# Quadratic Zeeman Effect for an s-electron Moving in a Screened Coulomb Potential 

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#### Abstract

: The s-level of a screened hydogenic atom in a uniform magnetic field of arbitrary strength B have been calculated accurately. First, the related spherical symmetric case $\frac{1}{2} \Omega^{2} r^{2}(\Omega \sim B)$ has been treated by using the linear variational method. The trial wave function is taken as linear mixture of hydogenic and 3D harmonic oscillator wave functions of s-symmetry.Second, the actual nonspherical case $\frac{1}{2} \Omega^{2}\left(x^{2}+y^{2}\right)$ is studied by adopting the correction proposed by Mustafa and Chhajlany [14]. The latter accounts for the difference between the two cases in an approximate way by utilizing the properties of the 2D and 3D harmonic oscillators. We start our study with the unsceened hydrogen case to recover their shifted $\frac{1}{N}$ results and deal then with the more involved case of the screened hydrogen atom covering wide ranges of the applied field and of the screening length.


Key words: Zeeman effect, screening, virial theorem.

## 1-Introduction:

A large number of physical problems require solving the schrodinger equation for spherically symmetric potential in order to determine the energy eigenvalues and eigenfunctions. Since only a handful of potentials is exactly solvable, in general, one has to resort to numerical techniques or approximation schemes. A typical example of such a potential the one that arise in an atomic system under the effect of an external magnetic field B. The treatment of the quadratic Zeeman effect $\left(\sim \mathrm{B}^{2}\right)$ attracts most available perturbativ techniques. These include the shifted $\frac{1}{N}$ expansion method based on logar perturbation theory ([7], [13], [14], [22-24], [19], [5]), the shifted 1-expansion technique ([15], [18], [14]) and the so called PT-symmetric pseudperturbation ([2, 3]), the semiclassical quantization method [6] and the so called $\hbar$ expansion method [4]. The eigenvalue problem for the Yukawa (or Debye-Huckel) potential fans into such a category ([20], [8]). When a uniform static magnetic field act on the screened electron, the problem becomes much more complicated even for very weak screenings [9]. For example, the application of such a field to the hydrogen atom breaks the orbital symmetry, thus destroying the angular momentum as a good constant of motion [14].Conventional perturbation treatment of this problem can handle the weak and strong field limits where the problem becomes almost separable. However, the experimentally most important situation is the one in which the magnetic and the Coulombic fields are comparable. To bridge the two limits
of the magnetic field, one has to rely on various approximations and interpolation techniques. One of such techniques is the linear variational method [23] which proves efficient in dealing with 2 D donor problems. The 3 D problem is much more complicated because the hamiltomian in this case is no longer spherically symmetric as in the 2 D . As we all know, that most numerical and perturbative techniques such as the shifted $1 / \mathrm{N}$ method as an example, require spherical symmetric potentials as a prerequisite initial condition. To overcome this difficulty reasonably well, we utilize the proposal due to [14]. This proposal enables one to consider initially a related spherical symmetric problem close to the real non-spherical one and then account for the nonspherical character in an approximate manner. In the next section, we outline the basic idea behind such strategy. In sec.3, we present our calculational procedure. In $\sec .4$, discuss our results and finally sec. 5 concludes the paper.

## 1- The method

## (i)- The Coulomb screening

Consider a positive point charge $(+Z)$ that is immersed in plasma. The Coulomb potential due to this charge, namely, $Z / r$ is screened by the electrons surrounding it so that an electron experiences the presence of this positive charge as if Z is modified by this screening into an effective charge $Z_{\text {eff }}=Z e^{-s r}$, where $s$ is a screening parameter. To see how this comes about, consider the expression [1]:
$V(r)=\frac{Z}{(2 \pi)^{3}} \int d^{3} q-\frac{4 \pi}{q^{2}(\bar{q})} e^{i \vec{q} \cdot \vec{r}}$
and that in the Thomas-Femi approximation, the dielectric constant is given by

$$
\varepsilon(\vec{q})=1+\frac{s^{2}}{q^{2}}
$$

$$
\left.\begin{array}{l}
\text { and }  \tag{2}\\
s^{2}=\frac{3 k_{F}}{\pi a_{o}}
\end{array}\right\}
$$

where
$a_{o}=\hbar^{2} \varepsilon / m e^{2}$
Is the Bohr radius. The screened potential by using (2) gives:

$$
V(r)=\frac{Z}{(2 \pi)^{3}} \int d^{3} q \frac{4 \pi}{\left(q^{2}+s^{2}\right)} e^{i \vec{q} \vec{r}}
$$

Therefore

$$
\begin{equation*}
V(r)=\frac{Z}{r} e^{-s r} \tag{3}
\end{equation*}
$$

with

$$
\begin{equation*}
s=\left(\frac{16}{3 \pi^{2}}\right)^{\frac{1}{3}}\left(\frac{r_{s}}{a_{0}}\right)^{\frac{1}{2}} K_{F}=\frac{2.95}{\left(r_{s} / a_{0}\right)^{\frac{1}{2}}} A^{0^{-1}} \tag{4}
\end{equation*}
$$

Corresponding to a screening length $\mathrm{D}=1 / \mathrm{s}$. Taking into account that for most of cases, the ratio $\mathrm{r}_{\mathrm{s}} / \mathrm{a}_{\mathrm{o}}$ is between 2 and 6 , we have that $1.2 \leq \mathrm{s} \leq 2.1$. The influence of the applied magnetic field R is a nontrivial problem and in the present article we assume that $s$ is the same for all values of $B$.

