Spatial enantioseparation of gaseous chiral molecules

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We explore the spatial enantioseparation of gaseous chiral molecules for the cyclic three-level systems coupled with three electromagnetic fields. Due to molecular rotations, the specific requirements of the polarization directions of the three electromagnetic fields lead to the space-dependent part of the overall phase of the coupling strengths. Thus, the overall phase of the coupling strengths, which differs with π for the enantiomers in the cyclic three-level model of chiral molecules, varies intensely in the length scale of the typical wavelength of the applied electromagnetic fields. Under the induced gauge potentials resulting from the space-dependent part of the overall phase and the space-dependent intensities of coupling strengths, we further show spatial enantioseparation for typical parameters of gaseous chiral molecules.

I. INTRODUCTION

The left- and right-handed chiral molecules (called enantiomers) coexist in many biologically active compounds. Usually, only one form of enantiomer is biologically beneficial, while the other one is useless or even harmful. Thus, the enantioseparation of chiral molecules is fundamentally significant in organic chemistry [1], pharmacology [2–5], and biochemistry [6–8]. It also becomes an important topic in atomic, molecular, and optical physics [9–30]. Among these methods [9–30], an interesting one is based on a cyclic three-level (Δ-type) configuration [24–30], which is formed by applying three electromagnetic fields to couple with the electric dipole transitions among three levels. In general, such a cyclic three-level system is forbidden in natural atoms whose states have certain (even or odd) parities, but it exists in chiral molecules and other symmetry-broken systems [31–35]. For chiral molecules, the cyclic three-level model is special since the product of the three corresponding coupling strengths of the electric dipole transition moments changes sign with enantiomers. Then the overall phase of the product of the three corresponding coupling strengths differs by π with two enantiomers [36, 37]. This property makes the cyclic three-level model feasible in the enantioseparation [37–45] and the enantiodiscrimination [46–53] of chiral molecules.

In the cyclic three-level systems, the inner-state enantioseparation (commonly called enantio-specific state transfer) [37–43] and the spatial enantioseparation [44, 45] have been discussed theoretically. The corresponding experiments of inner-state enantioseparation have also been investigated in experiments [54, 55]. For inner-state enantioseparation, a chiral mixture can be enantiopurified in one of the three inner states (i.e., with only one enantiomer occupying that state) [37–39]. The enantiopure molecules in that state can be further spatially separated from the initial chiral mixture by a variety of energy-dependent processes. For spatial enantioseparation [44, 45], the different overall phases of the coupling strengths for two enantiomers lead to the chirality-dependent induced gauge potentials for molecules. By these induced gauge potentials, a racemic molecular beam is split into subbeams, depending on the molecular chirality or inner state.

For real molecules, especially the gaseous chiral molecules used in recent experiments [55] for the cyclic three-level systems, the molecular rotations should be considered and may introduce some negative effects on inner-state or spatial enantioseparation in the original schemes [36–45]. Due to the magnetic degeneracy of the molecular rotational states, the ideal single-loop cyclic three-level system [36–45] is generally replaced by a complicated multiple-loop one [56]. This reduces the ability of inner-state enantioseparation and spatial enantioseparation of chiral molecules. Fortunately, for the asymmetric-top chiral molecules, the single-loop configuration is constructed by applying three electromagnetic fields with appropriate polarization vectors and frequencies [57, 58].

In this paper, for the single-loop cyclic three-level system, we discuss the spatial enantioseparation of asymmetric-top chiral molecules taking into consideration molecular rotations. In order to construct the single-loop cyclic configuration, the propagation directions of three electromagnetic fields may not be parallel. Then, under the three-photon resonance condition, molecules in different spatial positions (r) experience different space-dependent parts (δ⃗r = δ⃗k · ⃗r [59]) of the overall phases of the coupling strengths, which results in the problem of phase mismatching [60] in the inner-state enantioseparation [37–43, 54, 55] and the enantiodiscrimination [61–67]. We find that this space-dependent part of the overall phase (besides the spatial distribution of the intensity of the three coupling strengths) offers an additional resource that results in induced gauge potentials. In contrast, in the original research of the spatial enantioseparation of cyclic three-level systems [44, 45] (as well as other research of the inner-state enantioseparation [37–43]), the molecular rotations were not considered. Then the propagation directions of three electromagnetic fields are designed to be parallel for simplicity. Under the three-photon resonance condition, the
space-dependent part \( \phi_r \) of the overall phase of the coupling strengths is zero. So, in this case \([44,45]\), the induced gauge potentials originate only from the spatial distribution of the intensity of three coupling strengths.

