

Multi-ensemble metrology by programming local rotations with atom movements

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Current optical atomic clocks do not utilize their resources optimally. In particular, an exponential gain in sensitivity could be achieved if multiple atomic ensembles were to be controlled or read out individually, even without entanglement. However, controlling optical transitions locally remains an outstanding challenge for neutral-atom-based clocks and quantum computing platforms. Here we show arbitrary, single-site addressing for an optical transition via sub-wavelength controlled moves of atoms trapped in tweezers. The scheme is highly robust as it relies only on the relative position changes of tweezers and requires no additional addressing beams. Using this technique, we implement single-shot, dual-quadrature readout of Ramsey interferometry using two atomic ensembles simultaneously, and show an enhancement of the usable interrogation time at a given phase-slip error probability. Finally, we program a sequence that performs local dynamical decoupling during Ramsey evolution to evolve three ensembles with variable phase sensitivities, a key ingredient of optimal clock interrogation. Our results demonstrate the potential of fully programmable quantum optical clocks even without entanglement and could be combined with metrologically useful entangled states in the future.

Sensors based on quantum probes provide some of the most precise measurements in science^{1–5}. For many such systems, fundamental sensitivity limits can be improved through entanglement^{6–9}, but in the presence of noise, a practical advantage of such schemes is not guaranteed^{10,11}. A complementary approach studies optimal metrology with entanglement-free quantum control and readout methods. For both approaches, an important figure of merit is not just the sensitivity to a given observable but also the dynamic range over which that observable can be unambiguously estimated^{12–15}.

In the particular case of optical atomic clocks¹, the observable of interest is the stochastically evolving phase of a laser acting as a local oscillator, which is mapped into a population imbalance of an ultranarrow optical transition. The clock stability improves with the interrogation time, but the phase can only be unambiguously

mapped when it is in the range of $[-\pi/2, \pi/2]$; phases outside this range lead to phase-slip errors, which limits the attainable interrogation time at a given phase-slip error probability in the case of local-oscillator-limited clocks. Optimal readout schemes^{12,13,16} could exponentially improve the attainable interrogation time, but require local rotational control over sub-ensembles during the sensing protocol or local mid-circuit readout and reset, both of which have not been demonstrated to date.

Here we show local control of optical transitions in a tweezer-array clock^{17–19} by using rearrangement techniques^{20–25} on atoms in superposition states to precisely control the position-dependent phase imprinted by light–matter interaction. The scheme^{26,27} is experimentally simple and highly robust as it solely relies on the relative stability of the tweezer positions and does not

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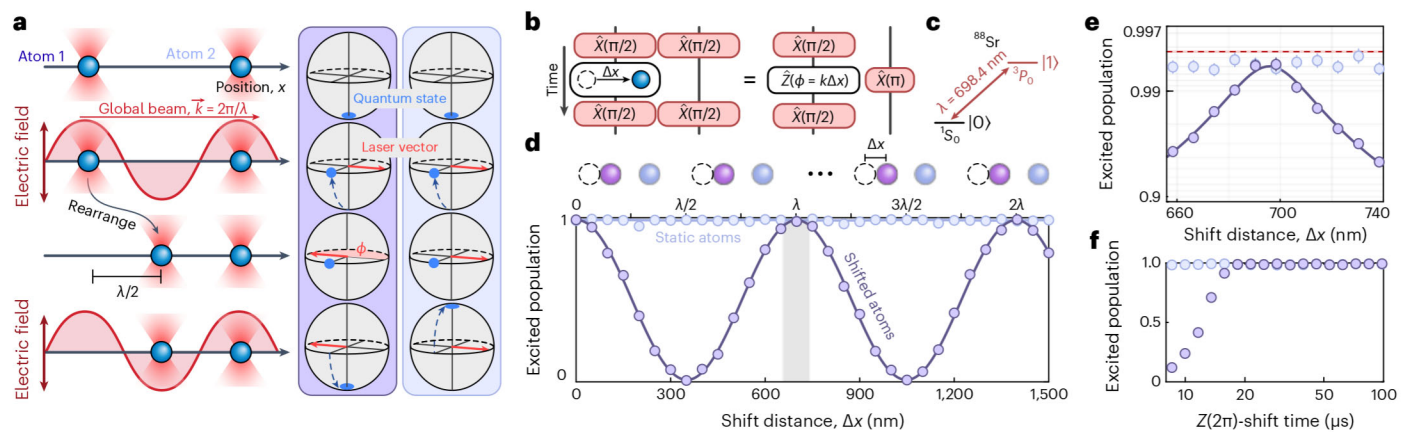


Fig. 1 | Single-site addressing with movement-induced phase shifts. **a**, We consider two atoms individually trapped in optical tweezers, both initially in the electronic ground state. Travelling light emitted from a global laser beam applies a $\pi/2$ rotation to both atoms, and is disabled, but remains phase coherent with the atomic transition. One of the atoms is then moved by half the laser wavelength (λ) from its initial position, rotating the effective local laser frame by an angle $\phi = \pi$. When the laser drive is restarted to apply another $\pi/2$ pulse, the moved atom now rotates back to the ground state, whereas the static atom rotates to the excited state. **b**, Control over the atom displacement Δx is equivalent to arbitrary local rotations of the laser drive by $\phi = k\Delta x$ about the $\hat{Z}$ axis. **c**, We implement this protocol with an array of ^{88}Sr atoms utilizing the ultranarrow $^1S_0 \leftrightarrow ^3P_0$ transition with $\lambda = 698.4 \text{ nm}$ for global driving. **d**, With an array of 39 tweezers in one dimension (top), we apply the protocol in **b**, shifting every odd site (purple markers) in the array and leaving all the even sites static

(blue markers) during the dynamics. A sinusoidal oscillation emerges in the excited state population of the shifted sites (bottom), with a period of $699(1) \text{ nm}$. **e**, Focusing on the region around $\Delta x = \lambda$ (grey-shaded region in **d**), we find that the shifted atom shows no measurable loss in fidelity compared with the unshifted atoms. Correcting for the bare fidelity for performing a global $\hat{X}(\pi)$ rotation (red dashed line; $0.9956(1)$), we find that the shift operation is performed with a fidelity of $0.9984(5)$. The ratio of the shifted to unshifted fidelities is $0.9998(5)$, suggesting that the dominant source of error comes from global laser phase noise during the finite wait time required to perform the shift, rather than the movement itself. From the data in **d**, we find that the crosstalk to the static atoms is $0.1(2)\%$, consistent with **0**. **f**, The shift to apply a $\hat{Z}(2\pi)$ rotation can be performed without a noticeable loss of fidelity down to shift times of $\sim 20 \mu\text{s}$; the data in **e** are taken with a shift time of $32 \mu\text{s}$, in addition to an extra wait time of $34 \mu\text{s}$ to account for finite jitter in the control timings.

involve any auxiliary addressing beams. Using this technique, we demonstrate arbitrary, parallel, single-site-resolved optical qubit rotations with high fidelity.

