Cancellation of Stark Shifts in Optical Lattice Clocks by Use of Pulsed Raman and Electromagnetically Induced Transparency Techniques

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We propose a combination of electromagnetically induced transparency--Raman and pulsed spectroscopy techniques to accurately cancel frequency shifts arising from electromagnetically induced transparency fields in forbidden optical clock transitions of alkaline earth atoms. At appropriate detunings, time-separated laser pulses are designed to trap atoms in coherent superpositions while eliminating off-resonance ac Stark contributions, achieving efficient population transfer up to 60% with inaccuracy <10^{-17}. Results from the wave-function formalism are confirmed by the density matrix approach.

DOI: 10.1103/PhysRevLett.97.233001

PACS numbers: 32.80.–t, 32.70.Jz, 42.50.Gy, 42.62.Eh

In the field of optical frequency standards and clocks, single trapped ions [1] and alkaline earth atoms [2–6] are advancing clock performances. The advantage arises from superhigh resonance quality factors of these optical transitions [7], which are expected to be 10^5 better than microwave fountains. These fountain clocks are already below the 10^{-15} relative fractional uncertainty [8]. Fermionic isotopes of alkaline earths trapped in optical lattices at the magic wavelength [9] offer ultranarrow linewidths of a few millihertz without recoil and Doppler effects, but remain potentially sensitive to systematic effects arising from the nuclear spin-related tensor polarizability [3–5]. On the other hand, bosonic isotopes with no nuclear spin and a higher natural isotopic abundance avoid multiple hyperfine components but lack direct excitation of the clock transition [1] ↔ [2] in Fig. 1(a). Indirect excitation via continuous-wave electromagnetically induced transparency (EIT) has been proposed to probe these forbidden transitions [10,11]. A similar scheme for the ^{174}Yb forbidden clock transition was implemented by applying a dc magnetic field for state mixing [6].

All such schemes can suffer from Stark shifts due to nonresonant electric-dipole couplings of the clock levels to other states induced by the applied electromagnetic fields [12,13]. Reference [10] provides some detailed calculations of these shifts. To further reduce this potential systematic error, we could apply an approach similar to that used for the determination of the magic wavelength [3,4] or the hyperpolarizability contribution to the ac Stark shifts [14]: Measurements at different field strengths are used to extrapolate the clock frequency to vanishing field. However, this simple approach does not apply to the EIT-related schemes where the applied field strength modifies also the optical pumping time required to prepare the atoms in a coherent superposition [15]. The preparation time required for optimal signal contrast and clock stability becomes impractically long at low field strengths. But using large fields increases the ac Stark shifts and limits the clock accuracy. To overcome these limits, the pulsed scheme proposed in this Letter [Fig. 1(c)] optimizes clock performance by using time-separated laser pulses to prepare and interrogate the optical clock transition [16]. It is an original mix of Ramsey spectroscopy and highly efficient population transfer under coherent population trapping (CPT) [17]. The first pulse prepares atoms in a coherent superposition and the second pulse probes the clock frequency. This configuration produces a large contrast in the detected clock signal. More importantly, as the detunings of the applied fields affect the phase evolution of the atomic wave-function, a proper combination of the common mode laser detuning Δ_0 and pulse durations τ, τ_m (Fig. 1) reduces the clock shift to ~10^{-17}. The discussion presented here reveals for the first time a general relation connecting the preparation time of the Raman

![FIG. 1 (color online). (a) Three-level atom-light configuration for an optical lattice clock based on time-separated laser pulses including relaxation and decoherence rates. The optical detunings Δ_1 = Δ_0 + η_1, Δ_2 = Δ_0 - Δ_1, Δ_0, Δ_1, Δ_2 include ac Stark shifts η_1 from off-resonant levels. Here Δ_0 is the common mode detuning and Δ_1 denotes deviation from the Raman condition. (b) The corresponding dressed-state representation of bright |Ψ⟩ and dark |Ψ_NC⟩ states defining the clock transition. (c) The probing pulse sequence.](https://example.com/figure1.png)
coherence and the signal contrast in the subsequent detection of this coherence, relevant to many EIT or CPT related experiments.

