a Physics interaction, in two dimensional space (2D), the pseudoharmonic potential play an important role in the history of molecular and Quantum Chemistry, furthermore, this potential used to describing molecular structures and interactions, in addition, the pseudoharmonic potential may be used for the energy spectrum of linear and non-linear systems [37]. The plan of the paper is as follows: In next section, we present and review the Boopp's shift method. In section 3 a two dimensional central potential known by pseudoharmonic potential is represented. Next, in section 4 we derive the deformed

molecular diatomic with the pseudoharmonic potential in noncommutative real space, we also obtain the deformed bound-state eigensolutions of molecular diatomic. A detailed discussion presented in section 5. Finally, the relevant conclusions are given in last section.

2. REVIEW OF BOOPP'S SHIFT METHOD

In this paper, a Boopp's shift method will be used, instead of solving the non commutative Schrödinger equation by using star product procedure, we replace new star product in the Schrödinger equation by the usual ordinary product by making a Boopp's shift and the translation of the noncommutative Schrödinger equation, respectively, [26,35,36]:

$$\hat{x}_i = x_i - \frac{\theta_{ij}}{2} p_j \text{ and } \hat{p}_i = p_i \quad (3)$$

$$\hat{H}(\hat{p}_i, \hat{x}_i) * \bar{\Psi}(\vec{r}) = E_{nc-ph} \bar{\Psi}(\vec{r}) \rightarrow H(\hat{p}_i, \hat{x}_i) \psi(\vec{r}) = E_{nc-ph} \psi(\vec{r})$$

where $\hat{H}(\hat{p}_i, \hat{x}_i)$, $\bar{\Psi}(\vec{r})$ and E_{nc-ph} , $\psi(\vec{r})$ are represents NC 2D-Hamiltonian, NC complex wave function, NC energy and ordinary complex wave function respectively. After a straightforward calculation, on can show that in NC 2D spaces:

$$\begin{aligned} \hat{x} &= x - \frac{\theta}{2} p_y, & \hat{y} &= y + \frac{\theta}{2} p_x \\ \hat{x}^2 &= x^2 - \theta x p_y, & \hat{y}^2 &= y^2 + \theta y p_x \\ \hat{p}_{x,y} &= p_{x,y} \end{aligned} \quad (4)$$

where $\theta \equiv \theta^{12}$ then, we can easily verify that, the commutators, between the operators position themselves, in NC 2D space, will be written as follows:

$$\begin{aligned} [\hat{x}, \hat{y}] &= i\theta \\ [\hat{x}, \hat{x}] &= [\hat{y}, \hat{y}] = 0 \\ [x, y] &= [x, x] = [y, y] = 0 \end{aligned} \quad (5)$$

Based to the eq. (4), we can show after a straightforward calculation, the following two-terms, which allow us to obtain the deformed potential in 4th section:

$$\begin{aligned}\hat{r}^2 &= r^2 - \theta L_z \\ \frac{1}{\hat{r}^2} &= \frac{1}{r^2} + \frac{\theta}{r^4} L_z\end{aligned}\quad (6)$$

As a direct result of eq. (4), it's clearly that, the kinetic term is invariant in NC 2D spaces:

$$\frac{\vec{p}^2}{2\mu} = \frac{\vec{\hat{p}}^2}{2\mu} \quad (7)$$

where μ is the reduced mass of the diatomic molecules.

3. THE PSEUDOHARMONIC POTENTIAL IN ORDINARY TWO DIMENSIONAL SPACES

In this section we shall review some fundamentals production of Schrödinger equation with pseudoharmonic potential while referring to [37] for more details. In particular, we shall have for previously central potential in commutative 2D space:

$$\left(-\frac{1}{2\mu} \left(r \frac{1}{r} \frac{\partial}{\partial r} \left(r \frac{\partial}{\partial r} \right) + \frac{1}{r^2} \frac{\partial^2}{\partial \varphi^2} \right) + V(r) \right) \Psi(\vec{r}) = E \Psi(\vec{r}), \quad (8)$$