## (ii) The Hamiltonian

In the nonrelativistic scheme, the Hamiltonian for a screened hydrogen atom when a constant magnetic field $B$ is applied perpendicular to the ( $\mathrm{x}, \mathrm{y}$ ) plane can be written as

$$
\begin{equation*}
H=\frac{1}{2 m}\left(\vec{p}+\frac{e}{2} \vec{B} \times \vec{r}\right)^{2}-\frac{Z e^{2}}{r} e^{-s r} \tag{5}
\end{equation*}
$$

Using atomic units (see the appendix) H becomes:
$H=-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}+\frac{1}{2} \Omega^{2} \rho^{2}$
Where Fl is a dimensionless parameter which can be taken as a measure of the magnetic field, (in cgs) $\Omega=\mathrm{B} / 4.7 \times 109 \mathrm{G}$. It is clear that H is not a spherical symmetric because of the presence of the last term in (6) and so the orbital quantum number 1 is no longer a good quantum number, The angular momentum is not conserved as a consequence of the noncommutivity between $\hat{\mathrm{H}}$ and $\dot{\mathrm{L}}^{2}$ namely $\left[\hat{\mathrm{H}}, \dot{\mathrm{L}}^{2}\right] \neq 0$. Furthermore, the solution of the eigenvalue problem with $H$ defined by (6) cannot be obtained in a closed form or even numerically by conventional techniques, instead: we start with the related spherical symmetric counterpart, namely;

$$
\begin{equation*}
H_{s p h}=-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}+\frac{1}{2} \Omega^{2} r^{2} \tag{7}
\end{equation*}
$$

and try to solve the corresponding Schrödinger equation

$$
\begin{equation*}
H_{s p h} \phi=E^{\prime} \phi \tag{8}
\end{equation*}
$$

Even so the exact solution of (8) cannot be expressed in a closed from in terms of special functions. In the present paper we choose to use a mixed-basis variational method with trial function as linear combination of screened hydrogenic and 3D harmonic basis eigenfunction. Such a trial function will reduce to the screened hydrogen atom when $\Omega=0$ and to that of the 30 harmonic oscillator for large values of $n$. In order to obtain an estimate to the actual energy E , we use the approximate formula (Mustafa and Chhajlany (7994))

$$
\begin{equation*}
E \approx \frac{2}{3}\left(E^{\prime}-E_{\text {coul }}\right)+E_{\text {coul }} \tag{9}
\end{equation*}
$$

Equation (9) is based on the fact that the ground state energies of an isotropic pure 2D harmonic oscillator, with the potential term $\Omega^{2}\left(x^{2}+y^{2}\right) / 2$, are equal to two-thirds of the eigenvalues of an isotropic pure 3D harmonic oscillator with the potential term $\Omega^{2}\left(x^{2}+y^{2}+z^{2}\right) / 2$. Thus for the around state and for isotropic harmonic oscillator, the eigenvalues of $\Omega^{2}\left(x^{2}+y^{2}\right) / 2$ contribute twothirds of those of the spherically symmetric one, $\Omega^{2} r^{2} / 2$, to the coulomb energy eigenvalues. To derive a formula for $\mathrm{E}_{\text {coul }}$, we start from
$H \psi=E \psi$ with H given by (6) and impose the normalization condition
$\langle\psi \mid \psi\rangle=1$, to obtain
$E=\langle\psi| \frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}+\frac{1}{2} \Omega^{2}\left(x^{2}+y^{2}\right)|\psi\rangle$
Using
$\left\langle x^{2}\right\rangle=\left\langle y^{2}\right\rangle=\frac{1}{3}\left\langle r^{2}\right\rangle$
We obtain
$\left\langle\rho^{2}\right\rangle \approx \frac{2}{3}\left\langle r^{2}\right\rangle$
Using(12) into (10) we get:

$$
\begin{align*}
& E \approx\langle\psi| \frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}+\frac{2}{3}\left(\frac{1}{2} \Omega^{2} r^{2}\right)|\psi\rangle \\
& =\frac{2}{3}\langle\psi|-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}+\frac{1}{2} \Omega^{2} r^{2}|\psi\rangle+\frac{1}{3}\langle\psi|-\frac{1}{2} \nabla^{2}-\frac{z e^{-s r}}{r}|\psi\rangle \tag{13}
\end{align*}
$$

Using equation (7) and (8), we obtain:
$E \approx \frac{2}{3} E^{\prime}+\frac{1}{3}\langle\psi|-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}|\psi\rangle$
If we denote the second term by $\mathrm{E}_{\text {coul }}$, we finally obtain (9) and

$$
\begin{equation*}
E_{\text {coul }}=\langle\psi|-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}|\psi\rangle \tag{14}
\end{equation*}
$$

It is clear that (13) cannot be evaluated exactly since $\Psi$ is unknown. If $\Psi$ is replaced by $\Phi$ from (8), we arrive at an approximate formula
$E_{\text {coul }} \approx\left\langle-\frac{1}{2} \nabla^{2}-\frac{Z e^{-s r}}{r}\right\rangle_{\text {sph }}$
Where $\left\rangle_{\text {sph }}\right.$ denotes the ground state expectation value by using $\Phi$ instead of $\Psi$ In the weak field limit $(\Omega \rightarrow 0)$, it can be shown (see the appendix) that
$E_{\text {coul }}(\Omega \rightarrow 0) \approx-\frac{Z}{2}\left\langle\frac{e^{-s r}}{r}\right\rangle_{\text {sph }}$
The evaluation of $\mathrm{E}_{\text {coul }}$ in either formula require the knowledge of the screened hydrogenic eigenfunctions $\phi_{s c H}$ satisfying
$\left(-\frac{1}{2} \nabla^{2}-\frac{Z}{r} e^{-s r}\right) \phi_{s c H}=\varepsilon_{s c H} \phi_{s c H}$
Unfortunately (16) cannot be solved analytically to obtain exact $\phi_{s c H}$ can, however, be solved by numerical integration [20] This would yield $\phi_{s c H}$ in a form of tables only.