We utilize such rotations to double the dynamic range of optical Ramsey spectroscopy by performing simultaneous evolution on two separate atomic ensembles within one tweezer array, each of which measures a different phase quadrature¹⁶; we extend the coherent interrogation time by a factor of $3.43(13)$ relative to the standard, single-ensemble sequence. Finally, we realize a proof-of-principle protocol for programming local dynamical decoupling sequences during Ramsey interrogation such that different ensembles within a single atom array have different sensitivities to phase variations, and discuss its implementation as part of a general protocol for improving clock stability^{12,13}.

Aside from clocks, our technique for implementing local, parallel rotations about arbitrary axes might also find use in neutral-atom quantum computing platforms utilizing optical transitions^{27,28}, where the local coherent control of optical qubits has not been demonstrated before. More generally, our results point to a future of fully programmable neutral-atom optical clocks that incorporate features of quantum computers.

The basic principle of our scheme is illustrated in Fig. 1a. We consider two atoms both initially in the ground electronic state $|0\rangle$, interacting with a global laser beam characterized by wavevector $k = 2\pi/\lambda$ and wavelength λ , propagating along the array axis. With the globally applied laser, we create an equal superposition state of $|0\rangle$ and the excited state $|1\rangle$; in a Bloch sphere picture, this corresponds to a $\pi/2$ rotation around the x axis ($\hat{X}(\pi/2)$). The laser beam is then extinguished with an optical modulator, but remains phase coherent with the atomic transition. Using atom rearrangement techniques^{20–25}, one of the atoms is shifted from its original position by Δx , applying an effective phase shift of $\phi = k\Delta x$ (Methods). In Fig. 1a, we first consider the special case of $\Delta x = \lambda/2$, or equivalently, a π rotation around the z axis ($\hat{Z}(\pi)$) for the shifted atom (Fig. 1a). Subsequently, we apply a second global $\hat{X}(\pi/2)$

rotation with the same laser as before; the shifted atom now rotates back to $|0\rangle$ because of the movement-induced phase shift, whereas the unshifted atom completes its rotation to $|1\rangle$.

The main principle behind this scheme is a locally controlled change of the relative phase between the atomic dipole oscillation and the phase of the laser while the atom is in a superposition state; in essence, our scheme realizes a locally controlled Ramsey sequence with global driving (Methods). Similar techniques have been used in the context of ion-trap experiments with two ions²⁶, but not in a scalable fashion, as is possible with tweezer arrays²⁷.

We show an experimental demonstration with our ^{88}Sr optical tweezer-array experiment^{17,29,30}. We employ a one-dimensional array of 39 optical tweezers generated via an acousto-optic deflector (AOD) driven by an arbitrary waveform generator (AWG). This allows for precise control over the relative tweezer positions at the nanometre level, enabling arbitrary $\hat{Z}(\phi)$ rotations (Fig. 1b). Global driving is performed on the ultranarrow $^1S_0 \leftrightarrow ^3P_0$ optical clock transition with a transition wavelength of $\lambda = 698.4 \text{ nm}$ (Fig. 1c).

In a first experiment, we globally apply an $\hat{X}(\pi/2)$ operation to the entire array, then shift every odd site by the same distance Δx , apply another global $\hat{X}(\pi/2)$ rotation and finally measure the excited state population in both shifted and unshifted sub-arrays. The excited state population of the shifted atoms P_s shows sinusoidal oscillations with a period of $699(1) \text{ nm}$ as a function of Δx , consistent with $\phi/2\pi = \Delta x/\lambda$, where λ is the transition wavelength. The quoted error on this measurement is purely statistical, and ignores potential systematic error arising from the independent distance calibration performed with an optical resolution test target. We note that the present measurement is likely a far more precise and accurate distance calibration tool, and could find use as an effective in situ laser-based ruler with applications for the precision determination of distance-dependent inter-atom effects, such as Rydberg interactions³¹.

To quantify the phase-shift fidelity, we focus on a narrow region around $\Delta x = \lambda$, corresponding to a $\hat{Z}(2\pi)$ rotation (Fig. 1e). A quadratic

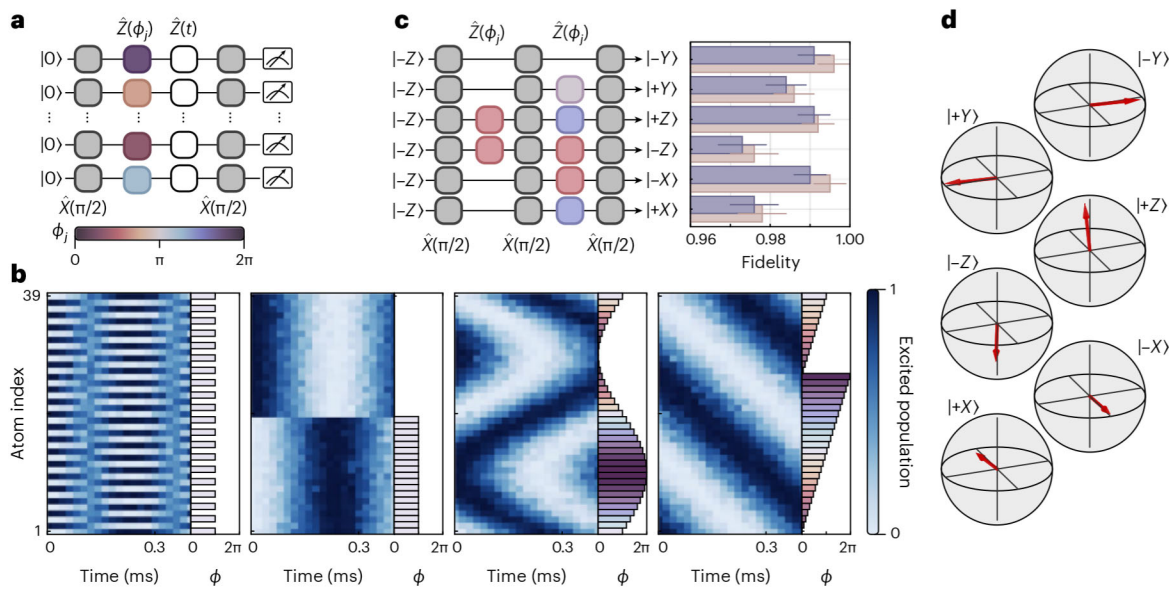


Fig. 2 | Arbitrary, parallel and local rotations. a, We implement site-resolved phase shifts ϕ_i during the dark time t of standard Ramsey interrogation by inserting arbitrary and parallel shifts of various distances to the array of atoms. **b**, Results of this operation as a function of Ramsey time (x axis) for different tweezers in the array (y axis). The corresponding programmed phase-shift pattern is shown on the right of each panel. **c**, By applying multiple global $\hat{X}(\pi/2)$ pulses (grey blocks), in tandem with local movement shifts (same colour scale as

in **a**), arbitrary local rotations can be performed. We show a demonstration by rotating an array of six atoms, initially in the $|0\rangle = | -Z \rangle$ state, in parallel to the six cardinal states ($| -Z \rangle$, $| +Z \rangle$, $| -Y \rangle$, $| +Y \rangle$, $| -X \rangle$ and $| +X \rangle$), achieving an average fidelity of 0.984(2) (blue bars) and 0.987(2) after SPAM correction (tan bars), limited by global $\hat{X}(\pi/2)$ fidelity and decoherence during the time needed for movement (Methods). **d**, Bloch sphere visualizations of the states measured with quantum state tomography in **c**.

