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Chandra M. Adhikari

Jonathan C. Canales

Thusitha P.W. Arthanayaka

Ulrich D. Jentschura

Missouri University of Science and Technology, ulj@mst.edu

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Article

Magic Wavelengths for $1S-nS$ and $2S-nS$ Transitions in Hydrogenlike Systems

Chandra M. Adhikari ^{1,*} , Jonathan C. Canales ^{1,2}, Thusitha P. W. Arthanayaka ² and Ulrich D. Jentschura ³ 

¹ Department of Chemistry, Physics and Materials Science, Fayetteville State University, Fayetteville, NC 28301, USA; jcanales@skidmore.edu

² Department of Physics, Skidmore College, Saratoga Springs, NY 12866, USA; twkulati@skidmore.edu

³ Department of Physics, Missouri University of Science and Technology, Rolla, MO 65409, USA; ulj@mst.edu

* Correspondence: cadhikari@uncfsu.edu

Abstract: We study the magic wavelength for two-photon $1S-nS$ transitions in a hydrogen and deuterium atom, as well as $2S-nS$ transitions, where the lower level is the metastable $2S$ state. At the magic wavelength, the dynamic Stark shifts of the ground and the excited state of the transition coincide, so that the transition frequency is independent of the intensity of the trapping laser field. Experimentally feasible magic wavelengths of transitions with small slopes in the atomic polarizabilities are determined; these are the most stable magic wavelengths against variations of the laser frequency. We provide data for the magic wavelengths for the $1S-nS$ and $2S-nS$ transitions in hydrogen and deuterium, with $n = 2, \dots, 8$. We also analyze the stability of the elimination of the ac Stark shift at the magic wavelength against tiny variations of the trapping laser frequency from the magic value.

Keywords: magic wavelength; polarizabilities; atomic transitions; optical trapping

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1. Introduction

Optical lattice-clocks and optical dipole traps are at the forefront of modern scientific research in ultracold atoms [1–4]. An oscillating (ac) electric field plays a vital role in trapping an atom by a laser. However, the ac Stark shift introduced by the trapping laser field constitutes an obstacle, because the frequency of the transition whose frequency is to be measured, gets shifted significantly by the trapping light field [5,6]. For a specific atomic reference state $|\phi\rangle$, the ac Stark shift is given as [7]:

$$\Delta E_{\text{ac}}(\phi, \omega_L) = -\frac{I_L}{2\epsilon_0 c} \alpha(\phi, \omega_L), \quad (1)$$

where I_L is the laser intensity, ϵ_0 is the permittivity of free space, c is the speed of light in a vacuum, and $\alpha(\phi, \omega_L)$ is the dynamic polarizability of the atomic state $|\phi\rangle$ as a function of the angular frequency ω_L of the incident (trapping) laser. The dynamic polarizability is generally different for the ground state of an atom in comparison to its excited states. There are, however, some distinct, so-called magic laser wavelengths, where the ac Stark shifts of the two atomic states involved in a transition become equal to each other. At a magic wavelength, the ac Stark shift does not alter the transition frequency as it cancels in the difference of the shifts of the ground and excited states [4,8–12].

This work concentrates on the $1S-nS$ and $2S-nS$ transitions in hydrogen and deuterium. Transitions between different hydrogen and deuterium energy levels, their comparison with hydrogen, and the study of the isotope effect represent topics of fundamental interest in atomic physics and precision measurements [13–17].

The International System of Units (SI), sometimes referred to as the SI mksA unit system, is used throughout the paper (base units are meter, second, kilogram, and Ampere). We organize this paper as follows: In Section 2, we first evaluate a few matrix elements useful in the calculation of the dynamic polarizability (see Refs. [18–30]). At first, angular components of the matrix elements are calculated as presented in Ref. [31], and then the radial components are evaluated. The evaluation of the radial components is done using the Sturmian decomposition of the Schrödinger Green function [19–21], and then the radial integrations are done. We take into account the summations over the discrete and continuous spectra. The continuum states significantly contribute to the Stark shift for the ground state of hydrogen [28,32]. In Section 3, magic wavelengths for $1S-nS$ and $2S-nS$ transitions are calculated for hydrogen and deuterium. In Section 4, we discuss the leading reduced-mass and other corrections to the magic wavelengths. Finally, conclusions are drawn in Section 5.