Specifically, we pay attention to the induced gauge potentials in two special directions, i.e., in the \( \delta \vec{k} \) direction (called the \( \hat{q} \) direction) and perpendicular to the \( \delta \vec{k} \) direction (called the \( \hat{p} \) direction). We make the transverse profile of each electromagnetic field have the Gaussian form. The radii of the Gaussian beams are chosen to be much larger than the wavelength of the electromagnetic field. Under the above condition, the induced gauge potentials in the \( \hat{q} \) direction exhibit the slowly varying chirality-independent envelope multiplied by a quickly varying chirality-dependent quasi periodic function. The induced gauge potentials in the \( \hat{p} \) direction are chirality dependent and slowly varying, similar to the previous case without considering the molecular rotations \([44]\).

Finally, we explore the center-of-mass motion of chiral molecules under such induced gauge potentials for two cases with initial velocity in the \( \hat{q} \) and \( \hat{p} \) directions, respectively. In a short time so that the molecules move only the distance of a typical wavelength scale, we find that the effect of the spatial enantioseparation in the former case is much better than the effect in the latter case. In a long time in which the molecules move a distance much longer than a typical wavelength, the effect of the spatial enantioseparation is comparable in the two cases that the initial velocity in the \( \hat{q} \) and \( \hat{p} \) directions.

\[ \phi_r = \delta \vec{k} \cdot \vec{r} \equiv \left( \vec{k}_{13} - \vec{k}_{23} - \vec{k}_{12} \right) \cdot \vec{r}, \]

where \( \delta \vec{k} = \vec{k}_{13} - \vec{k}_{23} - \vec{k}_{12} \) are the coupling strengths with \( \Omega_{jl} \exp(i\omega_{jl}t - i\vec{k}_{jl} \cdot \vec{r}) \) are the coupling strengths with \( \Omega_{jl} = \vec{\mu}_{jl} \cdot \vec{E}_{jl} \). Here \( \vec{\mu}_{jl} \) are the electrical dipole matrix elements and \( \vec{E}_{jl} \) are the electromagnetic fields coupling with the transition \( |j \rangle \leftrightarrow |l \rangle \).

The chirality dependence of the molecule is reflected as \([36,37]\)

\[ \Omega^L_{12} = -\Omega^R_{12}, \quad \Omega^L_{13} = \Omega^R_{13}, \quad \Omega^L_{23} = \Omega^R_{23}. \]

Here we have added the superscript \( L \) or \( R \) to denote the left-handed or right-handed chiral molecule. When referring to left-handed or right-handed chiral molecules, we add the superscript. When there is no superscript, we are referring to general molecules. It is clear from Eq. (4) that the overall phase of the product of the three coupling strengths for two enantiomers at the same position has a difference of \( \pi \) \([37]\).

Such a chirality dependence offers the possibility to realize the enantioseparation \([37-45,54-58]\), the enantiomutation \([46-53,61-67]\), and the enantiocconversion \([68-72]\). Besides the chirality-dependent part of the overall phase, the overall phase usually contains the space-dependent part \( \phi_r \) of the overall phase and the time-dependent part \( \phi_t \) of the overall phase:

\[ \phi_r = \delta \vec{k} \cdot \vec{r}, \]

\[ \phi_t = (\omega_{12} + \omega_{23} - \omega_{13}) t. \]

Usually, we take the three-photon resonance condition

\[ \omega_{12} + \omega_{23} - \omega_{13} = 0. \]

Then, the time-dependent part \( \phi_t \) of the overall phase is zero.

In the original investigations of the (inner-state or spatial) enantioseparation \([37-45]\), the molecular rotations are not considered, so the wave vectors of the three electromagnetic fields (\( \vec{k}_{12}, \vec{k}_{23}, \) and \( \vec{k}_{13} \)) are parallel to each other. Then, the phase matching is satisfied (i.e., \( \delta \vec{k} = \vec{k}_{13} - \vec{k}_{23} - \vec{k}_{12} = 0 \)), and the corresponding space-dependent part \( \phi_r \) of the overall phase is zero. However, for gaseous chiral molecules, the molecular rotations should be considered.

