

## Atom lens without chromatic aberrations

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We propose a lens for atoms with reduced chromatic aberrations and calculate its focal length and spot size. In our scheme a two-level atom interacts with a near-resonant standing light wave formed by two running waves of slightly different wave vectors, and a far-detuned running wave propagating perpendicularly to the standing wave. We show that within the Raman-Nath approximation and for an adiabatically slow atom-light interaction, the phase acquired by the atom is independent of the incident atomic velocity.

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A crucial element of the toolbox for atom optics [1] is a lens to focus atom waves. Such an atom lens plays a crucial role in the realm of atom lithography [2], which is important nowadays for a multitude of technological applications. For this reason, many theoretical suggestions [3] and their realizations in experiments [4] have been made using laser fields. However, most of these realizations suffer from chromatic aberrations. In the present paper we propose a lens that is free of this type of aberration by using a special combination of light waves. Our lens is the atom-optics analog of a conventional achromatic lens [5].

We start our analysis by recalling the key features of a conventional thin lens [2,6] where a two-level atom interacts with a standing light field detuned by  $\Delta$ , giving rise to the Rabi frequency  $\Omega_0$ . This interaction creates the optical potential  $U_{\text{opt}}(x) \equiv (\hbar\Omega_0^2/\Delta) \sin^2(k_x x)$  for the motion along the  $x$  axis, which we treat quantum-mechanically. In contrast, the velocity  $v_y$  of the atom in the direction of the  $y$  axis is large and remains almost constant during the scattering process. For this reason we consider this motion classically, which allows us to set  $y \equiv v_y t$ .

Moreover, due to the small interaction time  $w_0/v_y$  determined by the waist  $w_0$  of the standing wave and the longitudinal velocity  $v_y$ , and the large detuning  $\Delta$ , we neglect the spontaneous emission, provided that

$$\Gamma w_e \left( \frac{w_0}{v_y} \right) \lesssim 1, \quad (1)$$

where  $\Gamma$  and  $w_e$  are the spontaneous emission rate and the occupation probability of the excited state, respectively. In the case of  $|\Delta| > \Omega_0$ , the maximum value of the population probability is  $w_e \sim (\Omega_0/\Delta)^2$ .

In the Raman-Nath approximation [7] for the transverse center-of-mass motion of the atom the displacement of the atom along the  $x$  axis caused by atom-field interaction is small compared to  $1/k_x$ , corresponding to

$$\omega_r \Omega_0 \frac{w_0^2}{v_y^2} \ll 1, \quad (2)$$

where  $\omega_r \equiv \hbar k_x^2/(2M)$  denotes the recoil frequency.

Within these approximations we imprint the phase

$$\phi(x) \sim -\frac{U_{\text{opt}} w_0}{\hbar v_y} = -\frac{\Omega_0^2 w_0}{\Delta v_y} \sin^2(k_x x) \quad (3)$$

onto the wave function of the center-of-mass motion of the atom in the ground state, which for  $k_x |x| \ll 1$  is a quadratic function of  $x$ , that is,

$$\phi(x) \approx -\frac{\Omega_0^2 w_0}{\Delta v_y} k_x^2 x^2. \quad (4)$$

This quadratic variation is the origin of a thin lens [6,7] with the focal length

$$\mathcal{F}_0 = \frac{M v_y^2}{2\hbar\Omega_0} \frac{\Delta}{k_x^2 w_0 \Omega_0} \equiv \kappa v_y^2 \Delta. \quad (5)$$