2. Dynamic Polarizability of nS States

We consider the following matrix element of the Schrödinger–Coulomb propagator [23,33,34] of the hydrogen atom, for a reference state $|\phi_n\rangle$,

$$P(\phi_n; \omega_L) = \frac{e^2}{3} \left\langle \phi_n \left| \vec{r} \left(\frac{1}{H_s - E_n + \hbar\omega_L} \right) \vec{r} \right| \phi_n \right\rangle, \quad (2)$$

where \vec{r} is the electron position operator, and $H_s = \vec{p}^2 / (2\mu) - e^2 / (4\pi\epsilon_0 r)$ is the atomic Schrödinger Hamiltonian. The bound-state energy is $E_n = -(\alpha^2 \mu c^2) / (2n^2)$. The momentum operator is denoted as \vec{p} , and the reduced mass of the system is μ . Furthermore, E_n is the energy, and, in the context of the current investigation, ω_L is the angular frequency of the (trapping-field) laser. We use the well-known expression for the Schrödinger–Coulomb bound state $\Psi_{n\ell m}(\vec{r}) = \Psi_{n\ell m}(r, \theta, \varphi) = R_{n\ell}(r) Y_{\ell m}(\theta, \varphi)$ in the coordinate representation,

$$R_{n\ell}(r) = \left[\frac{(n - \ell - 1)!}{(n + \ell)!} \right]^{1/2} \frac{2^{\ell+1}}{n^2} \frac{1}{a_0^{3/2}} \left(\frac{r}{n a_0} \right)^\ell \exp\left(-\frac{r}{n a_0}\right) L_{n-\ell-1}^{2\ell+1}\left(\frac{2r}{n a_0}\right). \quad (3)$$

Here,

$$a_0 = \frac{\hbar}{\alpha \mu c} \quad (4)$$

is a generalized Bohr radius, adapted to the reduced mass of the system. The spherical harmonic of the angular part is $Y_{\ell m}(\theta, \varphi)$,

$$Y_{\ell m}(\theta, \varphi) = \left[\frac{(2\ell + 1)(\ell - m)!}{4\pi(\ell + m)!} \right]^{1/2} P_\ell^m(\cos(\theta)) e^{im\varphi}. \quad (5)$$

Here, $L_n^\ell(x)$ is the associated Laguerre polynomial, and $P_\ell^m(\cos(\theta))$ is the associated Legendre polynomial. Also, n, ℓ and m are the principal, orbital angular and magnetic orbital quantum numbers, respectively. The Green function $G(\vec{r}_1, \vec{r}_2, \omega_L)$ of the Schrödinger–Coulomb Hamiltonian H_s fulfills the second-order differential equation:

$$(H_s - E_n + \hbar\omega_L) G(\vec{r}_1, \vec{r}_2, \omega_L) = \delta^{(3)}(\vec{r}_1 - \vec{r}_2) \quad (6)$$

and can be expressed in terms of radial Green function $g_\ell(r_1, r_2, \nu_n)$, and spherical harmonics $Y_{\ell m}(\theta_i, \varphi_i)$ as [19]:

$$G(\vec{r}_1, \vec{r}_2, \nu_n) = \sum_{\ell=0}^{\infty} \sum_{m=-\ell}^{\ell} g_\ell(r_1, r_2, \nu_n) Y_{\ell m}(\theta_1, \varphi_1) Y_{\ell m}^*(\theta_2, \varphi_2). \quad (7)$$