## (3)- Calculations:

If we attempt to apply the variational method using only $\phi_{s c h}$ for the screened hydrogen atom. we will obtain a good agreement with accurate results for small values of $\Omega$, but this approach fails for large $\Omega$ even if we consider a basis with many terms. Analogous situation occurs if we use pure oscillator basis for large $\Omega$, which converges very slowly for small values of $\Omega$. In order to overcome such difficulties, we use mixed basis approach to represent $\Phi$ of equation (8)
$\phi=\sum_{i} C_{i} \phi_{i}$
$\phi_{\mathrm{i}}$ either belongs to the screened hydrogen atom $\left(\phi_{i}=\phi_{s c H}\right)$ or belongs to the 30 harmonic oscillator $\left(\phi_{i}=\phi_{\text {osc }}\right)$ where $C_{i}$ are varied so as to get minimum energy. It is worth noticing that our mixed basis are not orthogonal under inner product i,e $\left\langle\phi_{i} \mid \phi_{j}\right\rangle \neq 0$. We proceed to minimize the expectation value $\langle\phi| H_{s p h}|\phi\rangle$ with the normalization condition $\sum_{\mathrm{i}} \sum_{\mathrm{j}} \mathrm{C}_{\mathrm{i}}^{*} \mathrm{C}_{\mathrm{j}} \mathrm{S}_{\mathrm{ij}}=1$ fulfilled,
After performing a variation on the basis coefficients $C_{i}$ we reduce our problem to that of solving the matrix equation

$$
\begin{equation*}
\left\lfloor\mathrm{H}_{\mathrm{ij}}-\lambda \mathrm{s}_{\mathrm{ij}}\right\rfloor\left\lfloor\mathrm{C}_{\mathrm{j}}\right\rfloor=0 \tag{18}
\end{equation*}
$$

Where $H_{i j}=\left\langle\phi_{i}\right| H_{s p h}\left|\phi_{j}\right\rangle$ and $\quad S_{\mathrm{ij}}=\left\langle\phi_{i} \mid \phi_{j}\right\rangle$ Using the definition (7) in (18) and solving the secular equation $\left|H_{i j}-\lambda S_{i j}\right|=0$ to obtain the lowest value of $\lambda$. $\mathrm{E}^{\prime} \leq \lambda$ can be obtained in terms of $\mathrm{H}_{\mathrm{ij}}$ and $\mathrm{S}_{\mathrm{ij}}$.

The advantage of this approach is two fold. First we have a lower bound for our energy $\mathrm{E}^{\prime}$. Second we obtain a relatively simple expression for the normalized eigenfunction.

In this paper we choose to work with a two term mixed varaitional basis. In order Lo compute the binding energy for the ground state, we use a two term (1S) like states, namely $\phi_{s c H}^{1 s}$ and $\phi_{\text {osc }}^{1 s}$ i.e
$\phi_{1 s}=c_{1} \boldsymbol{\phi}_{s c H}^{1 s}+c_{2} \boldsymbol{\phi}_{o s c}^{1 s}$
substituting (19) inter (18): we get

$$
\left(\begin{array}{ll}
H_{11}-\lambda & H_{12}-\lambda s_{2}  \tag{20}\\
H_{21}-\lambda s_{21} & H_{22}-\lambda
\end{array}\right)\binom{C_{1}}{C_{2}}=\binom{0}{0}
$$

Where

$$
\begin{equation*}
H_{11}=\left\langle\phi_{s c H}^{1 s}\right| H_{s p h}\left|\phi_{s c H}^{1 s}\right\rangle \tag{21}
\end{equation*}
$$

$H_{22}=\left\langle\phi_{o s c}^{1 s}\right| H_{s p h}\left|\phi_{o s c}^{1 s}\right\rangle$
$s=s_{12}=s_{21}=\left\langle\phi_{o s c}^{1 s} \mid \phi_{s c h}^{1 s}\right\rangle$

$$
\begin{equation*}
H_{21}=H_{12}=\left\langle\phi_{s c h}^{1 s}\right| H_{s p h}\left|\phi_{o s c}^{1 s}\right\rangle \tag{24}
\end{equation*}
$$

The solution of the secular equation (18) is given by

$$
\begin{equation*}
\lambda=H_{11}+H_{22}-2 S H_{12}-\sqrt{\left(H_{11}+H_{22}-2 S H_{12}\right)^{2}-4\left(1-S^{2}\right)\left(H_{11} H_{22}-H_{12}^{2}\right) 2\left(1-S^{2}\right)} \tag{25}
\end{equation*}
$$

By solving (20) for $\mathrm{C}_{\mathrm{i}}$ with the normalization $\sum\left|C_{i}\right|^{2}=1$, w get:

$$
\begin{align*}
& C_{1}=\left[1+\left(\frac{H_{11}-\lambda}{H_{12}-\lambda S}\right)^{2}-2 S\left(\frac{H_{11}-\lambda}{H_{12}-\lambda S}\right)\right]^{-1 / 2}  \tag{26}\\
& C_{2}=\left(\frac{H_{11}-\lambda}{H_{12}-\lambda S}\right)\left[1+\left(\frac{H_{11}-\lambda}{H_{12}-\lambda S}\right)^{2}-2 S\left(\frac{H_{11}-\lambda}{H_{12}-\lambda S}\right)\right]^{-1 / 2} \tag{27}
\end{align*}
$$

$\mathrm{H}_{11}, \mathrm{H}_{12}$ and S for the screened hydrogen may by evaluated numerically since $\phi_{s c H}^{1 s}$ is in the form of tables.