The atomic evolution between $^1S_0$ and $^3P_0$ is properly described in the dressed-state picture [Fig. 1(b)],

$$|1\rangle = \frac{\Omega_1}{\sqrt{\Omega_1^2 + \Omega_2^2}} |\Psi_C\rangle + \frac{\Omega_2}{\sqrt{\Omega_1^2 + \Omega_2^2}} |\Psi_{NC}\rangle, \quad |2\rangle = \frac{\Omega_2}{\sqrt{\Omega_1^2 + \Omega_2^2}} |\Psi_C\rangle - \frac{\Omega_1}{\sqrt{\Omega_1^2 + \Omega_2^2}} |\Psi_{NC}\rangle,$$

(1)

where $\Omega_1$ and $\Omega_2$ are the Rabi frequencies for the transitions $^1S_0 \leftrightarrow ^1P_1$ and $^3P_0 \leftrightarrow ^1P_1$. For an ideal 3-level system described in (1), the dark state $|\Psi_{NC}\rangle$ remains insensitive to light shift, while the bright state $|\Psi_C\rangle$ is always coupled to the laser light. A realistic atomic clock has to deal with off-resonant ac Stark shifts acting on $|\Psi_C\rangle$ while atoms are pumped into $|\Psi_{NC}\rangle$ with a few spontaneous emission cycles. Thus, a judicious tradeoff between the short-time dynamics for a high-contrast signal (large optical pumping) and the reduced external ac shifts (and resonance power broadenings) under a low field strength needs to be found for practical realizations of these EIT or Raman-type clocks.

To describe our pulsed method, we start from a three-level configuration as shown in Fig. 1(a). The optical Bloch equations (OBEs) describe three-level dynamics including external shifts, relaxations, and decoherences between atomic states [18] in terms of the density matrix:

$$\dot{\rho} = -\frac{i}{\hbar} [H, \rho] + \mathcal{R} \rho.$$

(2)

In the interaction picture, the atom-light Hamiltonian $H$ and relaxation matrix $\mathcal{R} \rho$ become

$$\frac{H}{\hbar} = \begin{pmatrix} \Delta_1 & 0 & 0 \\ 0 & \Delta_2 & \Omega_2 \\ 0 & \Omega_1 & 0 \end{pmatrix}, \quad \mathcal{R} \rho = \begin{pmatrix} \Gamma_{31} \rho_{33} & -\gamma_c \rho_{12} & -\gamma_c \rho_{13} \\ -\gamma_{c1} \rho_{21} & \Gamma_{32} \rho_{33} & -\gamma_c \rho_{23} \\ -\gamma_{c1} \rho_{31} & -\gamma_{c2} \rho_{32} & -\Gamma_{33} \rho_{33} \end{pmatrix}. \quad \Gamma = \Gamma_{31} + \Gamma_{32}, \gamma_c = \gamma_{c1} + \gamma_{c2}, \text{ and the Raman decoherence } \gamma_c \text{ [see Fig. 1(a)]. Electric and/or magnetic dipole couplings determine the Rabi frequencies } \Omega_i \ (i = 1, 2).$$

Equation (2) describes the dynamics of a closed $\Lambda$ system where optical detunings $\Delta_i$ include ac Stark shifts $\eta_i$ from nonresonant electric-dipole couplings of $|1\rangle$ and $|2\rangle$ to other states. For $\Omega_1, \Omega_2 \approx \Gamma_{31}, \Gamma_{32}, \gamma_{c1}, \gamma_{c2}$, the population in state $|3\rangle$ is slaved to the population difference $\Delta n(t) = \rho_{22}(t) - \rho_{11}(t)$ and Raman coherence $\rho_{12}(t)$. This allows finding analytical solutions to Eq. (2) by adiabatic elimination of the intermediate state $|3\rangle$ [19,20].

The reduced two-level system dynamics are described by a Bloch-vector representation [21,22].