It is convenient to define the dimensionless energy variable $t_n = t_n(\omega_L)$, which parameterizes the laser frequency,

$$t_n \equiv t_n(\omega_L) = \left(1 + \frac{2n^2 \hbar \omega_L}{\alpha^2 \mu c^2}\right)^{-1/2}, \quad [n t_n(\omega_L)]^2 = -\frac{\alpha^2 \mu c^2}{2(E_n - \omega_L)}. \quad (8)$$

The quantity ν_n acts as a generalization of the principal quantum number n of the reference state, to non-integer ν_n , as follows,

$$\nu_n \equiv \nu_n(\omega_L) = n t_n = n t_n(\omega_L). \quad (9)$$

The low-frequency limits are $\nu_n \rightarrow n$ and $t_n \rightarrow 1$ for $\omega_L \rightarrow 0$, and the high-frequency limit is $t_n \rightarrow 0$ for $\omega_L \rightarrow \infty$. The fine-structure constant is denoted as α , and c is the speed of light. In the so-called Sturmian form, the radial Green function $g_\ell(r_1, r_2, \omega_L)$ can be written as [19–21]:

$$g_\ell(r_1, r_2, \omega_L) = \frac{2\mu}{\hbar^2} \left(\frac{2}{a_0 \nu_n}\right)^{2\ell+1} (r_1 r_2)^\ell \exp\left(-\frac{r_1 + r_2}{a_0 \nu_n}\right) \times \sum_{k=0}^{\infty} \frac{k! L_k^{2\ell+1}\left(\frac{2r_1}{a_0 \nu_n}\right) L_k^{2\ell+1}\left(\frac{2r_2}{a_0 \nu_n}\right)}{(k + 2\ell + 1)! (k + \ell + 1 - \nu_n)}, \quad (10)$$

where $\nu_n \equiv \nu_n(\omega_L)$. If the reference states is an nS state with $\ell = 0$, then one can express Equation (2) in integral form,

$$P(\phi_n; \omega_L) = \frac{e^2}{3} \int_0^\infty dr_1 r_1^3 \int_0^\infty dr_2 r_2^3 R_{n0}(r_1) g_{\ell=1}(r_1, r_2, \nu_n = n t_n) R_{n0}(r_2), \quad (11)$$

where t_n is defined in Equation (8). For a ground state hydrogen ($n = 1$), for example, the radial part of the wave function reads $R_{10}(r) = 2 a_0^{-3/2} \exp(-r/a_0)$. Substituting the radial part $R_{10}(r)$ and using the Sturmian form of the radial Green function from Equation (10) in Equation (11), one obtains:

$$P(1S; \omega_L) = \frac{64 \mu e^2}{3 \hbar^2 a_0^6 t_1^3} \int_0^\infty r_1^4 dr_1 \int_0^\infty r_2^4 dr_2 \exp\left(-\frac{r_1 + r_2}{a_0 t_1}\right) \times \exp\left(-\frac{r_1 + r_2}{a_0}\right) \sum_{k=0}^{\infty} \frac{k! L_k^3\left(\frac{2r_1}{a_0 t_1}\right) L_k^3\left(\frac{2r_2}{a_0 t_1}\right)}{(k + 3)! (k + 2 - t_1)}, \quad (12)$$

which after some algebra works out to [9,24,25,28,29]:

$$P(1S; \omega_L) = \frac{\hbar^2 e^2}{\alpha^4 \mu^3 c^4} \left[\frac{2 t_1^2 (-3 + 3 t_1 + 12 t_1^2 - 12 t_1^3 - 19 t_1^4 + 19 t_1^5 + 26 t_1^6 + 38 t_1^7)}{3 (-1 + t_1)^5 (1 + t_1)^4} - \frac{256 t_1^9}{3 (-1 + t_1)^5 (1 + t_1)^5} {}_2F_1\left(1, -t_1; 1 - t_1; \left(\frac{1 - t_1}{1 + t_1}\right)^2\right) \right]. \quad (13)$$