fit to $P_s(\Delta x)$ shows a maximum value of $P_s = 0.9940(5)$ (not corrected for state preparation and measurement (SPAM) errors), consistent with the mean excited state population of unshifted atoms, $P_u = 0.9942(2)$, in the same range. Correcting for the bare $\hat{X}(\pi)$ fidelity (Fig. 1e, red dashed line) of 0.9956(1) shows that the shift operation is performed with a fidelity of 0.9984(5). We note that applying SPAM correction on the bare fidelities maintains the shift fidelity largely unchanged as it is calculated from the ratio of the two. The ratio of the shifted to unshifted fidelities is 0.9998(5), suggesting that the dominant source of error comes from global laser phase noise during the finite wait time required to perform the shift, rather than the movement itself. We study the fidelity to perform the $\hat{Z}(2\pi)$ rotation as a function of shift time (Fig. 1f), and find that the fidelity remains constant down to shift times of $t_s = 20 \mu\text{s}$; data in Fig. 1e were taken with $t_s = 32 \mu\text{s}$, plus an additional $34 \mu\text{s}$ of wait time to account for jitter in the subsequent control timings. Importantly, for all shift distances (Fig. 1d), the excited state population of the neighbouring unshifted atoms is nearly constant, showing crosstalk of only 0.1(2)% (Methods).

Arbitrary rotation patterns can be imprinted on the array by shifting all of the atoms by varying distances such that rotations about the z axis with tweezer-resolved phase ϕ_i are applied (Fig. 2a). We show the results of time-resolved Ramsey spectroscopy for four different choices of single-site addressing patterns, demonstrating arbitrary, site-reversed and parallel $\hat{Z}$ rotations (Fig. 2b). Such addressing patterns could be used to negate variations in the transition frequency across the array, for instance due to gradients in magnetic field or from finite differences in the wavelengths of the tweezers generated by an AOD¹⁷. Combining these single-site $\hat{Z}(\phi_i)$ rotations with a series of global $\hat{X}(\pi/2)$ pulses allows for rotations about any axes, not just the z axis. As a demonstration (Fig. 2c,d), we choose a set of six contiguous atoms, initially in the ground state (denoted here as $| -Z \rangle$), and rotate them each in parallel into the six states $| -Z \rangle$, $| +Z \rangle$, $| -Y \rangle$, $| +Y \rangle$, $| -X \rangle$ and $| +X \rangle$, with an average fidelity of 0.984(2) (0.987(2) SPAM-corrected), as determined by state tomography (Methods). The dominant

limitations to this value are likely from global drive infidelity and dephasing during the finite shift times.

We note that while here we have demonstrated our protocol on a one-photon optical transition, it could be used to induce a similar effect for two-photon Raman transitions, for instance between hyperfine states³², assuming the two beams are counter-propagating. Further, the movement-induced phase shifts employed here rely solely on a relative change in tweezer position, in contrast to alternative techniques that apply additional addressing beams^{33–36}, where the phase shift is proportional to a local addressing beam's intensity. While the addressing beam intensity and alignment are prone to drifts on experimental timescales, relative atom movements are ultimately derived from the radio-frequency electronic output of an AWG, which is precise, consistent and robust. We emphasize that our results did not utilize noise-compensating composite pulse sequences and that all data were taken without any system realignments or recalibrations of the atom movements.

We now demonstrate that access to such robust, high-fidelity, single-site operations can enable enhanced sensing protocols for entanglement-free metrology. In particular, several protocols relying on local control have been proposed for improving the stability of phase estimation^{12,13,16,37} by increasing the dynamic range in which the stochastically evolving laser phase θ can be estimated.

Here we experimentally show one such proposal¹⁶, by splitting the array into two sub-ensembles using local addressing to perform Ramsey interferometry simultaneously in two orthogonal bases, X and Y , yielding populations $P^{(x)}$ and $P^{(y)}$. While readout in a single basis limits the invertible phase range to $\theta \in [-\pi/2, \pi/2]$, readout in both bases allows this range to be unambiguously extended to $[-\pi, \pi]$ (Fig. 3a). Consequently, we can afford a longer Ramsey interrogation time before θ drifts outside of the invertible range, which would cause a phase-slip error. Note that while the atom number in each quadrature has been halved, this typically does not increase the quantum projection noise (QPN)³⁸ from the dual-quadrature measurement compared to a single-basis measurement (Methods)^{12,16}.