Now we consider the (single-loop) cyclic three-level mod-
els of gaseous chiral molecules by considering the rotational degrees of freedom, following the methods in Refs. [55, 56]. We consider different rotational states $|J_{\tau M}\rangle$ with the same vibrational ground state $|v_0\rangle$. Here $J$ is the angular momentum quantum number, $M$ is the magnetic quantum number, and $\tau$ runs from $-J$ to $J$ in unit steps in the order of increasing energy [73]. We omit $|v_0\rangle$ in further discussions. We choose the working states as $|1\rangle = |0_{10}\rangle$, $|2\rangle = |1_{-1}\rangle$, and $|3\rangle = (|1_{11}\rangle + |1_{-1}\rangle)/\sqrt{2}$ (see Fig. 1). Correspondingly, these three electromagnetic fields are $\hat{\vec{k}}_1$, $\hat{\vec{k}}_2$, and $\hat{\vec{k}}_3$ of the three electromagnetic fields may not be parallel. For simplicity, we make $\hat{\vec{k}}_1 = |\hat{\vec{k}}_1| \hat{\vec{x}}$ propagate in the $\hat{\vec{x}}$ direction and $\hat{\vec{k}}_2 = |\hat{\vec{k}}_2| \hat{\vec{z}}$ and $\hat{\vec{k}}_3 = |\hat{\vec{k}}_3| \hat{\vec{z}}$ propagate in the $\hat{\vec{z}}$ direction (see the inset of Fig. 1).

Such a setup of the electromagnetic fields makes $\delta \hat{\vec{k}} \neq 0$, which means that the space-dependent part $\phi_r$ of the overall phase is nonzero. This introduces the problem of phase mismatching, which is the main limitation of the inner-state enantioseparation [54] and the enantiodiscrimination [61–67]. In the following we discuss how the nonzero space-dependent part $\phi_r$ of the overall phase affects the spatial enantioseparation of gaseous chiral molecules.

III. INDUCED GAUGE POTENTIALS: QUASI-PERIODIC STRUCTURE

In the preceding section we showed our cyclic three-level model taking into consideration molecular rotations. In this section, we show the induced gauge potentials of such a model.

To this end, we assume the conditions of one-photon resonance $(\omega_2 - \omega_1) - \omega_{12} = 0$ and large detuning as well as weak coupling $|\Delta| \gg |\Omega_{12}| \sim |\Omega_{23}| \gg |\Omega_{12}|$ with $\Delta \equiv (\omega_3 - \omega_1) - \omega_{13} = (\omega_3 - \omega_2) - \omega_{23}$. By adiabatically eliminating the excited state $|3\rangle$ in the large detuning condition [44], we have the effective inner Hamiltonian

$$H'_{\text{inn}} = \hbar \Lambda_1 |1\rangle \langle 1| + \hbar \Lambda_2 |2\rangle \langle 2| + \hbar (g e^{i\phi} |1\rangle \langle 2| + \text{H.c.}) \quad (7)$$

with

$$g e^{i\phi} \equiv e^{-i|\hat{\vec{k}}_1|z} \left( \Omega_{12} e^{i\phi_r} - \frac{\Omega_{13}\Omega_{23}^*}{\Delta} \right). \quad (8)$$

Here $g$ is a positive real constant and $\Phi$ is a real constant. They are both uniquely determined by the parameters on the right-hand side of Eq. (8). Actually, they will depend on the chirality of the (left- or right-handed) chiral molecules (with $\Omega_{jl} \rightarrow \Omega_{jl}^L$). Then we have the inner dressed states [44]

$$|\chi_1\rangle = \cos \theta |1\rangle + e^{i\phi} \sin \theta |2\rangle,$$
$$|\chi_2\rangle = -\sin \theta |1\rangle + e^{-i\phi} \cos \theta |2\rangle,$$

with the corresponding eigenvalues $\varepsilon_1 = h(\Lambda_1 + g \tan \theta)$ and $\varepsilon_2 = h(\Lambda_2 - g \tan \theta)$, respectively. Here $\Lambda_1 = -|\Omega_{12}|^2/\Delta$, $\Lambda_2 = -|\Omega_{23}|^2/\Delta$, and $\theta$ is given by $\tan 2\theta = 2g/(\Lambda_1 - \Lambda_2)$. The parameters $\theta$ and $\Phi$ are related to the intensities and space-dependent part $\phi_r$ of the overall phase of the coupling strengths. Meanwhile, they are both chirality dependent and space dependent.