We take the opportunity to correct a sign error in the term multiplying the hypergeometric function in Equation (3a) of Ref. [9]. In the static limit $t_1 \rightarrow 1$ ($\omega_L \rightarrow 0$), Equation (13) yields half the the static polarizability,

$$P(1S; \omega_L \rightarrow 0) = \frac{9 e^2 \hbar^2}{4 \alpha^4 \mu^3 c^4} + \mathcal{O}(\omega_L^2). \quad (14)$$

For $t_1 \rightarrow 0$ ($\omega_L \rightarrow \infty$), the leading asymptotic terms are:

$$P(1S; \omega_L \rightarrow \infty) = \frac{3 \hbar^2 e^2}{\alpha^2 \mu^2 c^2} \frac{1}{\hbar \omega_L} - \frac{3 \hbar^2 e^2}{2\mu} \frac{1}{\hbar^2 \omega_L^2} + \mathcal{O}(\omega_L^{-3}). \tag{15}$$

Following the same procedure for 2S and 3S states, one obtains [9,25,28,29]:

$$P(2S; \omega_L) = \frac{\hbar^2 e^2}{\alpha^4 \mu^3 c^4} \left[\frac{16t_2^2}{3(t_2 - 1)^6(t_2 + 1)^4} \left(1181t_2^8 - 314t_2^7 - 16t_2^6 - 166t_2^5 + 14t_2^4 + 138t_2^3 - 48t_2^2 - 42t_2 + 21 \right) - \frac{16384t_2^9(4t_2^2 - 1)}{3(t_2^2 - 1)^6} {}_2F_1 \left(1, -2t_2; 1 - 2t_2; \frac{(1 - t_2)^2}{(1 + t_2)^2} \right) \right], \tag{16}$$

for the 2S state. For the 3S state, one has the result:

$$P_{3S}(\omega_L) = \frac{\hbar^2 e^2}{\alpha^4 \mu^3 c^4} \left[\frac{54t_3^2}{(t_3 - 1)^8(t_3 + 1)^6} \left(15538t_3^{12} - 2852t_3^{11} - 13283t_3^{10} + 2090t_3^9 + 2871t_3^8 + 40t_3^7 - 62t_3^6 - 492t_3^5 + 128t_3^4 + 236t_3^3 - 95t_3^2 - 46t_3 + 23 \right) - \frac{6912t_3^9(7t_3^2 - 3)^2(9t_3^2 - 1)}{(t_3^2 - 1)^8} {}_2F_1 \left(1, -3t_3; 1 - 3t_3; \frac{(1 - t_3)^2}{(1 + t_3)^2} \right) \right]. \tag{17}$$

The dynamical polarizability $\alpha_{\phi_n}(\omega_L)$ for a reference state ϕ_n is the sum:

$$\alpha(\phi_n; \omega_L) = P(\phi_n; \omega_L) + P(\phi_n; -\omega_L). \tag{18}$$

As a check, in the static limit, we have $\alpha(\phi_n; \omega_L \rightarrow 0) = 2P(\phi_n; \omega_L \rightarrow 0)$, so that the well-known static polarizability of the ground state:

$$\alpha(1S; \omega_L \rightarrow 0) = \frac{9 e^2 \hbar^2}{2\alpha^4 \mu^3 c^4} \tag{19}$$

is confirmed. The above formalism is adapted to take into account the reduced mass of the system, which is different for hydrogen as opposed to deuterium. If one were to carry out the calculation with the reduced mass μ of the system being replaced by the electron mass m_e , then the appropriate scaling factor for the magic angular frequency is μ/m_e , where m_e is the electron mass. The polarizability itself contains two matrix elements of the position operator, which each scale with the factor m_e/μ , while the propagator denominator scales with μ/m_e , resulting in a total scaling factor $(m_e/\mu)^3$ for the polarizability.