## (4) Results and discussion

## (i) The s=0 case (the unscreened Coulomb potential)

As a prime nary stage let us start with the bare Coulomb potential $\left(-\frac{Z}{r}\right)$ i.e with $\mathrm{s}=0$. the computation of the energy by equation (25) is greatly simplified for this special case. Using $\phi_{H}^{1 s}$ and $\phi_{o s c}^{1 s}$ given by

$$
\begin{equation*}
\phi_{H}^{1 s}=\frac{1}{\sqrt{\pi}} e^{-r}, \phi_{o s c}^{1 s}=\left(\frac{\Omega}{\pi}\right)^{3 / 4} e^{-\frac{1}{2} \Omega r^{2}} \tag{28}
\end{equation*}
$$

We get:

$$
\begin{align*}
& H_{11}=\left\langle\phi_{H}^{1 s}\right|-\frac{1}{2} \nabla^{2}-\frac{1}{r}+\frac{1}{2} \Omega^{2} r^{2}\left|\phi_{H}^{1 s}\right\rangle=-\frac{1}{2}-\frac{3}{2} \Omega^{2}  \tag{29}\\
& H_{22}=\left\langle\phi_{o s c}^{1 s}\right|-\frac{1}{2} \nabla^{2}-\frac{1}{r}+\frac{1}{2} \Omega^{2} r^{2}\left|\phi_{o s c}^{1 s}\right\rangle=-\frac{3}{2} \Omega-2 \sqrt{\frac{\Omega}{\pi}}  \tag{30}\\
& H_{12}=\frac{3}{2} \Omega S+4\left(\frac{1}{\Omega \pi}\right)^{\frac{1}{4}}\left\{1-\sqrt{\frac{2}{\Omega}} e^{\frac{1}{2 \Omega}} \int_{1 / \sqrt{2 \Omega}}^{\infty} e^{-x^{2}} d x\right\} \tag{31}
\end{align*}
$$

Where $S$ is given by:
$s=4 \sqrt{2}\left(\frac{1}{\pi \Omega^{3}}\right)^{\frac{1}{4}}\left\{\left(1+\frac{1}{\Omega}\right) e^{\frac{1}{2 \Omega}} \int_{1 / \sqrt{2 \Omega}}^{\infty} e^{-x^{2}} d x-\frac{1}{\sqrt{2 \Omega}}\right\}$
The details for getting (31) and (32) appear in the appendix. The integral in (31) and (31), as a function of the applied field, can either be
evaluated numerically or taken from tables. Using equation (25), we can evaluate an upper bound for $E_{1 s}^{\prime}\left(E_{1 s}^{\prime} \leq \lambda\right)$ and the eigenfunction $\phi_{1 s}$ by using $\mathrm{C}_{1}$ and $\mathrm{C}_{2}$ from (26) and (27) respectively. $\mathrm{E}_{\text {coul }}$ Reduces to:

$$
\begin{align*}
& E_{\text {coul }} \approx\left\langle\phi_{1 s}\right|-\frac{1}{2} \nabla^{2}-\frac{Z}{r}\left|\phi_{s}\right\rangle \\
& =-\frac{1}{2} C \frac{2}{1}-C_{1} C_{2} S-C_{2}^{2}\left(\frac{3}{2} \Omega-2 \sqrt{\frac{\Omega}{\pi}}\right) \tag{33}
\end{align*}
$$

(see the appendix).
The result (33) is suitable for intermediate $\Omega$. For very small $\Omega$, $\mathrm{E}_{\text {coul }} \approx 0.167$, according to equation (15). For very large 51 the Coulomb effect is negligible and we approximately have $E \approx \frac{3}{2} E^{\prime}$

In table (1), we list the results of our calculations for the special case $\mathrm{s}=0$ according to equations (29-33) for various values of $\Omega . \mathrm{E}_{1 \mathrm{~s}}$ is evaluated by using equation (9) and listed in the last two columns where it has been obtained by using two sets of values for $\mathrm{E}_{\text {coul }}$ as stated above. As seen, these sets are very close in the range ( $\Omega=0.1-1.0$ )because the Coulomb interaction overwhelms the magnetic, As $\Omega$ grows up, the values calculated according to equation (33) becomes more valid. In table (2) these calculations are extended to higher values of $\Omega$. For such high values, $\mathrm{E}_{\text {coul }}$ becomes negligible compared with the effect of the magnetic field and so E is approximately equals to $2 \mathrm{E}^{\prime} / 3$. It is also shown in table (2) the results of other workers namely those due to Rosen [21] and Mustafa and Chhajlani [14], It is seen that our results ate very close to those of Rosen in the low field region but Slightly differ from Mustafa and Chhajlani being lower than ours in the low field region. The tatter authors used the shifted $1 / \mathrm{N}$ method due to Imbo et.a1 [7] and as pointed out by Villalba and Pino [22] this method overestimate the energy of the ground state for low field and also in the high field. The variational solution obtained using the hydrogen basis is in good agreement with the results obtained by Martin el.al. [12]. The impressive feature of our results as a whale is that the energies have the Coulomb limit of $E_{1 s}=-0.5$ at very low field and approach the outcome of the Londau level $\mathrm{E}_{1 \mathrm{~s}}=\Omega$ in tile high field, thus describing the spectrum of the electron in a uniform large magnetic field strength.