In the inner dressed states basis $\{ |\chi_1\rangle, |\chi_2\rangle \}$, we assume the adiabatic approximation [44, 45]. Then the two-level system reduces to two subsystems $H' = \sum_{\alpha=1}^2 H_{\alpha}|\chi_{\alpha}\rangle \langle \chi_{\alpha}|$ with [74–77]

$$H_{\alpha} = \frac{1}{2m}(-i\hbar \nabla - \vec{A}_{\alpha})^2 + V_{\alpha}. \quad (10)$$

Their corresponding induced vector potentials $\vec{A}_{\alpha}$ and the induced scalar potentials $V_{\alpha}$ are

$$\vec{A}_{\alpha} = i\hbar |\chi_{\alpha}| \nabla \chi_{\alpha},$$
$$V_{\alpha} = \varepsilon_{\alpha}, \quad (\alpha = 1, 2). \quad (11)$$

For real $\Omega_{12}$, $\Omega_{23}$, and $\Omega_{13}$, the induced scalar potentials are $V_1 = -R + W$ and $V_2 = -R - W$, with $R = (\Omega_{12}^2 + \Omega_{23}^2)/2\Delta$ and $W = (\Omega_{12}^2 - R^2 - 2 \cos \phi_{12}\Omega_{12}\Omega_{23}\Omega_{13}/\Delta)^1/2$. It is clear that the induced vector and scalar potentials depend on both the chirality and inner dressed states [44, 45] and change with the spatial position. We note that the subsequent results based on the Hamiltonian (10) with U(1) gauge fields are gauge invariant [74, 78].

It is observed from Eqs. (8-11) that the nonzero space-dependent part $\phi_r$ of the overall phase also contributes to the induced gauge potentials. This is very different from the original research [44, 45] without considering the molecular rotations, where the induced gauge potentials are only attributed to the spatial distribution of the intensities of the three coupling strengths.

In order to show how the space-dependent part $\phi_r$ of the overall phase affects the induced gauge potentials, we choose a new coordinate with $\hat{\vec{q}} = (\hat{\vec{z}} - \hat{\vec{x}})/\sqrt{2}$ and $\hat{\vec{p}} = (\hat{\vec{z}} + \hat{\vec{x}})/\sqrt{2}$. Since $\delta \hat{\vec{k}} = |\hat{\vec{k}}_1|(\hat{\vec{z}} - \hat{\vec{x}}) = \sqrt{2}\hat{\vec{k}}_1|\hat{\vec{q}}$ is in the $\hat{\vec{q}}$ direction, the space-dependent part $\phi_r$ of the overall phase only varies in the $\hat{\vec{q}}$ direction.

We take the 1,2-propanediol as an example as follows. The corresponding bare transition frequencies of the related three-level model are given by $\omega_{21}/2\pi = 6431.06\text{ MHz}$, $\omega_{32}/2\pi = 5781.09\text{ MHz}$, and $\omega_{31}/2\pi = 12212.15\text{ MHz}$ [79], respectively, where $\omega_{jl} = |\omega_{j} - \omega_{l}|$. The intensities $\Omega_{jl}$ of the coupling strengths are assumed to have the Gaussian forms

$$\Omega_{12} = \tilde{\Omega}_{12} e^{-(\omega_{21} - \omega_{12})^2/\sigma_{12}^2},$$
$$\Omega_{13} = \tilde{\Omega}_{13} e^{-(\omega_{13} - \omega_{12})^2/\sigma_{13}^2},$$
$$\Omega_{23} = \tilde{\Omega}_{23} e^{-(\omega_{23} - \omega_{12})^2/\sigma_{23}^2}. \quad (12)$$

Here $\tilde{\Omega}_{jl}$ are assumed to be positive real constants for simplicity. Considering the limitation of large detuning and weak coupling, we choose $\Omega_{12}/2\pi = 1\text{ MHz}$, $\Omega_{23}/2\pi = 10\text{ MHz}$, $\Omega_{13}/2\pi = 10\text{ MHz}$, and $\Delta/2\pi = 100\text{ MHz}$. In Eq. (12), $\sigma_{jl}$ represents the beam radius of the electromagnetic field. We assume that the beam radii of all the electromagnetic waves are equal for simplicity and choose $\sigma_{jl} = \sigma = 0.54\text{ m}$. In
we show the motion of two enantiomers in the quasi-periodic structure of chirality dependent and differs by a half-integer period. In order to discuss the difference in the quasi periodic function of two enantiomers, Figs. 2(a) and 2(b) are enlarged and shown together. (d) Induced scalar potentials $V_\alpha$ at fixed $q = 0$ (expressed as $V_\alpha|_{q=0}$) corresponding to two enantiomers. Here $V_\alpha|_{q=0}$, which is not affected by the space-dependent part $\phi_\alpha$ of the overall phase, presents chirality-dependent features similar to the case in Ref. [44], where the molecular rotations were not considered. Comparing Fig. 2(c) with Fig. 2(d) in a typical wavelength range (about the order of 0.01 m) of electromagnetic fields, we find that, while $V_\alpha|_{q=0}$ in Fig. 2(d) is almost unchanged in the $p$ direction, $V_\alpha|_{p=0}$ in Fig. 2(c) varies quickly in the $q$ direction. This affects the center-of-mass motion of chiral molecules.