The spot size  $S \equiv \alpha_0 \mathcal{F}$  is determined by the focal length  $\mathcal{F}$  and the angular divergence  $\alpha_0 \equiv \delta v/v_y$  of the atomic beam, with  $\delta v$  being the uncertainty of the transverse atomic velocity. For a Gaussian wave packet of width  $\delta x$  the uncertainty  $\delta v = \hbar/(M\delta x)$  gives rise to the angular divergence  $\alpha_0 = \hbar/(M v_y \delta x)$  and the spot size

$$S_0 = \frac{v_y \Delta}{2k_x^2 \delta x w_0 \Omega_0^2}. \quad (6)$$

According to Eqs. (5) and (6) in a thin conventional lens both the focal length  $\mathcal{F}_0$  and the spot size  $S_0$  depend on the atomic velocity  $v_y$ , namely,  $\mathcal{F}_0 \propto v_y^2$  and  $S_0 \propto v_y$ , resulting in large chromatic aberrations. These scaling laws serve as our motivation to engineer a phase element for atoms and, in particular, a lens with reduced chromatic aberrations. Our suggestion relies on the interaction of a two-level atom with a near-resonant standing wave, providing us with the optical potential inducing the focusing, and a far-detuned traveling light wave removing the achromatic aberrations.

We create the lens field by the superposition

$$\mathbf{E}_s(t, \mathbf{r}) = \mathbf{E}_l(\mathbf{r}) e^{-i\omega t} (e^{i\mathbf{k}_2 \mathbf{r}} - e^{i\mathbf{k}_1 \mathbf{r}}) + \text{c.c.} \quad (7)$$

of two traveling waves of wave vectors  $\mathbf{k}_1 \equiv (k_x, k_y) \equiv (k \cos \alpha, k \sin \alpha)$  and  $\mathbf{k}_2 \equiv (-k_x, k_y)$ , which form an angle  $\alpha$  relative to the  $x$  axis shown in Fig. 1. Here  $\mathbf{E}_l(\mathbf{r})$  describes the position-dependent real-valued amplitude of the waves, which, throughout the article, is assumed to be of the form of the TEM<sub>01</sub> Hermite-Gauss mode with a node along the  $y$

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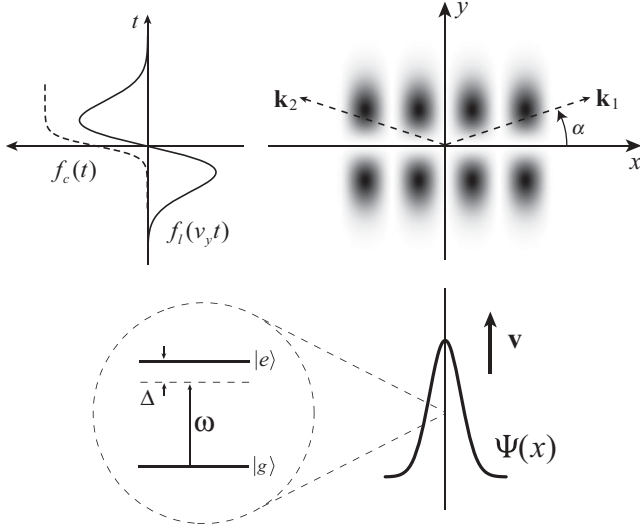


FIG. 1. Scattering of the wave packet  $\Psi = \Psi(x)$  of a two-level atom by a combination of a standing electromagnetic field formed by two propagating waves of wave vectors  $\mathbf{k}_1$  and  $\mathbf{k}_2$  and a traveling wave propagating orthogonal to the  $x$ - $y$  plane serving as a control field. As the atom propagates along the  $y$  axis with the velocity  $\mathbf{v} = v_y \mathbf{e}_y$ , the envelope of the two running waves translates according to the relation  $y = v_y t$  into the time-dependent function  $f_l = f_l(v_y t)$ . During the atom-field interaction the effective detuning of the field frequencies from the atomic transition changes its sign due to the control field with an envelope  $f_c(t)$  in the shape of a “top hat.”

axis. The frequency  $\omega$  is detuned from the frequency of the atomic transition between the ground  $|g\rangle$  and excited  $|e\rangle$  states of the corresponding energies  $E_g \equiv \hbar\omega_g$  and  $E_e \equiv \hbar\omega_e$  by an amount  $\Delta \equiv \omega - \omega_e + \omega_g$  as shown in Fig. 1.