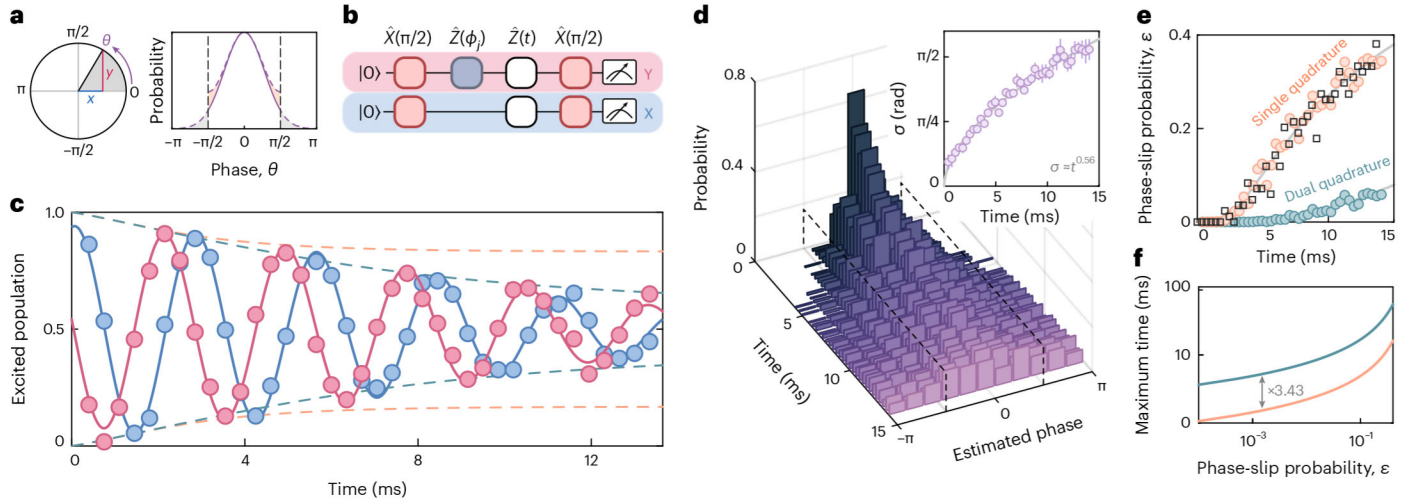


Fig. 3 | Enhanced sensing with dual-quadrature measurement. **a**, For a given phase angle θ , population measurement in only a single basis, for example Y , can only be inverted within a dynamic range of $-\pi/2 < \theta < \pi/2$. By measuring both quadratures X and Y , this dynamic range can be doubled to $-\pi < \theta < \pi$, allowing for interrogating larger spreads in phase, such as when measuring for longer times. **b**, We implement the dual-quadrature readout of Ramsey interrogation by applying local $\pi/2$ -phase shifts to all the odd sites in the array. **c**, With single-quadrature readout, the interrogation time is limited due to phase slips, visible by the separation between a decay envelope reconstructed from the single-quadrature phase spread (orange dashed line) and the averaged Ramsey signal (blue and red markers and lines). The equivalent reconstruction with dual-quadrature readout (green dashed line) is accurate up to longer times. **d**, To perform this reconstruction, we measure the time-resolved probability distributions of the estimated phase relative to the mean from dual-quadrature measurement. As the standard deviation (σ) of the phase distribution grows

(inset), the estimated phase begins exceeding the $-\pi/2 < \theta < \pi/2$ range for normal spectroscopy (black dashed lines), but is still resolvable via dual-quadrature measurement. Note that the time-dependent contribution from QPN to the standard deviation has been subtracted off in the inset (Methods). **e**, We estimate the phase-slip probability ϵ for single-quadrature (orange circles) and dual-quadrature (green circles) measurements by fitting a folded Gaussian to the time-resolved estimated phases in **d**. The fit is folded over at the boundaries of the dynamic range to account for the behaviour of phase slips, as that in **a**. For the single-quadrature case, we also directly estimate the probability from the underlying data (squares), which is in good agreement with the estimate from the fit. The solid lines are the predicted phase-slip probabilities from the fit in the inset of **d**. This fit is used to estimate the decay envelopes in **c**. **f**, For a given allowable phase-slip probability, the enhanced dynamic range of the dual-quadrature readout improves the maximum possible interrogation time. For our particular phase-growth profile (inset of **d**), the improvement is a factor of ~ 3.43 .

To implement this dual-quadrature readout, we perform Ramsey interferometry with the addition of a $\hat{Z}(\pi/2)$ rotation to all odd sites in the array before readout (Fig. 3b). The resultant oscillations in $P^{(x)}$ and $P^{(y)}$ show a $\pi/2$ -phase shift between the even (X) and odd (Y) sites in the array (Fig. 3c). For every repeated measurement (indexed by j) at time t , we estimate the phase as¹²

$$\theta_j(t) = \arg(z_j^{(x)}(t) + iz_j^{(y)}(t)), \quad (1)$$

where $z_j^{(x,y)}(t) = (2P_j^{(x,y)}(t) - 1)$ and $\arg$ is the argument function. We then calculate the difference, $\delta_j(t)$, of $\theta_j(t)$ from its mean phase (Methods).

We plot the probability distribution $\mathcal{P}(\delta_j(t))$ (Fig. 3d) and observe a continuous growth of its standard deviation (s.d.) σ (inset). We stress that we are interested in the distribution of the laser phase itself, which determines the phase-slip error probability. Hence, we have subtracted the contribution from QPN to our experimental data (Fig. 3d, inset) (Methods). We find that the laser phase s.d. grows with time as a power law: $\sigma = \beta t^\alpha$, with $\alpha = 0.56(2)$, which we attribute to a power spectral density composed of $1/f$ and white frequency noise. If this s.d. of the laser phase itself becomes too large compared to the dynamic range, frequent phase-slip errors occur. In Fig. 3e, we evaluate the phase-slip probability (ϵ) that the phase has exceeded the bounds of $[-\pi/2, \pi/2]$ (in emulation of a theoretical single-basis measurement; Fig. 3d, black dashed lines) or $[-\pi, \pi]$ (for the dual-quadrature readout); we find that the error probability for the single-basis case quickly becomes substantially larger at shorter interrogation times (Methods).

We further characterize the maximum interrogation time $T_{\max}(\epsilon)$ for which the phase-slip error probability is still below a threshold ϵ (Methods). We find that $T_{\max}(\epsilon)$ is significantly increased for the dual-quadrature case (Fig. 3f) by a factor of 3.43(13), the exact

numerical value of which is determined by the phase s.d. growth rate observed experimentally and is related to the laser noise spectrum (Methods). Such elongation in the attainable interrogation time can be directly translated to enhanced stability in a metrological setting. For example, in a zero dead-time optical clock, the stability is proportional to the square root of $T_{\max}(\epsilon)$, such that we can project an increase in stability by a factor of $\sqrt{3.43} \approx 1.8$ for our particular noise profile. This would constitute a practical improvement in phase estimation without increasing the probability of phase-slip errors, a common problem for entanglement-enhanced metrology schemes^{11,39}.

Even greater enhancements in dynamic range, and hence clock stability, could be possible through the use of multiple ensembles with different interrogation times by utilizing fast quantum non-demolition measurements^{40,41} or by explicitly programming ensembles with different sensitivities to the global laser phase^{12,13}. In the latter of these protocols, the total number of atoms is evenly divided into M ensembles, which are each further subdivided into two sub-ensembles for dual-quadrature measurement. One ensemble is used for normal phase measurement, while for the rest the free evolution time is reduced by factors of $2^{-1}, \dots, 2^{-M}$, or equivalently, their effective phase accumulation is reduced by the same amount. If this procedure is performed correctly, the effective ensemble coherence times will then be extended by factors of $2, \dots, 2^{M-1}$, meaning that slower-evolving ensembles can be used to probe for phase slips in the fastest ensembles. This then allows for phase estimation over a wider dynamic range beyond $[-\pi, \pi]$, and potentially allows for an improved scaling of the clock stability with atom number¹² at a fixed phase-slip probability (Fig. 4a).