with, pseudoharmonic potential:

$$V(r) = D_e \left(\frac{r}{r_e} - \frac{r_e}{r} \right)^2 \quad (9)$$

where D_e , is the dissociation energy and r_e is the equilibrium intermolecular separation. The potential (9) can be rewritten in the form of an isotropic Harmonic oscillator plus inverse quadratic potential [37]:

$$V(r) = ar^2 + \frac{b}{r^2} + c \quad (10)$$

where $a = D_e r_e^{-2}$, $b = D_e r_e^{+2}$ and $c = -2D_e$. The complete normalized wave function $\Psi(\vec{r})$ separated on radial function $R_{nl}(r)$ and a standard angular as follows:

$$\Psi(\vec{r}) = r^{-\frac{1}{2}} R_m(r) \exp(\pm im\varphi) \quad (11)$$

where $R_{nl}(r)$ satisfy the following equation:

$$\left(\frac{d^2}{dr^2} + 2\mu \left(E - ar^2 - \frac{b}{r^2} - c \right) - \frac{m^2 - \frac{1}{4}}{r^2} \right) R_m(r) = 0 \quad (12)$$

where E is the energy and m represent the orbital angular momentum Quantum number. The complete orthonormalization functions and the energy eigen values respectively in two dimensional spaces [37]:

$$\begin{aligned}\Psi_0(r, \varphi) &= N_0 \exp\left(-\frac{1}{2}\sqrt{2a\mu}r^2\right) r^{\sqrt{2b\mu}} \\ E_0 &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu}\right) \\ \Psi_2(r, \varphi) &= N_2 \exp\left(-\frac{1}{2}\sqrt{2a\mu}r^2\right) r^{\sqrt{2b\mu+4}} \exp(\pm i2\varphi) \\ E_1 &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu+1}\right) \\ \Psi_2(r, \varphi) &= N_2 \exp\left(-\frac{1}{2}\sqrt{2a\mu}r^2\right) r^{\sqrt{2b\mu+4}} \exp(\pm i2\varphi) \\ E_2 &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu+4}\right) \\ \Psi_m(r, \varphi) &= N_m \exp\left(-\frac{1}{2}\sqrt{2a\mu}r^2\right) r^{\sqrt{2b\mu+m^2}} \exp(\pm im\varphi) \\ E_m &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu+m^2}\right),\end{aligned}\quad (13)$$

where, the normalization constant N_m is given by:

$$N_m = \left[\frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+m^2}}}{\sqrt{2b\mu+m^2}!} \right]^{\frac{1}{2}} \quad (14)$$

As well know that, pseudoharmonic potential is a modified Harmonic oscillator were studied.

4. THE PSEUDOHARMONIC POTENTIAL IN NONCOMMUTATIVE TWO DIMENSIONAL REAL SPACES

If we apply the Boopp's shift method to the Schrödinger equation, on the molecular diatomic with the pseudoharmonic potential $V(\hat{r})$, we obtain:

$$\left(-\frac{1}{2\mu}\left(r\frac{1}{r}\frac{\partial}{\partial r}\left(r\frac{\partial}{\partial r}\right)+\frac{1}{r^2}\frac{\partial^2}{\partial \varphi^2}\right)+V(\hat{r})\right)\hat{\Psi}(\hat{r})=E_{NC}\hat{\Psi}(\hat{r}) \quad (15)$$

where $V(\hat{r})$ obtained from replaced operator $\hat{r}$ by the ordinary r as follows:

$$V(\hat{r})=a\hat{r}^2+b\hat{r}^{-2}+c \quad (16)$$

And E_{NC} is the energy in NC 2D space. Based on eq. (6) we can show that, the operator potential $V(\hat{r})$ take the following form:

$$V(\hat{r})=ar^2+\frac{b}{r^2}+c+V_{ph}(r) \quad (17)$$

where the supplementary term $V_{ph}(r)$ take the form:

$$V_{ph}(r)=\theta\left(\frac{b}{r^4}-a\right)L_z \quad (18)$$

Clearly, the term $V_{pot}(r)$, proportional with small parameter θ , then we considered as a perturbative terms, furthermore we can rewritten it to the equivalent form:

$$V_{ph}(r)=2\theta f(r)\vec{S}\vec{L} \quad (19)$$

where $\vec{S}=\frac{1}{2}$ and $f(r)=\frac{b}{r^4}-a$. The operator $\vec{S}\vec{L}$ traduced physically, the coupling between spin and orbital momentum, known by spin-orbit interaction. Furthermore we can rewrite it to the equivalent form:

$$V_{ph}(r)=\theta f(r)\left(\vec{J}^2-\vec{L}^2-\vec{S}^2\right) \quad (20)$$

The corresponding NC Hamiltonian H_{NC-1} will be form, as follows:

$$H_{NC-1}=-\frac{\Delta}{2\mu}+D_e\left(\frac{r}{r_e}-\frac{r_e}{r}\right)^2+\theta f(r)\left(\vec{J}^2-\vec{L}^2-\vec{S}^2\right) \quad (21)$$

It known that, in Quantum mechanics the operators $(\vec{J}^2, \vec{L}^2, \vec{S}^2$ and $J_z)$ are formed complete set of commuting observables, then $(\vec{J}^2-\vec{L}^2-\vec{S}^2)$ will be gives two eigen values: the first is $(L(j,l,s)=l(l+\frac{1}{2})(l+\frac{3}{2})-l(l+1)-\frac{3}{4})$ and

the second $(L'(j,l,s)=l(l-\frac{1}{2})(l+\frac{1}{2})-l(l+1)-\frac{3}{4})$, corresponded $j=l+\frac{1}{2}$ (spin up) and $j=l-\frac{1}{2}$ (spin down), respectively. Then, we can form a diagonal matrix of order (2×2) with elements H_{up} and H_d :

$$\begin{aligned} H_{up} &= -\frac{\Delta}{2\mu}+D_e\left(\frac{r}{r_e}-\frac{r_e}{r}\right)+\theta f(r)L(j,l,s) \text{ if } j=l+\frac{1}{2} \Rightarrow \text{spinup} \\ H_d &= -\frac{\Delta}{2\mu}+D_e\left(\frac{r}{r_e}-\frac{r_e}{r}\right)+\theta f(r)L'(j,l,s) \text{ if } j=l-\frac{1}{2} \Rightarrow \text{spindown} \end{aligned} \quad (22)$$

The modification to the energy levels, associate with spin up and spin down, E_{up} and E_d , respectively, at first order of θ , obtains by the applying the perturbation theory:

$$\begin{aligned} E_{up} &= \langle n, j, l, s | H_{up} | n, j, l, s \rangle \\ E_d &= \langle n, j, l, s | H_d | n, j, l, s \rangle \end{aligned} \quad (23)$$

In NC 2D real space, the modification of the energy levels associated with spin up, in the first order of θ correspond the m^{th} order of excited states E_m determined by using Eqs. (19), (20) and (23):

$$\frac{E_{mup}}{|N_m|^2}=\theta L(j,l,s)\int_0^{+\infty}\left[\exp\left(-\frac{1}{2}\sqrt{2a\mu}r^2\right)r^{\sqrt{2b\mu+m^2}}\right]^2\left(\frac{b}{r^4}-a\right)rdr \quad (24)$$

this can be written as follows:

$$E_{lmup} = 2\pi\theta L(j, l, s) \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+m^2}}}{\sqrt{2b\mu+m^2}!} \theta(bT_1 - aT_2) \quad (25)$$

where the two terms T_1 and T_2 are given by:

$$\begin{aligned} T_1 &= \int_0^{+\infty} \exp(-\sqrt{2a\mu}r^2) r^{2\sqrt{2b\mu+m^2}-3} dr \\ T_2 &= \int_0^{+\infty} \exp(-\sqrt{2a\mu}r^2) r^{1+2\sqrt{2b\mu+m^2}} dr \end{aligned} \quad (26)$$