To remove ac Stark shifts while maintaining a high signal contrast, we apply the Ramsey technique for EIT or Raman fields to this effective two-level system, minimizing systematic frequency shifts over the free-evolution time $T$. The Ramsey-like sequence of preparation, free-evolution, and probe, followed by the final destructive detection of the ground-state population, is indicated in Fig. 1(c). This eliminates power broadening of the clock transition which is always present for continuous excitation [23]. By solving for the two-level system as in [22] we can express the populations as

$$\rho_{ii} = \alpha_{ii}(\tau, \tau_m)[1 + \beta_{ii}(\tau, \tau_m)e^{-\gamma_c T}\times \cos[\delta_{i} T - \Phi(\tau, \tau_m)]]\quad (4)$$

where $\alpha_{ii}(\tau, \tau_m)$ is the overall envelope function and $\beta_{ii}(\tau, \tau_m)$ is the amplitude of fringes, both containing exponential decays $e^{-\gamma_c T}$ and $e^{-\gamma_m T}$ to their steady states [21]. $\tau_p$ is the characteristic optical pumping time. The atomic phase shift $\Phi$ produces an approximated clock frequency shift assuming $\tau, \tau_m \ll T$:

$$\delta \nu = \frac{\Phi(\tau, \tau_m)}{2\pi T(1 + \tau_p \tau_m)}, \quad (5)$$

which includes all ac Stark contributions accumulated during the pulsed interactions. Hence, a longer free-evolution time $T$ reduces the light shifts on the clock transition. Furthermore, as will be shown below, a special value $\Delta_0$ of the common detuning which can be found to suppress ac Stark effects on the clock frequency. Study of the population dynamics from Eq. (4) leads to an expression for the time $\tau_p$ that is required to pump atoms into their final steady state, simplified for $\Delta_0 = \Delta_1 = \Delta_2$:

$$\tau_p(\Delta_0) = \frac{2 \Delta_0^2 + \Gamma^2/4}{\Gamma(\Omega_1^2 + \Omega_2^2)}[1 - \gamma_c \left(\frac{\Omega_1^2 - \Omega_2^2}{\Omega_1^2 + \Omega_2^2}\right)]^{-1}. \quad (6)$$

Here $Y = (\Gamma_{31} - \Gamma_{32})/\Gamma$ is the branching ratio difference for the intermediate state which scales the contribution of each Rabi frequency to the pumping rate $\tau_p^\text{-1}$. This time scale $\tau_p$ sets the EIT or CPT response in either transient or steady states. Previous work on EIT or CPT concentrates mainly on the symmetric case with $Y = 0$. But in the case of alkaline earths where $Y = \pm 1$, Eq. (6) shows that the Rabi frequency associated with the weaker transition dictates $\tau_p$. For the $^{88}\text{Sr}$ lattice clock where $\Gamma_{31} = 2\pi \times 32 \text{ MHz} \gg \Gamma_{32} = 2\pi \times 620 \text{ Hz}$ (i.e., $Y \sim 1$), the pumping time at resonance $\tau_p(0)$ is determined by the magnetic dipole coupling $\Omega_2$ between $^3P_0$ and $^1P_1$. Figure 2 shows the dependence of $\tau_p(0)$ on each Rabi frequency while keeping the other one fixed. The dotted lines are the corresponding differential ac Stark shift of the clock frequency in the steady state regime. Note that small ac Stark shifts correspond to long optical pumping times conflicting with realistic clock duty cycles. For instance, the proposal by [10] with an ac Stark shift below 21.7 mHz for
an accuracy of $2 \times 10^{-17}$ leads to a signal contrast of a few percent only after 160 s. The scheme presented here finds a combination of parameters that maximizes contrast while suppressing ac Stark shifts, exploiting the transient dynamics for short pulses and detuned laser fields. The highly asymmetric $Y$ allows this scheme to uniquely exploit ground-state detection with a high-contrast narrow resonance manifested in the atomic population transfer. In the region of detuning between Raman spectroscopy ($\Delta_0/\Gamma \gg 1$) and EIT/CPT ($\Delta_0/\Gamma \ll 1$), we find contrasts of up to 60\%, even though $\tau \ll \tau_p(\Delta_0) \approx 100$ s. This same approach could be extended easily to the four level scheme [11], the magnetic induced optical transition [6] or any other clock configurations involving dark states.