A running control wave

$$\mathbf{E}_r(t, \mathbf{r}) = \mathbf{E}_c(x, y) e^{i(k_c z - \omega_c t)} + \text{c.c.} \quad (8)$$

with the position-dependent amplitude  $\mathbf{E}_c(x, y)$  and the shape of a top-hat propagates along the  $z$  axis perpendicular to the  $x$ - $y$  plane. The frequency  $\omega_c$  is far detuned by  $\Delta_c \equiv \omega_c - \omega_{e'} + \omega_e$  from the atomic transition between the excited state  $|e\rangle$  and some other state  $|e'\rangle$ . We suppose that the control field is weak enough to be considered perturbatively, resulting in the Stark shift  $\Delta E_e = |\tilde{\rho} \mathbf{E}_c|^2 / (\hbar \Delta_c)$  of the atomic excited state  $|e\rangle$ , where  $\tilde{\rho} \equiv \langle e | \mathbf{d} | e' \rangle$  is the dipole matrix element.

The time evolution of the state vector

$$|\Psi(t)\rangle = a_e(t; \mathbf{r}) e^{-i\omega_e t} |e\rangle + a_g(t; \mathbf{r}) e^{-i\omega_g t} |g\rangle \quad (9)$$

follows from the Schrödinger equation. Indeed, within the rotating-wave approximation the time-dependent amplitudes  $a_g$  and  $a_e$ , which depend on the position  $\mathbf{r}$  of the atom as a parameter, obey the system of equations

$$i\hbar \frac{d}{dt} \begin{pmatrix} a_e \\ a_g \end{pmatrix} = \hat{H} \begin{pmatrix} a_e \\ a_g \end{pmatrix}. \quad (10)$$

The Hamiltonian

$$\hat{H} \equiv \begin{pmatrix} \Delta E_e & V_l^* e^{-i\Delta t} \\ V_l e^{i\Delta t} & 0 \end{pmatrix} \quad (11)$$

contains the complex-valued coupling matrix elements

$$V_l(\mathbf{r}) = 2\tilde{\rho} \mathbf{E}_l(x, y) e^{-ik_y y} \sin(k_x x), \quad (12)$$

with  $\tilde{\rho} \equiv \langle g | \mathbf{d} | e \rangle$  being the dipole matrix element.

We assume that the  $x$  and  $y$  dependence of  $\mathbf{E}_l$  and  $\mathbf{E}_c$  can be separated and, since the atomic motion along the  $y$  axis is treated classically,  $y = v_y t$ , we find the forms  $\mathbf{E}_l(x, y) \equiv \mathcal{E}_l(x) f_l(y) = \mathcal{E}_l(x) f_l(v_y t)$  and  $\mathbf{E}_c(x, v_y t) = \mathcal{E}_c(x) f_c(t)$  for the electric field amplitudes. Here the envelope function

$$f_l(y) \equiv \frac{\sqrt{2}}{\pi^{1/4} w_0} \exp\left(-\frac{y^2}{2w_0^2}\right) \quad (13)$$

of the standing wave results from the TEM<sub>01</sub> Hermite-Gauss mode along the  $y$  axis and satisfies the normalization conditions

$$\int_{-\infty}^{\infty} dy f_l^2(y) = w_0. \quad (14)$$

In contrast, the envelope

$$f_c(t) \equiv \theta(t) = \begin{cases} 1, & t \geq 0, \\ 0, & t < 0, \end{cases} \quad (15)$$

of the control field has a top-hat profile, that is, a stepwise dependence as expressed by the Heaviside function  $\theta(t)$  [8].

Moreover, the detunings  $\Delta$  and  $\Delta_c$  are assumed to have the same sign and we can then set the amplitude of the Rabi frequency  $\Omega_c \equiv |\tilde{\rho} \mathcal{E}_c| / \hbar$  of the control field to  $\Omega_c = \sqrt{2\Delta\Delta_c}$ . As a result, the Stark shift  $\Delta E_e$  induced by the control light field is given by  $\Delta E_e = 2\hbar\Delta f_c^2(t)$ .