Table: (2). $\mathrm{E}_{1 \mathrm{~s}}$ tin atomic units with $\mathrm{z}=1$ ), the energy of the ground state of an electron in (1s) state in the hydrogen atom that is subjected to an external magnetic field $\Omega$ compared the results of the references [14, 21 ]

| $\boldsymbol{\Omega}$ | Mustafa and <br> Chhajlany <br> Ref [14] | Rosen <br> Ref [21] | Present Work |
| :---: | :---: | :---: | :---: |
| .05 | -0.49754 | -0.498800 | -0.49751 |
| 0.25 | -0.44872 | -0.470300 | -0.44202 |
| 0.50 | -0.33685 | -0.331200 | -0.25039 |
| 1.00 | -0.04429 | -0.165200 | -0.12408 |
| 1.25 | 0.12078 | 0.152450 | 0.24045 |
| 2.50 | 1.03356 | 1.119600 | 1.14126 |
| 5.00 | 3.05150 | 2.623000 | 3.15018 |
| 25.0 | 21.00130 | 19.87000 | 20.62900 |
| 50.0 | 44.46250 | 42.80000 | 43.95300 |
| 500 | 483.08610 | 477.50000 | 482.23900 |
| 1000 | 976.19370 |  | 975.27000 |
| 2000 | 1966.45250 |  | 1965.29000 |
| 3000 | 2958.96550 |  | 2957.90700 |
| 5000 | 4947.10917 |  | 4945.52300 |
| 10000 | 9925.3057 |  | 9925.48000 |

Table (3) $\mathbf{E}_{1 s}$ of an electron that is moving in a screened Coulomb potential under the action of an external magnetic field of strength $\Omega$ for various screening lengths $D=1 / s$ (All energies are measured in Hartrees)

| $\Omega$ | D = 10 | $D=5$ | D = 3 | $\mathrm{D}=2$ | D=1 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 0.05 | -0.40447 | -0.32414 | -0.23383 | -0.14445 | 0.01514 |
| 0.10 | -0.39732 | -0.31663 | -0.22558 | -0.13450 | 0.05858 |
| 0.15 | -0.38609 | -0.30491 | -0.21263 | -0.118120 | 0.09291 |
| 0.20 | -0.37096 | -0.28896 | -0.19442 | -0.09322 | 0.12441 |
| 0.25 | -0.35122 | -0.26774 | -0.16910 | -0.05720 | 0.15531 |
| 0.30 | -0.32529 | -0.23935 | -0.13455 | -0.01228 | 0.18627 |
| 0.35 | -0.29127 | -0.23937 | -0.09112 | 0.03406 | 0.21750 |
| 0.40 | -0.24879 | -0.15715 | -0.04364 | 0.07600 | 0.24911 |
| 0.45 | -0.20102 | -0.10910 | 0.00173 | 0.11267 | 0.28110 |
| 0.50 | -0.18392 | -0.11974 | 0.04204 | 0.14364 | 0.31346 |
| 0.55 | -0.13793 | -0.07332 | 0.07766 | 0.17658 | 0.34619 |
| 0.60 | -0.09423 | -0.03328 | 0.11013 | 0.20664 | 0.37928 |
| 0.65 | -0.05491 | 0.00396 | 0.14087 | 0.23650 | 0.41270 |
| 0.70 | -0.01829 | 0.03367 | 0.17084 | 0.26644 | 0.44644 |
| 0.75 | 0.01661 | 0.07240 | 0.20064 | 0.29673 | 0.48049 |
| 0.80 | 0.05065 | 0.10778 | 0.23058 | 0.32740 | 0.51482 |
| 0.85 | 0.08442 | 0.14459 | 0.26084 | 0.35850 | 0.54943 |
| 0.90 | 0.11829 | 0.17881 | 0.29149 | 0.39009 | 0.58430 |
| 0.95 | 0.15249 | 0.20880 | 0.32257 | 0.42206 | 0.61941 |
| 1.00 | 0.18713 | 0.24530 | 0.35409 | 0.45440 | 0.65476 |
| 1.25 | 0.36804 | 0.42513 | 0.49289 | 0.62175 | 0.83472 |
| 2.5 | 1.23670 | 1.46963 | 1.54269 | 1.62470 | 1.82363 |
| 5.00 | 3.24622 | 3.33500 | 3.9349 | 4.02440 | 4.25427 |
| 25.0 | 21.16220 | 21.26080 | 21.3746 | 21.50500 | 21.83344 |
| 50.0 | 44.61220 | 44.70410 | 44.82000 | 44.95300 | 45.29440 |
| 500 | 483.110 | 483.202 | 483.313 | 483.455 | 483.543 |
| 1000 | 976.130 | 976.217 | 976.310 | 976.410 | 976.876 |