and we use the following form of special integral [39]:

$$\int_0^{+\infty} x^{\nu-1} \exp(-\beta x^2 - \gamma x) dx = (2\beta)^{-\frac{\nu}{2}} \Gamma(\nu) \exp\left(\frac{\gamma^2}{8\beta}\right) D_{-\nu}\left(\frac{\gamma}{\sqrt{2\beta}}\right) \quad (27)$$

where $D_{-\nu}$, is the parabolic cylinder function, as a condition ($\text{Re}(\beta) > 0$ and $\text{Re}(\nu) > 0$), we set $\beta = \sqrt{2a\mu}$, $\gamma = 0$ and ($\nu = 2\sqrt{2b\mu+m^2} - 2$ in T_1 and $\nu = 2 + 2\sqrt{2b\mu+m^2}$ in T_2), which allow us to obtaining:

$$\begin{aligned} T_1 &= (2\sqrt{2a\mu})^{\left(\sqrt{2b\mu+m^2}-1\right)} \Gamma\left(2\left(\sqrt{2b\mu+m^2}-1\right)\right) D_{-2\left(\sqrt{2b\mu+m^2}-1\right)}(0) \\ T_2 &= (2\sqrt{2a\mu})^{\left(-\sqrt{2b\mu+m^2}+1\right)} \Gamma\left(2\left(\sqrt{2b\mu+m^2}+1\right)\right) D_{-2\left(\sqrt{2b\mu+m^2}+1\right)}(0) \end{aligned} \quad (28)$$

the above equations are simplified to the following form:

$$E_{up} = 2\pi L(j, l, s) |N_m|^2 \theta \delta, \quad (29)$$

Where α , β and δ are given by:

$$\begin{aligned} \alpha &= \sqrt{2b\mu+m^2} - 1, \beta = \left(\sqrt{2b\mu+m^2} + 1\right) \\ \delta &= \left(b(2\sqrt{2a\mu})^{-\alpha} \Gamma(2\alpha) D_{-2\alpha}(0) - a(2\sqrt{2a\mu})^{-\beta} \Gamma(2\beta) D_{-2\beta}(0)\right) |N_m|^2 \end{aligned} \quad (30)$$

Similarly, we can easily obtain the modification the energy levels, corresponding electron with spin down E_{md} in the first

order of θ , corresponding m^{th} order of excited states, as follows:

$$E_{md} = 2\pi L'(j, l, s) |N_m|^2 \theta \delta \quad (31)$$

We conclude, from Eqs. (13), (29) and (31) that, the total energy of an electron with spin up and spin down (E_{NCmup} E_{NCmd}) corresponded (H_{up} and H_d) are given by, respectively:

$$\begin{aligned} E_{NCmup} &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu+m^2}\right) + 2\pi L(j, l, s) |N_m|^2 \theta \delta \\ E_{NCmd} &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu+m^2}\right) + 2\pi L'(j, l, s) |N_m|^2 \theta \delta \end{aligned} \quad (32)$$

5. DISCUSSIONS

As a good example, of applications for obtained results, we can estimate the emission energy ΔE_{NCd} , corresponding electron with spin down, when translate from first excited states to the stationary state, as follows:

$$\Delta E_{NCd} = E_{NC1d} - E_{NC0d} \quad (33)$$

Using eq. (32), we obtain easily:

$$\frac{\Delta E_{NCd} - \Delta E}{2\pi\theta} = L'\left(\frac{3}{2}, 1, \frac{1}{2}\right) |N_1|^2 - L'\left(\frac{3}{2}, 0, \frac{1}{2}\right) |N_0|^2 \quad (34)$$

where represent the emission energy in the ordinary commutative space:

$$\Delta E = \sqrt{\frac{2a}{\mu}} \left(\sqrt{2b\mu+1^2} - \sqrt{2b\mu}\right) \quad (35)$$