3. Magic Wavelength

Whenever the wavelength of an incident laser matches its magic value, the ac Stark shifts of the ground and excited states of a transition become identical. For example, in the transition $\phi_i \rightarrow \phi_f$ (initial to final states), we have according to Equation (1),

$$\Delta E_{ac}(\phi_i; \omega_L = \omega_M) = \Delta E_{ac}(\phi_f; \omega_L = \omega_M), \tag{20}$$

where the magic angular frequency ω_M and the the magic frequency ν_M are given by:

$$\omega_M = 2\pi c/\lambda_M, \quad \nu_M = c/\lambda_M, \tag{21}$$

respectively. In addition, λ_M is the magic laser wavelength. In the following, we will use the terms of magic wavelength and magic frequency interchangeably, with the conversion being understood according to Equation (21). For a chosen laser intensity, the ac Stark

shift of two levels depends on the polarizabilities of two atomic states. Thus, the magic wavelength λ_M of the $1S-nS$ transition can be determined by the condition:

$$\alpha_\Delta(\lambda_M) \equiv \alpha(\phi_f; \omega_M) - \alpha(\phi_i; \omega_M) = 0, \tag{22}$$

where $|\phi_f\rangle = |nS\rangle$, and $|\phi_i\rangle = |1S\rangle$ or $|2S\rangle$. The quantity $\nu_M = c/\lambda_M = \omega_M/(2\pi)$ gives the magic frequency in Hz (see also Tables 1 and 2).

Table 1. The most stable magic wavelengths and slopes at that magic wavelengths for $1S-nS$ and $2S-nS$ transitions for hydrogen.

Transition	ν_M [10^{14} Hz]	λ_M [nm]	ζ [Hz ($\frac{\text{cm}^2}{\text{kW}}$)]	ξ [$\frac{\text{Hz}}{\text{GHz}}$ ($\frac{\text{cm}^2}{\text{kW}}$)]	χ [$\frac{e^2 a_0^2/E_h}{\text{nm}}$]
1S–2S	5.82521	514.646	−221.58	−1.3563	−5.2186
1S–3S	2.18531	1371.85	−212.65	−20.192	−10.934
1S–4S	1.06583	2812.77	−211.59	−179.26	−23.089
1S–5S	0.60703	4938.67	−211.37	−1271.8	−53.138
1S–6S (I)	0.47495	6312.10	−211.33	−1058.6	−27.077
1S–6S (II)	0.42254	7094.95	−211.32	+2445.1	+49.501
1S–7S	0.32391	9255.47	−211.30	−5859.6	−69.707
1S–8S	0.22944	13,066.4	−211.28	−18966.	−113.21
2S–3S	2.20479	1359.73	−7063.5	−24.202	−13.340
2S–4S	1.06779	2807.60	−5909.3	−185.94	−24.039
2S–5S	0.60730	4936.47	−5719.7	−1287.3	−53.831
2S–6S (I)	0.47527	6307.82	−5686.1	−1077.7	−27.603
2S–6S (II)	0.42241	7097.28	−5675.0	+2494.1	+50.459
2S–7S	0.32397	9253.80	−5657.9	−5887.3	−70.062
2S–8S	0.22946	13,065.4	−5645.9	−19,007	−113.47

Table 2. The most stable magic wavelengths and slopes at that magic wavelengths for $1S-nS$ and $2S-nS$ transitions for deuterium.

Transition	ν_M [10^{14} Hz]	λ_M [nm]	ζ [Hz ($\frac{\text{cm}^2}{\text{kW}}$)]	ξ [$\frac{\text{Hz}}{\text{GHz}}$ ($\frac{\text{cm}^2}{\text{kW}}$)]	χ [$\frac{e^2 a_0^2/E_h}{\text{nm}}$]
1S–2S	5.82680	514.506	−221.40	−1.3549	−5.2129
1S–3S	2.18590	1371.48	−212.48	−20.171	−10.922
1S–4S	1.06612	2812.00	−211.42	−179.06	−23.065
1S–5S	0.60720	4937.32	−211.20	−1270.4	−53.080
1S–6S (I)	0.47508	6310.38	−211.16	−1057.5	−27.048
1S–6S (II)	0.42266	7093.02	−211.15	+2442.5	+49.447
1S–7S	0.32400	9252.95	−211.12	−5853.2	−69.631
1S–8S	0.22950	13,062.9	−211.11	−18946.	−113.08
2S–3S	2.20539	1359.36	−7057.7	−24.175	−13.325
2S–4S	1.06808	2806.84	−5904.5	−185.74	−24.013
2S–5S	0.60747	4935.12	−5715.1	−1285.9	−53.773
2S–6S (I)	0.47540	6306.11	−5681.4	−1076.5	−27.573
2S–6S (II)	0.42252	7095.35	−5670.4	+2491.4	+50.404
2S–7S	0.32406	9251.28	−5653.3	−5880.9	−69.985
2S–8S	0.22952	13,061.8	−5641.2	−18,986.	−113.34