The small difference between the field-free clock detuning $\delta$, and the ac Stark shifted detuning $\Delta_1 - \Delta_2 = (\eta_1 - \eta_2) + \delta$, under laser fields leads to a small phase shift of the Ramsey-EIT fringe defined by $\Phi(\tau, \tau_m)$ in Eq. (5) [19,20]. Solving Eq. (2) numerically, we find that a judicious choice of the laser detuning ($\Delta_0)_m$ cancels the external ac Stark shifts, minimizing the influence to the clock transition when high field strengths are used to rapidly drive EIT resonances. To confirm these results, we also establish an analytical expression for $\Phi(\tau, \tau_m)$ based on the atomic wave-function formalism [24], using the Hamiltonian of Eq. (3) adding only the term $-i\Gamma/2$ associated with spontaneous relaxation [25,26] and neglecting all lattice decoherences. By adiabatic elimination of state [3], within an effective two-level system including only the clock states [1] and [2], the amplitudes evolve with a matrix $M$, generalized from Ref. [27] by assuming $\Delta_1 \neq \Delta_2$:
of \( \langle \Delta_0 \rangle_m \) where clock shift is suppressed, as indicated in Fig. 3(b). When different free-evolution times \( (T) \) or pulse durations \( (\tau = \tau_m) \) are used, the accumulated phase shift changes, leading to variations in the dependence of \( \delta \nu \) on \( \Delta_0 \). Hence, the curves depicting \( \delta \nu \) versus \( \Delta_0 \) for different \( T \) rotate around \( \langle \Delta_0 \rangle_m \), with no changes in the signal contrast. In the second approach [Fig. 3(d)], we find the values of \( \Delta_0 \) where \( \delta \nu \) for \( \tau = 1 \) s is the same as that for some other values of \( \tau \). These values of \( \Delta_0 \) can be plotted as a function of \( \tau \) and extrapolated to \( \tau = 0 \) to find \( \langle \Delta_0 \rangle_m \). However, the signal contrast under smaller \( \tau \) is reduced due to the effect of pulse preparation on population transfer.

From Eq. (4) we find spectral line shapes and transition probabilities as a function of the experimental parameter \( \delta_\nu \), shown in Fig. 4(a). Since \( \tau \ll \tau_{\nu}(\Delta_0) \), the two-photon resonance has a Fourier transform linewidth given by the duration \( \tau \) where power broadening effects have been eliminated. The spectra also exhibit the typical coherent Ramsey nutations with period \( \sim 1/2T \) and a central fringe free from systematic shifts. We have also determined the sensitivity of \( \delta \nu \) to laser intensities \( (\Omega_i) \) and detunings, demonstrating that the uncertainty of the optical clock frequency \( <5 \) mHz \( (\sim 10^{-17}) \) is achievable by controlling \( \Delta_0 \) at the 100 kHz level around \( \langle \Delta_0 \rangle_m \). Meanwhile, \( \Omega_i \) fluctuations should be controlled \( <0.5\% \). We note that for a given set of \( \tau \) and \( \Omega_i \), different values of \( \langle \Delta_0 \rangle_m \) can be found. For example, \( \langle \Delta_0 \rangle_m = 200 \) MHz is another optimum value for larger \( \Omega_i \) [Fig. 4(b)]. In this case, the signal contrast is further improved with a population transfer of up to 60%, leading to enhanced clock stability but also slightly larger uncertainty.

In summary, our method achieves the \( 10^{-17} \) accuracy expected for a “light-insensitive” lattice clock with pulsed EIT or Raman techniques. A signal contrast of 20% to 60% (Fig. 4) can be achieved, including realistic lattice decoherence times [4]. Extensions are possible to the proposal of [11] by replacing the \( \left| P_1 \right\rangle \) state with \( \left| P_1 \right\rangle \) to magnetic field induced optical transitions [6], for other species such as \( ^{52}\text{Cr} \) [30], and for nuclear clock transitions [31].

We thank J. Dalibard, T. Ido, T. Zelevinsky, and C. Oates for discussions. This work is supported by ONR, NIST, and NSF. T. Z-W. thanks Observatoire de Paris and Délégation Générale de l’Armement for support.

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