As an outlook, we demonstrate a proof of principle of local control techniques towards such protocols by performing local dynamical decoupling such that three ensembles experience different effective Ramsey evolution times of T , $T/2$ and $T/4$. This is accomplished by

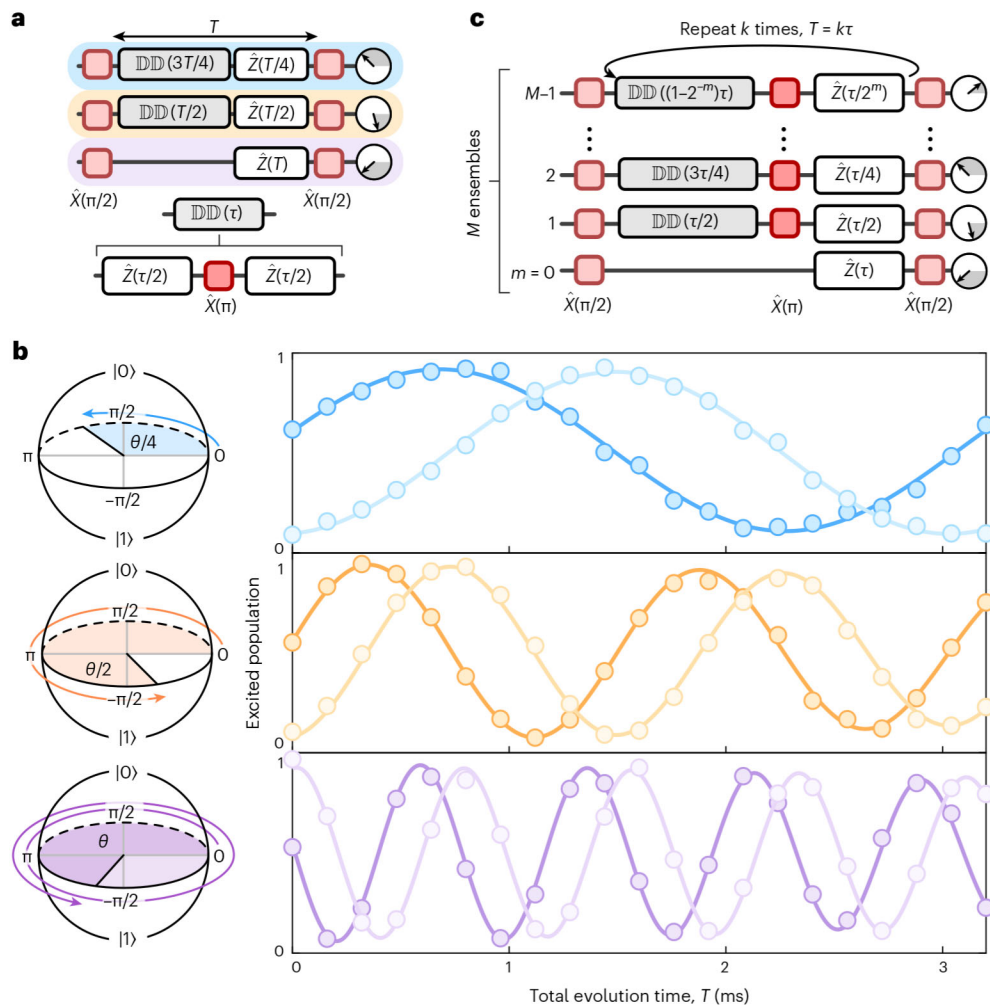


Fig. 4 | Local dynamical decoupling towards optimal metrology. a, We split the array into three ensembles, and perform a local dynamical decoupling (DD) sequence such that even though the total Ramsey dark time is T , individual ensembles experience different effective evolution times of $T/4$, $T/2$ and T . The phase of each ensemble is then measured using dual-quadrature readout. **b**, Slower-evolving ensembles (those which experience less evolution time) can be used to detect phase slips in faster-evolving ensembles, extending the effective interrogation time of optical clocks. Following the sequence in **a**, we find the three ensembles evolve at relative rates of 1.00:1.99(1):4.10(4) with

respect to the total evolution time T . The demonstrated scheme in **a** and **b** is effective for the case of slow frequency noise where the corresponding noise correlation time is longer than the total evolution time. **c**, To handle generic time-dependent noise with shorter correlation times, we envision breaking the total evolution time into k kernels of length τ , each of which is composed of local dynamical decoupling and free evolution. In this way, as long as τ is shorter than the correlation time of any time-dependent noise affecting the system, the different M ensembles (indexed by $m = 0, \dots, M-1$) can accumulate phase in a correlated manner over the interleaved Ramsey interrogation periods.

inserting local $\hat{X}(\pi)$ pulses (using techniques shown in Fig. 2c) during evolution at time $T/4$ for the second-fastest ensemble and time $3T/8$ for the slowest ensemble. Each ensemble is then further subdivided into two sub-ensembles for dual-quadrature readout (Fig. 4b). Resultant Ramsey oscillations versus the total evolution time T show a frequency ratio of 1.00:1.99(1):4.10(4), very close to the desired 1:2:4 ratio.

Following this experimental demonstration, we now discuss two limitations (and possible solutions) of this scheme, specifically related to the frequency noise profile and the atom number per ensemble. First, for the simplest case of shot-to-shot noise of laser frequencies that are otherwise constant during the interrogation, our scheme would allow the clock stability to be improved exponentially¹² by a factor of $\sqrt{2^{M-1}/M}$; the factor of $\sqrt{1/M}$ stems from increased QPN in the ensemble used for phase estimation and assumes that the total number of atoms is uniformly distributed across the M ensembles. However, for more general time-dependent frequency noise, the situation is more complex, requiring a higher-order pulse sequence¹². We propose one such pulse sequence in Fig. 4c, by breaking the total

evolution time T into multiple kernels of length τ . Within each kernel, each ensemble experiences a combination of local dynamical decoupling and free evolution, such that the net phase evolution time is $T, T/2, \dots, T/2^{M-1}$. This scheme could handle noise profiles where the local phase accumulation period τ is shorter than the correlation time of noise. We numerically find that exponential scaling of the maximum interrogation time is then possible up to a saturation point set by the effective decoupling bandwidth (Extended Data Fig. 2).

Second, multi-ensemble estimation schemes in general require sufficient atom number per ensemble to be useful¹². When the number of atoms per ensemble is limited, QPN can negate any advantage, i.e. when the error probability in estimating a phase slip by using a slower-evolving ensemble exceeds the actual phase-slip probability in the fastest-evolving ensemble. For the present experimental demonstration with $N \approx 6$ per ensemble, we do not expect a metrological gain (Extended Data Fig. 2), but we note that a generalization of our addressing scheme to two-dimensional tweezer clock systems¹⁹ is straightforward. For example, we imagine a realistic scenario of a

10×20 atom array with column-by-column control of the tweezer positions, such that it could be generated with crossed AODs or an AOD combined with a spatial light modulator. In this case, each pair of columns could realize one ensemble with dual-quadrature readout. Finally, we note that such exponential scaling is possible only up to a timescale where decoherence is dominated by local oscillator noise. Beyond that, the interrogation time will be limited by atomic coherence and ultimately by atomic state lifetime⁴².