The polarizabilities of the ground state and the 2S and 3S excited states of hydrogen, as a function of the laser wavelength, are shown in Figure 1. The polarizabilities are expressed in atomic units, i.e., in units of $e^2 a_0^2/E_h$. Wavelengths are presented in nanometers. Images (a) and (b) of Figure 1 present a general picture of the dynamic polarizabilities with the magic frequencies indicated. Images (b) and (d) of Figure 1 show the polarizabilities

near their intersections, i.e., near the magic wavelength. Note that the polarizabilities of the excited states vary over many orders of magnitude over the range of investigated frequencies and becomes singular at resonances, while the ground-state polarizability remains of the same order-of-magnitude as the static value as discussed in Equations (14), (18) and (19).

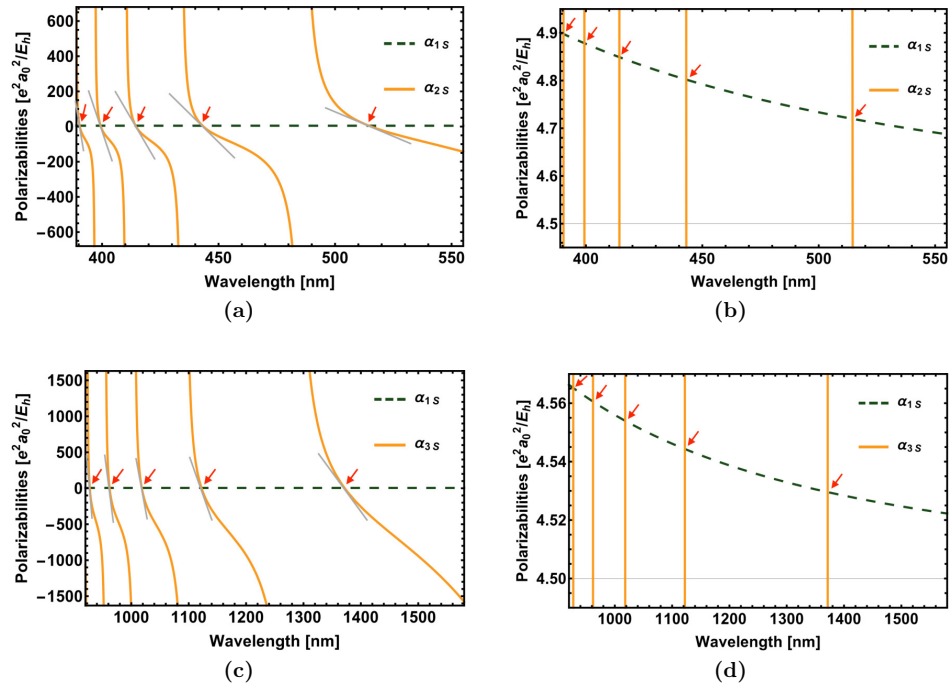


Figure 1. We investigate the polarizabilities of 1S and 2S states of hydrogen as a function of incident wavelength of the incident laser (a) and the same for 1S and 3S states of hydrogen (c). Polarizabilities near the points of intersections are shown in (b,d), with the scale of the ordinate axis decreased. Polarizabilities are expressed in atomic units, i.e., in units of $e^2 a_0^2/E_h$, where $a_0 = \hbar/(\alpha m_e c)$ is the Bohr radius (not adapted to the reduced mass of the system). We recall that the Bohr radius, adapted to the reduced mass, is $a_0 = \hbar/(\alpha \mu c)$ (see Equation (4)). Also, m_e is the electron mass, and c is the speed of light. The Hartree energy is $E_h = \alpha^2 m_e c^2$. The wavelengths corresponding to the points of intersection are the magic wavelengths.