The Hamiltonian Eq. (11) finally takes the form

$$\hat{H} \cong \begin{pmatrix} 2\hbar\Delta f_c^2(t) & \hbar\Omega(x) f_l(v_y t) e^{-i(\Delta - \omega_\alpha)t} \\ \hbar\Omega(x) f_l(v_y t) e^{i(\Delta - \omega_\alpha)t} & 0 \end{pmatrix}, \quad (16)$$

where

$$\omega_\alpha \equiv k_y v_y \quad (17)$$

and

$$\Omega(x) \equiv (2|\tilde{\rho} \mathcal{E}_l| / \hbar) \sin(k_x x) \equiv \Omega_0 \sin(k_x x) \quad (18)$$

are the velocity-dependent Doppler and position-dependent Rabi frequencies, respectively.

We now solve the Schrödinger equation (10) with the Hamiltonian Eq. (16) in the case of an adiabatically slow atom-field interaction. For this purpose we substitute the second equation of the system (10) for the amplitude  $a_g$  into the first one for  $a_e$  and get the second-order differential equation

$$\frac{d^2}{dt^2} a_g - \left( i\tilde{\Delta} + \frac{1}{f_l} \frac{df_l}{dt} \right) \frac{d}{dt} a_g + \Omega^2 f_l^2 a_g = 0 \quad (19)$$

with the initial conditions

$$a_g(t_0) = 1, \quad \left. \frac{da_g}{dt} \right|_{t_0} = 0 \quad (20)$$

at time  $t_0$ . Here we have introduced the time-dependent effective detuning

$$\tilde{\Delta}(t) \equiv \Delta - \omega_\alpha - 2\Delta f_c^2(t) = \Delta - \omega_\alpha - 2\Delta\theta(t). \quad (21)$$

In the case of a slowly varying envelope  $f_l(t)$  with

$$\left| \frac{1}{f_l} \frac{df_l}{dt} \right| \ll |\tilde{\Delta}| \quad \text{or} \quad |\tilde{\Delta}| \frac{w_0}{v_y} \gg 1, \quad (22)$$

we can neglect its time derivative in the second term of Eq. (19) and arrive at the approximate equation

$$\frac{d^2}{dt^2} a_g - i \tilde{\Delta} \frac{d}{dt} a_g + \Omega^2 f_l^2 a_g \simeq 0. \quad (23)$$

For each time interval, that is, for  $-\infty < t \leq 0$  and  $0 \leq t < \infty$ , when the detuning  $\tilde{\Delta}$  is constant, the solution of Eq. (23) with the initial conditions Eq. (20) reads

$$a_g(t) = \frac{1}{2} \left[ 1 - \frac{\tilde{\Delta}(t_0)}{\lambda(t_0)} \right] e^{i\phi_+(t)} + \frac{1}{2} \left[ 1 + \frac{\tilde{\Delta}(t_0)}{\lambda(t_0)} \right] e^{i\phi_-(t)} \quad (24)$$

with

$$\phi_{\pm}(t) = \frac{1}{2} \int_{t_0}^t dt' [\pm \lambda(t') + \tilde{\Delta}(t')] \quad (25)$$

and

$$\lambda(t) \equiv \sqrt{\tilde{\Delta}^2(t) + 4\Omega^2 f_l^2(v_y t)}. \quad (26)$$

For the two time intervals  $-\infty < t \leq 0$  and  $0 \leq t < \infty$  the initial time  $t_0$  corresponds to  $t_0 = -\infty$  and  $t_0 = 0$ , respectively, where the envelope  $f_l(v_y t)$  vanishes. From Eq. (26) we find  $\lambda(t_0) = |\tilde{\Delta}(t_0)|$  and

$$\frac{1}{2} \left[ 1 \pm \frac{\tilde{\Delta}(t_0)}{\lambda(t_0)} \right] = \frac{1}{2} \left[ 1 \pm \frac{\tilde{\Delta}(t_0)}{|\tilde{\Delta}(t_0)|} \right] = \theta[\pm \tilde{\Delta}(t_0)]. \quad (27)$$