We see from Eq. (11) that the induced vector potentials in the system under consideration are $\vec{A}_\alpha = A_{\alpha,q}(q,p)\hat{q} + A_{\alpha,p}(p,q)\hat{p}$. We obtain the effective magnetic field $\vec{B}_\alpha = \nabla \times \vec{A}_\alpha$, which has only one component $B_{\alpha,y}(q,p)$ at fixed $p = 0$ presents chirality-dependent quasi periodic features with the chirality-independent envelope, while $B_{\alpha,y}|_{q=0}$ [i.e., $B_{\alpha,y}(q,p)$ at fixed $q = 0$] presents chirality-dependent features very similar to those of the case in Ref. [44]. The feature of $B_{\alpha,y}$ is similar to that of $V_\alpha$, but its effect on the center-of-mass motion of molecules is much smaller than that of $V_\alpha$ under the typical parameters considered here. In the following we show the center-of-mass motion of the chiral molecule governed by the above-mentioned induced gauge potentials.

**IV. SPATIAL ENANTIOSEPARATION**

So far, we have shown how the space-dependent part $\phi_\alpha$ of the overall phase affects the induced gauge potentials. In this section we explore the center-of-mass motion of chiral molecules. Due to the large molecular mass and weak effective gauge potentials, the molecular propagation trajectory is treated classically [44]. Then the center-of-mass motion is governed by [44]

$$\dot{q} = v_q - \frac{A_{\alpha,q}}{m},$$

$$\dot{p} = v_p - \frac{A_{\alpha,p}}{m},$$

(13)

$$v_q = \frac{1}{m} \left[ (v_q - \frac{A_{\alpha,q}}{m}) \partial_q A_q + \left( v_p - \frac{A_{\alpha,p}}{m} \right) \partial_p A_p - \partial_q V \right],$$

$$v_p = \frac{1}{m} \left[ (v_q - \frac{A_{\alpha,q}}{m}) \partial_p A_q + \left( v_p - \frac{A_{\alpha,p}}{m} \right) \partial_p A_p - \partial_p V \right].$$

Here $v_q$ and $v_p$ represent the components of the molecular velocity $v$ in the $\hat{q}$ and $\hat{p}$ directions, respectively. Since the equations of the motion for different inner dressed states are independent, we omit the subscript $\alpha$ for $q_\alpha$, $p_\alpha$, $v_q$, $\alpha$, $v_p$, $A_{\alpha,q}$, $A_{\alpha,p}$, and $V_\alpha$. The molecules are initially placed at $(q(0), p(0)) = (0, 0)$ with the initial speed $|\vec{v}(0)| = 0.1$ m/s.

In Fig. 3 we show the motion of two enantiomers in the time interval $t \in [0, 0.1]$ s, where the chiral molecules move in the distance of a typical wavelength scale. We are interested in two cases with the initial velocity along $\hat{q}$ direction [shown in Figs. 3(a1) and 3(a2)] and in the $\hat{p}$ direction [shown

**Figure 2:** Induced scalar potentials $V_\alpha$ at fixed $q = 0$ (expressed as $V_\alpha|_{q=0}$) corresponding to two enantiomers. Here $V_\alpha|_{q=0}$, which is not affected by the space-dependent part $\phi_\alpha$ of the overall phase, presents chirality-dependent features similar to the case in Ref. [44], where the molecular rotations were not considered. Comparing Fig. 2(c) with Fig. 2(d) in a typical wavelength range (about the order of 0.01 m) of electromagnetic fields, we find that, while $V_\alpha|_{q=0}$ in Fig. 2(d) is almost unchanged in the $p$ direction, $V_\alpha|_{p=0}$ in Fig. 2(c) varies quickly in the $q$ direction. This affects the center-of-mass motion of chiral molecules.