In summary, we have demonstrated arbitrary local rotations for optical transitions through robust phase-sensitive position control in neutral-atom arrays, with sub-diffraction-limited precision. We have used such rotations to simultaneously interrogate two atomic ensembles for dual-quadrature readout of a Ramsey interferometry signal with demonstrable metrological gain, and have shown a proof of principle for controlling many ensembles with variable sensitivity during Ramsey evolution, a key ingredient of proposals for optimal clocks. Further, these methods could be naturally combined with metrologically useful entangled states^{16,39,43} to simultaneously enable high sensitivity with a large dynamic range. More generally, our results are an important step towards a fully programmable quantum optical clock based on neutral atoms, which would incorporate quantum computing techniques towards metrological gains, similar to work done with ion-trap devices^{43,44} but likely in a more scalable fashion. Such a universal neutral-atom clock system would ideally combine arbitrary local rotations, as shown here, with two-qubit entangling operations for optical transitions⁴⁵, and mid-circuit readout and reset, which has not been demonstrated so far.

Note added in proof: During the completion of this work, we became aware of a related work performing local $\hat{Z}$ rotations and studying entanglement-enhanced metrology in an optical tweezer-array clock experiment⁴⁶.

Online content

Any methods, additional references, Nature Portfolio reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41567-023-02323-w>.

References

- Ludlow, A. D., Boyd, M. M., Ye, J., Peik, E. & Schmidt, P. O. Optical atomic clocks. *Rev. Mod. Phys.* **87**, 637 (2015).
- Safronova, M. et al. Search for new physics with atoms and molecules. *Rev. Mod. Phys.* **90**, 25008 (2018).
- ACME Collaboration. Improved limit on the electric dipole moment of the electron. *Nature* **562**, 355–360 (2018).
- Roussy, T. S. et al. An improved bound on the electron's electric dipole moment. *Science* **381**, 46–50 (2023).
- Degen, C., Reinhard, F. & Cappellaro, P. Quantum sensing. *Rev. Mod. Phys.* **89**, 35002 (2017).
- Pezzè, L., Smerzi, A., Oberthaler, M. K., Schmied, R. & Treutlein, P. Quantum metrology with nonclassical states of atomic ensembles. *Rev. Mod. Phys.* **90**, 35005 (2018).
- Pedrozo-Peñafiel, E. et al. Entanglement on an optical atomic-clock transition. *Nature* **588**, 414–418 (2020).
- Macieszczak, K., Fraas, M. & Demkowicz-Dobrzański, R. Bayesian quantum frequency estimation in presence of collective dephasing. *New J. Phys.* **16**, 113002 (2014).
- Kaubuegger, R., Vasilyev, D. V., Schulte, M., Hammerer, K. & Zoller, P. Quantum variational optimization of Ramsey interferometry and atomic clocks. *Phys. Rev. X* **11**, 41045 (2021).
- Huelga, S. F. et al. Improvement of frequency standards with quantum entanglement. *Phys. Rev. Lett.* **79**, 3865 (1997).
- Schulte, M., Lisdat, C., Schmidt, P. O., Sterr, U. & Hammerer, K. Prospects and challenges for squeezing-enhanced optical atomic clocks. *Nat. Commun.* **11**, 5955 (2020).
- Rosenband, T. & Leibbrandt, D. R. Exponential scaling of clock stability with atom number. Preprint at <https://arxiv.org/abs/1303.6357> (2013).
- Borregaard, J. & Sørensen, A. S. Efficient atomic clocks operated with several atomic ensembles. *Phys. Rev. Lett.* **111**, 90802 (2013).
- Colombo, S., Pedrozo-Peñafiel, E. & Vuletić, V. Entanglement-enhanced optical atomic clocks. *Appl. Phys. Lett.* **121**, 210502 (2022).
- Demkowicz-Dobrzański, R., Kołodyński, J. & Guţă, M. The elusive Heisenberg limit in quantum-enhanced metrology. *Nat. Commun.* **3**, 1063 (2012).
- Li, W., Wu, S., Smerzi, A. & Pezzè, L. Improved absolute clock stability by the joint interrogation of two atomic ensembles. *Phys. Rev. A* **105**, 53116 (2022).
- Madjarov, I. S. et al. An atomic-array optical clock with single-atom readout. *Phys. Rev. X* **9**, 41052 (2019).
- Norcia, M. A. et al. Seconds-scale coherence on an optical clock transition in a tweezer array. *Science* **366**, 93–97 (2019).
- Young, A. W. et al. Half-minute-scale atomic coherence and high relative stability in a tweezer clock. *Nature* **588**, 408–413 (2020).
- Endres, M. et al. Atom-by-atom assembly of defect-free one-dimensional cold atom arrays. *Science* **354**, 1024–1027 (2016).
- Barredo, D., de Leseleuc, S., Lienhard, V., Lahaye, T. & Browaeys, A. An atom-by-atom assembler of defect-free arbitrary two-dimensional atomic arrays. *Science* **354**, 1021–1023 (2016).
- Bluvstein, D. et al. A quantum processor based on coherent transport of entangled atom arrays. *Nature* **604**, 451–456 (2022).
- Dordevic, T. et al. Entanglement transport and a nanophotonic interface for atoms in optical tweezers. *Science* **373**, 1511–1514 (2021).
- Lengwenus, A., Kruse, J., Schlosser, M., Tichelmann, S. & Birkel, G. Coherent transport of atomic quantum states in a scalable shift register. *Phys. Rev. Lett.* **105**, 170502 (2010).
- Beugnon, J. et al. Two-dimensional transport and transfer of a single atomic qubit in optical tweezers. *Nat. Phys.* **3**, 696–699 (2007).
- Schaetz, T. et al. Quantum dense coding with atomic qubits. *Phys. Rev. Lett.* **93**, 40505 (2004).
- Chen, N. et al. Analyzing the Rydberg-based optical-metastable-ground architecture for ¹⁷¹Yb nuclear spins. *Phys. Rev. A* **105**, 52438 (2022).
- Wu, Y., Kolkowitz, S., Puri, S. & Thompson, J. D. Erasure conversion for fault-tolerant quantum computing in alkaline earth Rydberg atom arrays. *Nat. Commun.* **13**, 4657 (2022).
- Cooper, A. et al. Alkaline-earth atoms in optical tweezers. *Phys. Rev. X* **8**, 41055 (2018).
- Choi, J. et al. Preparing random states and benchmarking with many-body quantum chaos. *Nature* **613**, 468–473 (2023).
- Béguin, L., Vernier, A., Chicireanu, R., Lahaye, T. & Browaeys, A. Direct measurement of the van der Waals interaction between two Rydberg atoms. *Phys. Rev. Lett.* **110**, 263201 (2013).
- Levine, H. et al. Dispersive optical systems for scalable Raman driving of hyperfine qubits. *Phys. Rev. A* **105**, 032618 (2022).
- Weitenberg, C. et al. Single-spin addressing in an atomic Mott insulator. *Nature* **471**, 319–324 (2011).
- Levine, H. et al. High-fidelity control and entanglement of Rydberg-atom qubits. *Phys. Rev. Lett.* **121**, 123603 (2018).
- Graham, T. M. et al. Multi-qubit entanglement and algorithms on a neutral-atom quantum computer. *Nature* **604**, 457–462 (2022).
- Wang, Y., Kumar, A., Wu, T.-Y. & Weiss, D. S. Single-qubit gates based on targeted phase shifts in a 3D neutral atom array. *Science* **352**, 1562–1565 (2016).