## (ii) The screened electron

In order to obtain $\phi_{s c h}^{1 s}$ in (19), we have to solve equation (16) numerically $\phi_{\text {scH }}^{1 s}$ is given by

$$
\begin{equation*}
\phi_{s c H}^{1 s}=\frac{1}{\sqrt{4 \pi}} \frac{1}{r} R_{1 s}(r) \tag{34}
\end{equation*}
$$

Where $\mathrm{R}_{1 \mathrm{~s}}$ satisfies the radial Schrodinger equation
$\left(-\frac{1}{2} \frac{d^{2}}{d r^{2}}-\frac{Z e^{-s r}}{r}\right) R_{1 s}=\varepsilon_{s c H}^{1 s} R_{1 s}$
Equation (35) has no analytic solution in a closed form in terms of special functions; It assumes an approximate analytic solution as a series in powers of the screening parameter, $s$ but such solution is only valid in the asymptotic region [10]. In order to compute $\mathrm{R}_{1 \mathrm{~s}}$ and $\varepsilon_{\text {sch }}^{1 s}$, Eg. (35) has to be solved numerically using for example Numerov method [11]. All integrals involving $\phi_{s c h}^{1 s}$ will be evaluated numerically accordingly. To calculate $\mathrm{E}_{1 \mathrm{~s}}$ variationally, we follow similar step using the relations $(21-27)$ to obtain the matrix elements $H_{i j}$ and $S_{i j}$, and then $\lambda$ by numerical integration. To calculate $\mathrm{E}_{\text {coul }}$ we use equation (33) with suitable replacements, in the form:

$$
E_{\text {coul }}=\varepsilon_{\text {sch }}^{1 s} C_{1}^{2}-C_{1} C_{2} S+C_{2}^{2}\left\{\frac{3}{2} \Omega-\left\langle\phi_{\text {osc }}^{1 s}\right| \frac{Z e^{-s r}}{r}+\frac{1}{2} \Omega^{2} r^{2}\left|\phi_{\text {osc }}\right\rangle\right\}
$$

where $\varepsilon_{s c h}^{1 s}$ and , S are obtained numerically and $\mathrm{C}_{\mathrm{i}}$ are calculated accordingly,

Evidently, the $\mathrm{E}_{1 \mathrm{~s}}$, depends on two, parameters. the screening length $\mathrm{D}=1 / \mathrm{s}$ and the magnetic field strength through $\Omega$. The range of the former is taken as $1 \leq \mathrm{D} \leq 10$.

In table (4). We have listed the results of our calculations for the values $s=0.1,0.2, \ldots ., 1.0$ for a wide range of $\Omega\left(0.05-10^{3}\right)$.It can be seen that the Yukawa potential shifts up the energy level as long as the screening parameters increases For s-1 the presence of the magnetic field shifts the $\mathrm{E}_{1 \mathrm{~s}}$ level away words the continuum. As s increases, the contribution of the magnetic field becomes more important and the energy eigenvalues are closer to those given by the oscillator energy expression i.e to Landau levels. Therefore a critical value for s (or $\mathrm{D} \approx 1$ ) for which it stops binding the electron in the s-states.

As a whole, the above results show that the presence of bounded states in a Yukawa hydrogenic atom strongly depends on the strength of the screening, The inclusion of the magnetic field permits one to recover the Landau energy levels. In summary, the mixed variational approach gives the most accurate numerical results even for large $\Omega$ values.

## 5- Conclusions

The Mustafa and Chhajlany proposal for calculating energies in the presence of a magnetic field on a screened hydrogenic atom is successful in treating such peculiar nonseprable mixed Hamiltonian. It permits one to start with far easier spherical symmetric Hamiltonian instead. Thus presents a direct and fairly easy way for correcting and accounting for the most crucial effect of the applied field. In this way it is easy to treat a much more realistic potential such as the Yukawa potential by using a
much simpler and straight forward approach as compared with much more involved approaches such as the shifted $1 / \mathrm{N}$ method for example. The mixed variational method with only two basis function remarkably yields highly accurate results as compared with shifted $1 / \mathrm{N}$ approach. This would encourage one to extend these calculations to excited states such tile 2 s , $2 p$, ..etc.

## Appendix

(i) Derivation of Equation (6)

By using $\hat{P}=-i \hbar \hat{\nabla}$ in Equation (5), we obtain

$$
\begin{equation*}
H=\frac{1}{2 m}\left[-i \hbar \hat{\nabla}+\frac{1}{2} e(\vec{B} \times \vec{r})\right]-\frac{Z e^{2} e^{-s r}}{r} \tag{A1}
\end{equation*}
$$

If the applied magnetic field is in the z - direction, then
$\vec{B}=B \vec{k}$
and

$$
\begin{equation*}
\vec{B} \times \vec{r}=B(-y, x, 0)=B \rho \overrightarrow{e_{\phi}} \tag{A2}
\end{equation*}
$$

Noting $\hat{\nabla} \cdot\left(\rho \overrightarrow{e_{\phi}}\right)=0$ and employing $\hat{L}_{z}=-i \hbar \frac{\partial}{\partial \varphi}$, we get:

$$
\begin{equation*}
\hat{H}=-\frac{\hbar^{2}}{2 m} \hat{\nabla^{2}}+\frac{e B}{2 m} \hat{L_{z}}+\frac{e^{2}}{8 m} B^{2} \rho^{2}-\frac{Z e^{2} e^{-s r}}{r} \tag{A4}
\end{equation*}
$$