For reference, in Tables 1 and 2, we also give the ac Stark shifts, converted to frequency units, of each individual level at the magic wavelength, normalized to the incident trapping laser intensity,

$$\zeta = \frac{1}{h} \frac{1}{I_L} \Delta E_{ac}(\phi_f, \omega_M) = \frac{1}{h} \frac{1}{I_L} \Delta E_{ac}(\phi_i, \omega_M). \quad (23)$$

It is essential to ensure the stability of the resonance frequency against small deviations of the trapping laser field from the magic wavelength. The deviation of the ac Stark shift from the zero value attained at the magic wavelength, due to a slight variation of the trapping laser frequency from the magic value, is given as:

$$\zeta = \frac{1}{h} \frac{1}{I_L} 2\pi \left. \frac{\partial}{\partial \omega_L} (\Delta E_{ac}(\phi_f, \omega_L) - \Delta E_{ac}(\phi_i, \omega_L)) \right|_{\omega_L = \omega_M} \quad (24)$$

where we refer to Equation (1). The prefactor $1/I_L$ divides out the intensity of the trapping laser, while the other prefactor $1/h$ converts the result to frequency units. Finally, the prefactor 2π converts the derivative with respect to the angular frequency to a derivative with respect to the trapping laser frequency. Our quantity ζ defined in Equation (24) differs from the quantity η defined in Equation (9) of Ref. [9] by a factor 2π . We also take the opportunity to point out that the factor $1/I_L$ was inadvertently omitted from the expression

in the middle of Equation (9) of Ref. [9] but is present in the final numerical value presented in Ref. [9]. The unit of ζ is the ac Stark shift, measured in Hz, divided by the deviation of the laser frequency from the magic value, measured in GHz, and the intensity of the exciting laser, measured in kW/cm². The unit of ζ is thus Hz/(GHz kW/cm²).

An alternative method of measuring the deviation is by calculating the change in the polarizability α_Δ , defined in Equation (22), against variations around the magic wavelength. For convenience, one can divide by the atomic unit of polarizability, which is $e^2 a_0^2 / E_h$, where E_h is the Hartree energy. Then,

$$\chi = \frac{E_h}{e^2 a_0^2} \left\{ \left. \frac{\partial \alpha_\Delta(\omega_L)}{\partial \omega_L} \right|_{\omega_L = \omega_M} \right\} \frac{\partial \omega_M}{\partial \lambda_M} = \frac{E_h}{e^2 a_0^2} \left\{ \left. \frac{\partial \alpha_\Delta(\omega_L)}{\partial \omega_L} \right|_{\omega_L = \omega_M} \right\} \left(-\frac{2\pi c}{\lambda_M^2} \right). \quad (25)$$

The unit of χ is thus the inverse nanometer, because a dimensionless quantity is differentiated with respect to a wavelength.