37. Bužek, V., Derka, R. & Massar, S. Optimal quantum clocks. *Phys. Rev. Lett.* **82**, 2207 (1999).
 38. Itano, W. M. et al. Quantum projection noise: population fluctuations in two-level systems. *Phys. Rev. A* **47**, 3554 (1993).
 39. Kessler, E. M., Lovchinsky, I., Sushkov, A. O. & Lukin, M. D. Quantum error correction for metrology. *Phys. Rev. Lett.* **112**, 150802 (2014).
 40. Bowden, W., Vianello, A., Hill, I. R., Schioppo, M. & Hobson, R. Improving the Q factor of an optical atomic clock using quantum nondemolition measurement. *Phys. Rev. X* **10**, 41052 (2020).
 41. Kohlhaas, R. et al. Phase locking a clock oscillator to a coherent atomic ensemble. *Phys. Rev. X* **5**, 21011 (2015).
 42. Bothwell, T. et al. Resolving the gravitational redshift across a millimetre-scale atomic sample. *Nature* **602**, 420–424 (2022).
 43. Marciniak, C. D. et al. Optimal metrology with programmable quantum sensors. *Nature* **603**, 604–609 (2022).
 44. Schmidt, P. O. et al. Spectroscopy using quantum logic. *Science* **309**, 749–752 (2005).
 45. Schine, N., Young, A. W., Eckner, W. J., Martin, M. J. & Kaufman, A. M. Long-lived Bell states in an array of optical clock qubits. *Nat. Phys.* **18**, 1067–1073 (2022).
 46. Eckner, W. J. et al. Realizing spin squeezing with Rydberg interactions in an optical clock. *Nature* **621**, 734–739 (2023).
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Methods

Light–matter interaction Hamiltonian

After applying the rotating-wave approximation, the light–matter interaction Hamiltonian considered in this work is given by⁴⁷

$$\hat{H} = \frac{\hbar|\Omega|}{2} (|0\rangle\langle 1| e^{ikx} + |1\rangle\langle 0| e^{-ikx}), \quad (2)$$

where Ω is the Rabi frequency, k is the global laser wavevector, x is the atomic position along the beam propagation axis, $|0\rangle$ is the ground state and $|1\rangle$ is the excited state. Atom displacements by Δx correspond to phase shifts of $\phi = k\Delta x$, as described in the main text.

Note that the choice of the phase for the initial $\hat{X}(\pi/2)$ rotation is a local gauge freedom, and thus all atoms in the array can be said to experience rotation about the same local axis, for example the x axis, despite the spacing between atoms generically not being perfectly commensurate with the driving wavelength. When the atom is shifted, it can be thought of as changing $\hat{X} \rightarrow \hat{X} \cos(\phi) + i\hat{Y} \sin(\phi)$. For the case of only a single global $\hat{X}$ rotation after the movement, this is equivalent to an effective $\hat{Z}(\phi)$ rotation of the quantum state; however, in general, if multiple global $\hat{X}$ operations are performed, the equivalence with an effective $\hat{Z}$ rotation breaks down.

Data analysis

Discrimination between $|0\rangle = |^1S_0\rangle$ and $|1\rangle = |^3P_0\rangle$ is performed by strongly driving the $^1S_0 \leftrightarrow ^1P_1$ transition for 10 μ s, which heats and ejects all atoms in 1S_0 (details on the strontium level structure are provided elsewhere^{29,48}). Atoms in 3P_0 are then pumped back into 1S_0 , and imaged with a lower power on the $^1S_0 \leftrightarrow ^1P_1$ transition; imaging is performed within 120 ms.

For all the data in the main text, we interleave³⁰ data-taking with feedback to the global $|0\rangle \leftrightarrow |1\rangle$ drive frequency every ~ 4 min to counteract slow (~ 40 min period) oscillations arising from environmental drifts. 1σ error bars in the main text are typically smaller than the marker size in all figures; this includes Figs. 1d–f, 3c and 4b.

Operation fidelities

Error modelling suggests that the global $X(\pi)$ fidelity of 0.9956(1) is primarily limited by measurement errors (see below), finite temperature (the average motional occupation of atoms along the radial axis is $\bar{n} \approx 0.2$, leading to an infidelity of 2×10^{-3}) and frequency noise on our laser; the latter of these, we believe, is also the dominant limitation to our Ramsey coherence time (Fig. 3c). The Rabi frequency is ~ 2.5 kHz for all measurements in this work, which allows for fast operations compared to the timescale of decay from 3P_0 (~ 550 ms (ref. 49)). Measurement errors are dominated by the vacuum-limited atom survival during imaging (0.9995(4)), the imaging fidelity for detecting the presence of an atom (0.9997(2)) and the likelihood of ejecting atoms in 1S_0 from the tweezers to perform state discrimination (0.9967(5)) before final imaging. We note that not all of these measurement errors contribute equally, and their relative importance generically depends on the amount of excited state population in the measured state.

The wavelength of the oscillation λ_{osc} (Fig. 1d) is found from fitting the excited state population of the shifted atoms with a sinusoid of the form $A \sin(2\pi \Delta x / \lambda_{\text{osc}}) + B$, from which we determine the quoted value of $\lambda_{\text{osc}} = 699(1)$ nm. Per our independent calibration of distances within the array, we expect there is an additional ~ 5 nm of systematic uncertainty on this measurement. Further systematic uncertainty could arise if the beam propagation is not perfectly coaxial with the array. The crosstalk fidelity of 0.1(2)% is found by fitting the unshifted-atom excited state populations (Fig. 1d) with a sinusoid of the same period as was determined for the shifted atoms; the quoted crosstalk is then the amplitude A of this sinusoid. We note that we have also checked for any residual linear phase shifts by repeating this experiment with

a global phase shift of $\pi/2$ on the final Ramsey pulse, which yields linear sensitivity to small phase shifts. We find the phase of unshifted atoms to still be consistent with zero over a range of more than one wavelength.

To determine the fidelity of arbitrary local rotations (Fig. 2c), we perform quantum state tomography by reading out the produced states in the x , y and z bases by rotating the state with a global $\pi/2$ pulse of a given global laser phase as necessary. The fidelity is then estimated as $F = \langle \psi_{\text{target}} | \rho | \psi_{\text{target}} \rangle$, where ρ is the experimental state determined by quantum tomography. Due to the choice of the six arbitrary rotations, the fidelity estimation coincides with the population of the excited state or ground state along the corresponding axis. For instance, the fidelity of $|+Y\rangle$ is determined by the population of the excited state when the prepared state is rotated and measured in the y basis. To access the intrinsic infidelity induced by arbitrary local rotations, we extract SPAM errors and correct them. SPAM sources are dominated by the same detection infidelities as for the global $\hat{X}(\pi/2)$ fidelity (see above), and the finite SPAM-corrected readout $\pi/2$ pulse fidelity of 0.9982(4).