For states with s-symmetry, $\left(l=0, m_{l}=0\right) \hat{L}_{z} \phi_{s}=0$, we get

$$
\begin{equation*}
\hat{H}=-\frac{\hbar^{2}}{2 m} \nabla^{2}+\frac{e B}{2 m} \rho^{2}-\frac{Z e^{2} e^{-s r}}{r} \tag{A5}
\end{equation*}
$$

Expressing $\hat{H}$ in atomic units (energy in $\frac{m e^{4}}{\hbar^{2}}$ and distances in Bohr units $\frac{\hbar^{2}}{m e^{2}}$ ), we get:

$$
\begin{equation*}
\hat{H}=-\frac{1}{2} \nabla^{2}+\frac{1}{2} \Omega^{2} \rho^{2}-\frac{Z e^{-s r}}{r} \tag{A6}
\end{equation*}
$$

Where $\Omega$ is measure of the strength of B and is given (in CGS) by

$$
\begin{equation*}
\Omega=\frac{B}{4.7 \times 10^{9} G} \tag{A7}
\end{equation*}
$$

(ii) Derivation of Equations $(12,15)$

The virial theorem $\left.2<-\nabla^{2} / 2\right\rangle=\langle\vec{r} \cdot \vec{\nabla} V>$
With respect to Equation $(14,17)$ yield respectively:

$$
\begin{align*}
& <\nabla^{2}+\frac{Z e^{-s r}}{r}+\Omega^{2}\left(x^{2}+y^{2}\right)>=0  \tag{A8}\\
& <\nabla^{2}+\frac{Z e^{-s r}}{r}+\Omega^{2} r^{2}>_{s p h}=0 \tag{A9}
\end{align*}
$$

Using the Hellmann - Feynman theorem on Equations (14, 17) we respectively obtain

$$
\left.\begin{array}{l}
\frac{\partial E(\Omega, z)}{\partial z}=-<\frac{e^{-s r}}{r}>  \tag{A10}\\
\frac{\partial E(\Omega, z)}{\partial \Omega}=\Omega\left\langle x^{2}+y^{2}\right\rangle
\end{array}\right\}
$$

and

$$
\left.\begin{array}{l}
\frac{\partial E^{\prime}(\Omega, z)}{\partial z}=-\left\langle\frac{e^{-s r}}{r}\right\rangle_{s p h}  \tag{A11}\\
\frac{\partial E^{\prime}}{\partial z}=\Omega\left\langle r^{2}\right\rangle_{s p h}
\end{array}\right\}
$$

Provided all wave functions are normalized.
The Euler homogeneity condition associated with Equations (A8, A9) yields:
$\left\langle\nabla^{2}\right\rangle=z\left(\frac{\partial E^{\prime}}{\partial z}\right)-\Omega\left(\frac{\partial E^{\prime}}{\partial z}\right)$
$\left\langle\nabla^{2}\right\rangle_{\text {sph }}=z\left(\frac{\partial E^{\prime}}{\partial z}\right)-\Omega\left(\frac{\partial E^{\prime}}{\partial z}\right)$
Respectively.
Using Equations $(14,17)$ we obtain

$$
\begin{equation*}
E=\frac{1}{2} z \frac{\partial E}{\partial z}+\Omega \frac{\partial E}{\partial \Omega} \tag{A14}
\end{equation*}
$$

$E=\frac{1}{2} z \frac{\partial E^{\prime}}{\partial z}+\Omega \frac{\partial E^{\prime}}{\partial \Omega}$
By substituting Equations (A14, A15) into Equation (15), we obtain
$\frac{1}{2} z \frac{\partial E}{\partial z}+\Omega \frac{\partial E}{\partial \Omega} \approx \frac{2}{3}\left(\frac{1}{2} z \frac{\partial E^{\prime}}{\partial z}+\Omega \frac{\partial E^{\prime}}{\partial \Omega}\right)+\frac{1}{3} E_{\text {coul }}$
Substituting Equations A12 and A13 into A16, we get:
$\left.\frac{1}{2} z<-\frac{e^{-s r}}{r}\right\rangle+\Omega^{2}\left\langle x^{2}+y^{2}\right\rangle \approx \frac{2}{3}\left(\frac{-z}{2}\left\langle\frac{e^{-s r}}{r}\right\rangle_{\text {sph }}+\Omega^{2}\left\langle r^{2}\right\rangle_{\text {sph }}\right)+\frac{1}{3} E_{\text {coul }}$
Consider now two limiting cases for Equation (A17).
(a) The limit $\Omega \rightarrow 0$ : Equation (A17) reads

$$
E_{\text {coul }} \approx 3 z\left\{\frac{1}{3}\left\langle\frac{e^{-s r}}{r}\right\rangle_{\text {sph }}-\frac{1}{2}\left\langle\frac{e^{-s r}}{r}\right\rangle\right\}
$$

Assuming
$\left\langle\frac{e^{-s r}}{r}\right\rangle_{s p h} \approx\left\langle\frac{e^{-s r}}{r}\right\rangle$
Provided that
$H \approx-\frac{1}{2} \nabla^{2}-\frac{z e^{-s r}}{r}$
We obtain

$$
\begin{equation*}
E_{\text {coul }} \approx-\frac{z}{2}\left\langle\frac{e^{-s r}}{r}\right\rangle_{\text {sph }} \tag{A18}
\end{equation*}
$$