With the definitions Eqs. (21), (25), and (26) of  $\tilde{\Delta}$ ,  $\phi_{\pm}$  and  $\lambda$  together with the explicit form Eq. (27) for coefficients, we can cast Eq. (24) into the compact form  $a_g \equiv \exp(i\varphi_g)$ , where the phase

$$\varphi_g(t; x) = \frac{1}{2} \int_{-\infty}^t dt' [\tilde{\Delta} - \text{sgn}(\tilde{\Delta}) \sqrt{\tilde{\Delta}^2 + 4\Omega^2(x) f_l^2(v_y t')}] \quad (28)$$

depends on the transverse coordinate  $x$  of the atom. Since we are interesting in engineering a lens for matter waves, we can ignore the phase  $-\omega_g t$  in the Schrödinger picture, which is independent of  $x$ , and the total phase  $\Phi_g$  acquired by the atom during its interaction with the two light fields is the sum

$$\Phi_g \equiv \varphi_g(t \rightarrow \infty; x) = \varphi_g(x; \Delta - \omega_\alpha) + \varphi_g(x; -\Delta - \omega_\alpha) \quad (29)$$

of the two contributions determined by the time intervals  $-\infty < t \leq 0$  and  $0 \leq t < \infty$ , where

$$\varphi_g(x; \delta) \equiv \frac{\text{sgn}(\delta)}{2} \int_0^\infty dt [|\delta| - \sqrt{\delta^2 + 4\Omega^2(x) f_l^2(v_y t)}]. \quad (30)$$

In the case of  $\omega_\alpha \ll |\Delta|$ , the total phase  $\Phi_g$  given by Eq. (29) reduces to

$$\Phi_g(x) = \omega_\alpha \int_0^\infty dt \left[ \frac{|\Delta|}{\sqrt{\Delta^2 + 4\Omega^2(x) f_l^2(v_y t)}} - 1 \right]. \quad (31)$$

When we introduce the integration variable  $y \equiv v_y t$ , and recall the definitions Eqs. (17) and (18) of  $\omega_\alpha$  and  $\Omega$ , we arrive at

$$\Phi_g(x) = k_y \int_0^\infty dy \left\{ \left[ 1 + 4 \frac{\Omega_0^2}{\Delta^2} \sin^2(k_x x) f_l^2(y) \right]^{-1/2} - 1 \right\}. \quad (32)$$

We emphasize that  $\Phi_g$  is proportional to  $k_y$ , which is a consequence of the noncollinearity of the two wave vectors  $\mathbf{k}_1$  and  $\mathbf{k}_2$ . Moreover,  $\Phi_g$  is independent of  $v_y$ . Hence, the combination of the lens field and the control wave acting on the atom creates an achromatic phase element.

We now use this phase element to construct a lens with reduced chromatic aberrations. For this purpose we consider a position of the atom close to a node of the standing wave, which allows us to expand the square root in Eq. (32), and we arrive at

$$\Phi_g(x) \approx \frac{k_y v_y}{\Delta} \left( -\frac{\Omega_0^2 w_0}{\Delta v_y} k_x^2 x^2 \right) = \frac{k_y v_y}{\Delta} \phi(x). \quad (33)$$

Here we have recalled the normalization condition Eq. (14) for the profile function  $f_l$  and the form Eq. (4) of the phase  $\phi$  induced by a regular optical potential.