The two values $L'\left(\frac{3}{2}, 1, \frac{1}{2}\right) = -2$ and $L'\left(\frac{3}{2}, 0, \frac{1}{2}\right) = -1$, then, the eq. (34) reduce to the simple form:

$$\Delta E_{NCd} = \Delta E + 2\pi \left(|N_0|^2 - 2|N_1|^2\right) \theta \delta \quad (36)$$

thus, the noncommutativity modified ΔE_{up}

by $2\pi \left(-2 \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+1}}}{\sqrt{2b\mu+1}} + \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu}}}{\sqrt{2b\mu}} \right) \theta \delta$,
it's important to calculate the absolute incertitude:

$$\frac{|\Delta E_{NCd} - \Delta E|}{\Delta E_d} = \frac{2\pi \left(\frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu}}}{\sqrt{2b\mu}} - 2 \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+1}}}{\sqrt{2b\mu+1}} \right) \theta \delta}{\sqrt{\frac{2a}{\mu}} (\sqrt{2b\mu+1}^2 - \sqrt{2b\mu}) + 2\pi (|N_0|^2 - 2|N_1|^2) \theta \delta} \quad (37)$$

after a straightforward calculation, we get the absolute incertitude:

$$\frac{|\Delta E_{NCd} - \Delta E|}{\Delta E_d} = 2\pi \left(\frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu}}}{\sqrt{2b\mu}} - 2 \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+1}}}{\sqrt{2b\mu+1}} \right) \theta \delta \ll \Delta E \quad (38)$$

we observe that the obtained result

$$\left(2\pi \left(\frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu}}}{\sqrt{2b\mu}} - 2 \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+1}}}{\sqrt{2b\mu+1}} \right) \theta \delta \right)$$

proportional to the infinitesimal parameter, then, the absolute incertitude is justified. Regarding to the eq. (32), the eigen values of energies are real, and then the NC Hamiltonian is Hermitian ($H_{NC-1} = (H_{NC-1})^+$), which can be decomposed as follows:

$$H_{NC-1} = H_{ph-com} + H_{ph-so} \quad (39)$$

where H_{ph-com} and H_{ph-so} are determined from, the following relation, respectively:

$$H_{ph-com} = \left[-\frac{\Delta}{2\mu} + D_e \left(\frac{r}{r_e} - \frac{r_e}{r} \right)^2 \right] I_{2*2} \quad (40)$$

$$H_{ph-so} = \theta f(r) \begin{pmatrix} L(j, l, s) & 0 \\ 0 & L'(j, l, s) \end{pmatrix}$$

It's clearly H_{ph-com} represent a particle of spin $(1/2)$ interacted exactly with the pseudoharmonic potential in ordinary 2D space, while the matrix H_{ph-so} represent the spin-orbit interaction.

Furthermore, if we choose, the parameter θ and the vector of a magnetic field, which oriented with (Oz.) axes, as follows [35, 36]:

$$\theta = \alpha B, \quad \vec{B} = B \vec{k} \quad (41)$$

$$\theta L_z = \alpha (\vec{J} \vec{B} - H_Z)$$

where α is real proportional constant and H_Z is the usual Zeeman operator in commutative space, Substituting Eq. (41) into the NC Hamiltonian equation (40), we find:

$$H_{NC-2} = -\frac{\Delta}{2\mu} + D_e \left(\frac{r}{r_e} - \frac{r_e}{r} \right)^2 + H_{ph-mag} \quad (42)$$

where the obtained magnetic operator H_{ph-mag} is given by:

$$H_{ph-mag} = -\alpha f(r) (H_Z - \vec{J} \vec{B}) \quad (43)$$

Physically, the eq. (43) represents the new magnetic interactions between electron with spin $(1/2)$ and an external magnetic field, including the ordinary Zeeman effect. In another hand, it's evidently to consider the infinitesimal parameter θ , the sum of 2-infinitesimal parameters θ_1 and θ_2 , then the complete NC Hamiltonian equal the sum of three fundamentals parts, the first one and the second part, are was represented from eq. (40), while the last part, it has seen from the eq. (43). Thus, the final operator of NC Hamiltonian H_{Ph-nc} , which obtain by applying the superposition principal:

$$H_{Ph-nc} = H_{ph-com} + H_{ph-so} + H_{ph-mag} \quad (44)$$

where, the 3-operators H_{ph-com} , H_{ph-com} and H_{ph-mag} are given by:

$$\begin{aligned}
H_{ph-com} &= -\frac{\Delta}{2\mu} + D_e \left(\frac{r}{r_e} - \frac{r_e}{r} \right)^2 \\
H_{ph-so} &= \alpha \left(\frac{b}{r^4} - a \right) (\vec{J}^2 - \vec{L}^2 - \vec{S}^2) \\
H_{ph-mag} &= \theta \left(\frac{b}{r^4} - a \right) (\vec{J} \vec{B} - H_z)
\end{aligned} \quad (45)$$

Regarding the eq. (33), we can deduce the complete NC Quantum spectrum of diatomic molecules with pseudoharmonic potential: $E_{NCup}(j,l,s,m)$ and $E_{NCd}(j,l,s,m)$ corresponding spin up and spin down respectively, evidently equal the sum 3 values of energies: $(E_m, E_{NC-so}, E_{NC-m})$ corresponding $(H_{ph-com}, H_{ph-so}$ and $H_{ph-mag})$, respectively, as follows:

$$\begin{aligned}
E_{NCup}(j,l,s,m) &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu + m^2} \right) + \\
&+ 2\pi L \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+m^2}}}{\sqrt{2b\mu+m^2}!} \delta(L(j,l,s)\theta_1 + m\theta_2) \\
E_{NCd}(j,l,s,m) &= c + \sqrt{\frac{2a}{\mu}} \left(1 + \sqrt{2b\mu + m^2} \right) + \\
&+ 2\pi L \frac{(2\sqrt{2a\mu})^{1+\sqrt{2b\mu+m^2}}}{\sqrt{2b\mu+m^2}!} \delta(L'(j,l,s)\theta_1 + m\theta_2)
\end{aligned} \quad (46)$$

6. CONCLUSIONS

In this paper, the Schrödinger equation for pseudoharmonic potential was solved by applying the Boopp's shift method to first order of the parameter in non commutative two dimensional spaces and ordinary stationer perturbation theory, we derived two new parts of NC Hamiltonian operator, the full modified of the energies levels corresponding electron with spin up and spin down were established. From this recent work, we concluded that, the NC Hamiltonian interpreting automatically two fundamentals Physics phenomenon: the new Zeeman magnetic effect including ordinary Zeeman effect and the spin-orbit interaction. Thus, the noncommutativity properties for pseudoharmonic potential were prolonged to

include new symmetries can be considers as a good candidate to gives new applications in another Physics fields.

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التشوه في مستويات الطاقة للكمون التوافقي الكاذب: الميكانيك الكمي اللانسبي

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الملخص:

يصف البحث الحلول الدقيقة لمعادلة شرودينغر لجزيئ ثنائي الذرة يتأثر بكمون توافقي كاذب $V(r) = D\left(\frac{r}{r_e} - \frac{r_e}{r}\right)$ ، تم حلها بتطبيق طريقة Boopp's shift في المرتبة الأولى للوسيط اللاتبادلي θ ، ونظرية الاضطرابات المستقرة. وقد وجد أن عصابات الطاقة (القيم الذاتية) كانت متحللة. كما وجد أن الهاملتونيان اللاتبادلي يصف ذاتياً جسيم فرميوني يدور نصف دورة ويتفاعل مع مجال مغناطيسي خارجي يتضمن تأثير زيمان المعتاد في الفضاء التبادلي. بالإضافة لذلك لوحظ أن الهاملتونيان اللاتبادلي بإمكانه وصف ظاهرة الدوران في مدار التفاعل تلقائياً و تم تبرير النتائج من خلال حساب الإرتياب المطلق.