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$$\dot{q} = v_q - \frac{A_{\alpha,q}}{m},$$

$$\dot{p} = v_p - \frac{A_{\alpha,p}}{m},$$

(13)

$$v_q = \frac{1}{m} \left[ (v_q - \frac{A_{\alpha,q}}{m}) \partial_q A_q + \left( v_p - \frac{A_{\alpha,p}}{m} \right) \partial_p A_p - \partial_q V \right],$$

$$v_p = \frac{1}{m} \left[ (v_q - \frac{A_{\alpha,q}}{m}) \partial_p A_q + \left( v_p - \frac{A_{\alpha,p}}{m} \right) \partial_p A_p - \partial_p V \right].$$

Here $v_q$ and $v_p$ represent the components of the molecular velocity $v$ in the $\hat{q}$ and $\hat{p}$ directions, respectively. Since the equations of the motion for different inner dressed states are independent, we omit the subscript $\alpha$ for $q_\alpha$, $p_\alpha$, $v_q$, $\alpha$, $v_p$, $A_{\alpha,q}$, $A_{\alpha,p}$, and $V_\alpha$. The molecules are initially placed at $(q(0), p(0)) = (0, 0)$ with the initial speed $|\vec{v}(0)| = 0.1$ m/s.

In Fig. 3 we show the motion of two enantiomers in the time interval $t \in [0, 0.1]$ s, where the chiral molecules move in the distance of a typical wavelength scale. We are interested in two cases with the initial velocity along $\hat{q}$ direction [shown in Figs. 3(a1) and 3(a2)] and in the $\hat{p}$ direction [shown
in Figs. 3(b1) and 3(b2)], respectively. Figures 3(a1) and 3(b1) show the change of the displacement perpendicular to the initial velocity direction over time in these two cases, respectively. We find that these displacements in Figs. 3(a1) and 3(b1) are much smaller than the corresponding displacement in the initial velocity direction (about 0.01 m). Then the movement is regarded as one dimensional in the direction of the initial velocity.

The one-dimensional movement in the direction of the initial velocity reflects some characteristic of the induced scalar potentials in this direction. Thus, we show the change of the displacement increment in the direction of the initial velocity over time in these two cases in Fig. 3(a2) and 3(b2), respectively. For the same enantiomer with the same inner dressed state, comparing the displacement increment in the initial velocity direction in Figs. 3(a2) and 3(b2), the former [i.e., \( q(t) - v_q(0)t \)] is thousands of times longer than the latter [i.e., \( p(t) - v_p(0)t \)]. Comparing the separation distance between two enantiomers with the same inner dressed state in Fig. 3(a2) with the one in Fig. 3(b2), the former is also thousands of times longer than the latter. In this sense, we conclude that, when the chiral molecules move the distance of a typical wavelength scale, the separation distance between two enantiomers corresponding to the initial velocity along the \( \hat{q} \) axis (affected by \( \phi_\nu \)) is about 1000 times longer than that corresponding to the initial velocity along the \( \hat{p} \) axis (not affected by \( \phi_\nu \)).

The propagation trajectory of chiral molecules in the time interval \( t \in [0, 10] \) s is shown in Fig. 4, where the chiral molecules move a distance much longer than a typical wavelength. The initial velocity is in the \( \hat{q} \) direction and a \( \hat{p} \) direction. When we let the chiral molecules move in the distance much longer than a typical wavelength, the separation distance between two enantiomers corresponding to the initial velocity along the \( \hat{q} \) axis (affected by \( \phi_\nu \)) is comparable to that corresponding to the initial velocity along the \( \hat{p} \) axis (not affected by \( \phi_\nu \)).

V. CONCLUSION

We have discussed the spatial enantioseparation of gaseous chiral molecules taking into consideration molecular rotations. Comparing with the original works [44, 45] without consideration of molecular rotations, the overall phase of the coupling strengths has the space-dependent feature due to the specific requirements of the polarization directions of the three electromagnetic fields [57, 58]. This space-dependent feature provides an additional resource to generate the chirality-dependent induced gauge potentials. Specifically, the induced gauge potentials vary intensely in the length scale of the typical wavelength of the applied electromagnetic fields. Taking the 1,2-propanediol as an example and using the typical parameters, we have shown that this property of induced gauge potentials provides some advantages for spatial enantioseparation, especially when the chiral molecules move the distance of a typical wavelength scale.

We would like to point out that here we have focused only on the case of large detuning in the discussion for simplicity. In fact, our results of spatial enantioseparation can be extended to general cases without the condition of large detuning [45]. In addition, we have only considered that the molecular ensemble is initially located at the same point. In realistic cases, the molecular ensemble has an initial position distribution in the initial position. Consideration of the specific limit to the molecular ensemble is left for future work.
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