Due to the control field  $\Phi_g$  is the product of the phase  $\phi$  corresponding to a conventional optical potential and the ratio  $(k_y v_y)/\Delta$ . Hence, the focal length and the spot size of our lens read

$$\mathcal{F} = \frac{\Delta}{k_y v_y} \mathcal{F}_0 \quad \text{and} \quad \mathcal{S} = \frac{\Delta}{k_y v_y} \mathcal{S}_0. \quad (34)$$

Since according to Eqs. (5) and (6)  $\mathcal{F}_0$  and  $\mathcal{S}_0$  depend quadratically and linearly on  $v_y$ , in our lens the focal length  $\mathcal{F}$  is proportional to  $v_y$  and the spot size  $\mathcal{S}$  is independent of it. This scaling implies a reduction of the chromatic aberrations in comparison with the conventional technique of focusing atoms. Moreover, in our lens both the focal length  $\mathcal{F}$  and the spot size  $\mathcal{S}$  are larger by a factor  $|\Delta|/(k_y v_y)$  than those of the conventional lens.

The reduced chromatic aberrations are due to the symmetry of the TEM<sub>01</sub> Hermite-Gauss mode with respect to a node at  $y = 0$ . Indeed, our combination of light waves acts as two thin optical lenses covering the domains  $-\infty < y \leq 0$  and  $0 \leq y < \infty$  and contributing to the total phase  $\Phi_g$  given by Eq. (29). Depending on the sign of  $\Delta$ , the first lens is converging, whereas the second one is diverging, or vice versa. According to Eq. (5), the corresponding focal lengths

$$\mathcal{F}_{\pm} \equiv 2\kappa v_y^2 (\pm \Delta - \omega_\alpha) = 2\kappa v_y^2 (\pm \Delta - k_y v_y) \quad (35)$$

give rise with the familiar identity

$$\frac{1}{\mathcal{F}} = \frac{1}{\mathcal{F}_+} + \frac{1}{\mathcal{F}_-} \quad (36)$$

to the total focal length

$$\mathcal{F} = 2\kappa v_y^2 \frac{(\Delta - k_y v_y)(\Delta + k_y v_y)}{2k_y v_y} \approx \frac{\Delta}{k_y v_y} \mathcal{F}_0, \quad (37)$$

which coincides with Eq. (34). Here we have recalled the definition Eq. (5) and used the fact that  $k_y v_y \ll |\Delta|$ . Thus,

the suggested atom lens is designed in a similar manner as a conventional achromatic lens in optics [5].

The conditions Eqs. (1) and (2) are satisfied in experiments [2]. Indeed, for metastable helium with the velocity  $v_y = 2000$  m/s, the angle  $\alpha = 10^{-3}$ , the waist  $w_0 = 50$   $\mu\text{m}$ , the wavelength  $\lambda = 1083$  nm, the wave packet width  $\delta x = \lambda$ , the detuning  $\Delta = 2\pi \times 30$  MHz, the rate  $\Gamma = 10^7$  s $^{-1}$ , and the Rabi frequency  $\Omega_0 = |\Delta|$ , we obtain the interaction time  $2w_0/v_y = 50$  ns and the Doppler frequency  $k_y v_y = 2\pi \times 1.85$  MHz. For these values, we obtain the focal length  $\mathcal{F}_0 = 390$   $\mu\text{m}$ , the spot size  $S_0 = 3$  nm, and  $\mathcal{F}_0/\mathcal{F} = S_0/S = (k_y v_y)/\Delta = 0.06$ .

In summary we have proposed a lens with reduced chromatic aberrations. Our scheme differs from a conventional lens by the use of *two* rather than a single light field. The improvement factor is given by the ratio of the detuning and a Doppler shift.

It is interesting to note that we can interpret [9]  $\Phi_g$  as a sum of two Berry phases [10–12] acquired by the atom during the two interaction regions  $-\infty < y \leq 0$  and  $0 \leq y < \infty$ . Since the Berry phase is purely of geometrical nature, it is insensitive to small perturbations in the control parameters [13]. For this reason, we expect that our lens designed in this way is more robust against small fluctuations of the system parameters, such as the intensity of the light fields.

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