According to the intersection points indicated with arrows in Figure 1a, some magic wavelengths for the 2S–1S transition are close to 514 nm, 443 nm, 414 nm, 399 nm, and 390 nm. One desires a magic wavelength, in which the polarizability has the minimum slope [9]. The polarizability has the minimum slope for the magic wavelength value near 514 nm. Hence it is the most stable one against small deviations of the laser frequency from the magic value (see also Tables 1 and 2). The wavelength 514 nm lies in between the 2S–3P hydrogen transition of 656.275 nm and the 2S–4P transition of 486.132 nm [13,14,35]. This indicates that there is no loss of atoms due to resonant driving. For the 1S–3S transition in hydrogen, some magic wavelengths are near 1371 nm, 1122 nm, 1018 nm, 961 nm, and 927 nm (see the intersection points indicated with arrows in Figure 1c). The 12-3S hydrogen polarizability difference has the minimum slope value for the 1371.85 nm wavelength, and hence it is the most stable one. Wavelength 1371.85 nm lies in between the 3S–4P hydrogen transition of 1875.07 nm and the 3S–5P hydrogen transition of 1281.80 nm [13,14,35], indicating that there is no loss of atoms due to resonant driving. For the 1S–6S and 2S–6S transitions, we indicate two alternative magic wavelengths, with commensurate stability coefficients ζ and χ . The smallest magic frequency, for 1S–6S and 2S–6S, has a somewhat worse stability against deviations frequency instabilities of the trapping laser.

The most stable magic wavelengths and slopes for a various 1S–*n*S and 2S–*n*S hydrogen and deuterium transitions are listed in Tables 1 and 2. The magic wavelengths in deuterium are almost the same but slightly smaller than the reduced-mass-corrected value in the analogous transitions in hydrogen. This is because the reduced mass of deuterium is slightly larger than the reduced mass of hydrogen.

4. Relativistic and Field–Configuration Corrections to Magic Wavelengths

A discussion on the accuracy of the result we have obtained is in order. One of the most important corrections comes from the reduced mass of the system, which is in the order of 10^{−3}; this correction is taken into account by the reduced-mass adapted treatment outlined in Section 2. Another important correction is of a relativistic origin. In order to account the relativistic corrections, one should make the following substitution [23] in the \vec{r} -matrix element of the polarizability as:

$$\left\langle \phi_n \left| \vec{r} \left(\frac{1}{H_s - E_n + \hbar\omega_L} \right) \vec{r} \right| \phi_n \right\rangle \rightarrow \left\langle \phi_n + \delta\phi_n \left| \vec{r} \left(\frac{1}{H_s + \delta H - E_n - \delta E_n + \hbar\omega_L} \right) \vec{r} \right| \phi_n + \delta\phi_n \right\rangle, \quad (26)$$

where δH is the relativistic Hamiltonian correction to the Schrödinger Hamiltonian H_s , and δE_n is the corresponding relativistic energy correction. Furthermore, $|\delta\phi_n\rangle$ is the relativistic wave function correction. All of these corrections have been discussed in the context of Lamb shift calculations [23]. The relativistic correction is of relative order $\alpha^2 \sim 10^{-4}$

and thus suppressed in comparison to the reduced-mass correction [36,37]. The dynamic polarizability and, hence, the magic wavelength also receive field-configuration dependent corrections [5], which depend on the exact position of the atom in the laser field (e.g., whether it is located at a minimum or maximum of the oscillating laser electric field). The field-configuration dependent corrections are also of the order of $\alpha^2 \sim 10^{-4}$ [5]. We refrain from a discussion of these effects because the mentioned corrections are not expected to limit experimental precision, in view of the relatively small slopes of the dynamic polarizabilities near the magic wavelengths, as given by the quantities ζ and χ given in Tables 1 and 2. The same applies to the hyperpolarizabilities, which give contributions to the fourth-order ac Stark shift [38].

5. Conclusions

We studied the magic wavelengths for the $1S-nS$ and $2S-nS$ transitions of hydrogen and deuterium. We determined the magic wavelengths in these transitions by calculating the polarizabilities of the atomic states of interest and then finding wavelengths at which the polarizabilities of the respective states cancel in the difference. The polarizabilities are provided by the nS states \vec{r} -matrix elements of the Schrödinger–Coulomb propagator, as discussed in Section 2. Following the determination of the magic wavelengths and corresponding trapping laser frequencies, we calculated the slopes of the dynamic Stark shifts with respect to the frequency of the trapping laser and summarized our results in Tables 1 and 2. We find that the stability of the elimination of the ac Stark shift against tiny deviations of the trapping laser frequency from the magic value decreases with the increasing quantum number of the excited final state of the transition.

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