(a) The limit $\Omega \rightarrow \infty$ : Equation (A17) reads

$$
\begin{equation*}
\Omega^{2}<x^{2}+y^{2}>\approx \frac{2}{3} \Omega^{2}<r^{2}>_{s p h} \tag{A19}
\end{equation*}
$$

Provided that

$$
H \approx \frac{1}{2} \nabla^{2}+\frac{1}{2} \Omega^{2} r^{2}
$$

Equation (A19) furnish a sound justification for the validity of approximate formula (15) and equation (A18) provides an approximation for $\mathrm{E}_{\text {coul }}$ at low field (iii) Evaluation of $\mathrm{S}_{12}$ and $\mathrm{H}_{12}$

$$
\begin{align*}
& S_{12}=\left\langle\phi_{H}^{1 S}\right| \phi_{o s c}^{1 S}>=4\left(\frac{\Omega^{3}}{\pi}\right)^{\frac{1}{4}} \int_{0}^{\infty} r^{2} e^{-\left(\frac{1}{2} \Omega r^{2}+r\right)} d r  \tag{A20}\\
& H_{12}=\frac{3}{2} S_{12} \Omega-4\left(\frac{\Omega^{3}}{\pi}\right)^{\frac{1}{4}} \int_{0}^{\infty} r^{2} e^{-\left(\frac{1}{2} \Omega r^{2}+r\right)} d r \tag{A21}
\end{align*}
$$

Integrating the second integral by parts once and the second integral twice, we finally obtain single integral of the form

$$
\begin{equation*}
\int_{0}^{\infty} r^{2} e^{-\left(a x^{2}+b x+c\right)} d x=\frac{1}{2} \sqrt{\frac{\pi}{a}} e^{\left(b^{2}-4 a c\right) / 2 a} \operatorname{erfc}\left(\frac{b}{2 \sqrt{a}}\right) \tag{A22}
\end{equation*}
$$

Where

$$
\begin{equation*}
\operatorname{erfc}(p)=\frac{2}{\sqrt{\pi}} \int_{p}^{\infty} e^{-x^{2}} d x \tag{A23}
\end{equation*}
$$

Using (A22), (A23) in (A20), (A21) obtain:

$$
\begin{align*}
& S_{12}=\sqrt{32}\left(\frac{1}{\pi \Omega^{3}}\right)^{\frac{1}{4}}\left\{(1+1 / \Omega) e^{\frac{1}{2 \Omega}} \int_{\frac{1}{\sqrt{2 \Omega}}}^{\infty} e^{-x^{2}}-\frac{1}{\sqrt{2 \Omega}} d x\right\}  \tag{A24}\\
& H_{12}=\frac{3}{2} \Omega S_{12}-4\left(\frac{1}{\pi \Omega^{3}}\right)^{\frac{1}{4}}\left\{1-\sqrt{\frac{2}{\Omega}} e^{\frac{1}{2 \Omega}} \int_{\frac{1}{\sqrt{2 \Omega}} e^{-x^{2}}}^{\infty}\right\} \tag{A25}
\end{align*}
$$

(iv) Derivation of Equation (46):

$$
\begin{equation*}
E_{\text {coul }}=\langle\psi| h|\psi\rangle \tag{A26}
\end{equation*}
$$

where

$$
\begin{align*}
& h=\frac{1}{2} \nabla^{2}-\frac{z}{r}  \tag{A27}\\
& \psi=c_{1} \phi_{H}^{1 s}+c_{2} \phi_{o s c}^{1 s}  \tag{A28}\\
& E_{\text {coul }} \approx \\
& \left.\left.c_{1}^{2}<\phi_{H}^{1 S}|h| \phi_{H}^{1 S}>+2 c_{1} c_{2}<\phi_{H}^{1 S}\left|h_{0}\right| \phi_{o s c}^{1 S}\right\rangle+c_{2}^{2}<\phi_{o s c}^{1 S}|h| \phi_{o s c}^{1 S}\right\rangle  \tag{A29}\\
& \\
& \\
& \quad=-\frac{1}{2} c_{1}^{2}-c_{1} c_{2} S_{12}+\frac{3}{2} c_{2}^{2} \Omega-c_{2}^{2}\left\langle\phi_{o s c}^{1 S}\right| \frac{z}{r}+\frac{1}{2} \Omega^{2} r^{2}\left|\phi_{o s c}^{1 S}\right\rangle
\end{align*}
$$

The integral in the last step maybe evaluated as follows:
$<\phi_{\text {osc }}^{1 S}\left|\frac{z}{r}+\frac{1}{2} \Omega^{2} r^{2}\right| \phi_{\text {osc }}^{1 S}>=4\left(\frac{\Omega^{3}}{\pi}\right)^{1 / 2}\left\{z \int_{0}^{\infty} r e^{-\Omega r^{2}} d r+\frac{1}{2} \Omega^{2} \int_{0}^{\infty} r^{4} e^{-\Omega r^{2}} d r\right\}$
Using the result:
$\int_{0}^{\infty} x^{m} e^{-a x^{2}} d x=\frac{\Gamma\left(\frac{m+1}{2}\right)}{2 a^{\left(\frac{m+1}{2}\right)}}$
we finally obtain

$$
\begin{equation*}
E_{\text {coul }} \approx \frac{1}{2} c_{1}^{2}-c_{1} c_{2} S_{12}+c_{2}^{2}\left(\frac{3}{2} \Omega-2 \sqrt{\frac{\Omega}{\pi}}\right. \tag{A32}
\end{equation*}
$$

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ملخص:
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