Estimating prior width of the laser phase

Here we detail how we find and isolate the prior width of the laser phase distribution from the experimentally measured values. We note that while the phase-slip probability depends only on this prior width, the measured distribution is further affected by QPN. The QPN itself is a function of the fraction of excited atoms measured in the Ramsey sequence. One thus finds that the relative contribution of the QPN term varies with the central phase of the laser and the prior width of the phase distribution. Assuming a system of $N = 20$ atoms, we plot the calculated phase distribution width as a function of central laser phase $\bar{\theta}$, but with the underlying prior phase distribution having a width of zero (Extended Data Fig. 1a). We repeat this calculation for both single- and dual-quadrature phase estimation. We note that these estimators are affected slightly differently by QPN. Specifically, while the optimal single-quadrature working point in terms of minimal QPN is around a mean phase of 0 (which corresponds to measurement with an excitation fraction of 0.5), we find that the optimal working point for a dual-quadrature estimation is around a mean phase of $\pi/4$ (corresponding to the two quadratures having excitation fractions of 0.85), although we note that the dual-quadrature value is relatively flat over the entire bandwidth. For longer interrogation times, the prior width grows as a power law, which depends on the laser noise spectrum. Thus, the contribution from projection noise is, in general, time dependent.

To isolate this effect and learn the true laser phase distribution as a function of interrogation time, we first calculate the total observable width including QPN, σ_{tot} , for a range of prior widths σ_δ at a given laser central phase $\bar{\theta}$. This is done by sampling random phases from a normal distribution, followed by sampling the observed phase from a binomial random process representing the projection uncertainty. Repeating this process over a million draws, we obtain the observed distribution as a function of prior width (Extended Data Fig. 1b). We then invert the function to obtain $\sigma_\delta(\sigma_{\text{tot}})$ and interpolate the latter to find the prior width at the given measured σ_{tot} .

Fitting the phase-slip probability

To find the phase deviation from the mean phase for a given shot, we fit the Ramsey oscillations (Fig. 3c) with a decaying sinusoid, and at each time we define the fitted mean populations as $\bar{P}^{(x)}(t)$ and $\bar{P}^{(y)}(t)$. These populations are inverted via equation (1) into the mean phase $\bar{\theta}(t)$, and finally we calculate the phase deviation from the mean as $\delta_j(t) = \text{mod}(\theta_j(t) - \bar{\theta}(t), \pi)$.

For the phase-slip probability (Fig. 3e), we fit the probability densities $\mathcal{P}(\delta_j(t))$ with $G(\mathcal{P})$, where G is a Gaussian distribution folded into the range $[-\pi, \pi]$. This fit provides an estimation of the true standard

deviation σ of the $\delta_j(t)$ distribution. With this in hand, for a given half-dynamic range B , we then find the phase-slip probability ϵ as

$$\epsilon = 2 \int_B^\infty G(\mathcal{P}) d\mathcal{P} = \operatorname{erfc}\left(\frac{B}{\sqrt{2}\sigma}\right), \quad (3)$$

where erfc is the complementary error function. Note that here $B = \pi/2$ corresponds to a single-basis measurement whereas $B = \pi$ corresponds to a dual-quadrature measurement.

To calculate the maximum interrogation time, we first find the prior width of the laser phase (as described earlier) and then fit the time-resolved profile of $\sigma(t)$, as $\sigma(t) = \beta t^\alpha$. We find $\beta = \pi \times 0.119(6)$ and $\alpha = 0.56(2)$. The growth of σ over time can then be used to predict the Ramsey decay envelope C for different choices of B , as $C = e^{-\sigma^2/2}$. In Fig. 3c, we show this envelope estimation for the cases of $B = \pi/2$ and $B = \pi$ (orange and green dashed lines, respectively).

We then analytically calculate the maximum interrogation time at a fixed phase-slip error probability ϵ from equation (3) as

$$T_{\max}(\epsilon) = \left(B/(\sqrt{2}\beta\operatorname{erfc}^{-1}(\epsilon))\right)^{1/\alpha}. \quad (4)$$

Limits in multi-ensemble metrology

To study the possible limitations of the multi-ensemble scheme, we simulate stochastic phase evolution of a local oscillator with $1/f$ frequency noise, whose overall power sets a characteristic single-ensemble $1/e$ Ramsey coherence time T_{10} . We numerically find the maximum interrogation time at a fixed phase-slip probability ϵ (here we use $\epsilon = 5 \times 10^{-3}$) with increasing ensemble number M by iteratively correcting for phase slips, as described elsewhere¹². We repeat this calculation for different dynamical decoupling block lengths τ . In Extended Data Fig. 2a, we plot the results for up to $M = 9$ ensembles, assuming infinite atom number. We find that adding more ensembles indeed enables exponential scaling of the interrogation time up to a saturation point set by the effective dynamical decoupling bandwidth (expressed in terms of τ/T_{10}).

We further study the effect of QPN on the efficacy of the scheme in the case of low atom number per ensemble. For the optimal dynamical decoupling sequence found previously, we vary the number of atoms per ensemble N and repeat the calculation, which is now affected by QPN. Specifically, the use of slower-evolving ensembles with limited atom number for the iterative correction of phase slips in the fastest-evolving ensembles is prone to errors due to the increased variance in such estimation. For a small number of atoms per ensemble, this negates any advantage. However, we note that $N \simeq 20$ atoms per ensemble suffice for efficient operation of the scheme, with the interrogation times and noise strength tested here.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Code availability

The codes used for analysis of all figures are available from the corresponding author upon reasonable request.

References

47. Steck, D. A. Quantum and atom optics; <http://steck.us/teaching> (2007).
48. Stellmer, S., Schreck, F. & Killian, T. C. Degenerate quantum gases of strontium. *Annu. Rev. Cold At. Mol.* **1**, 1–80 (2014).
49. Shaw, A. L. et al. Dark-state enhanced loading of an optical tweezer array. *Phys. Rev. Lett.* **130**, 193402 (2023).

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Author contributions

A.L.S., R.F. and M.E. conceived the idea and experiment. A.L.S., R.F., R.B.-S.T. and J.C. performed the experiments, data analysis and numerical simulations. A.L.S., R.F., R.B.-S.T., P.S., T.H.Y. and J.C. contributed to the experimental setup. A.L.S., R.F. and M.E. wrote the manuscript with input from all authors. T.H.Y. and M.E. supervised this project.

Competing interests

The authors declare no competing interests.

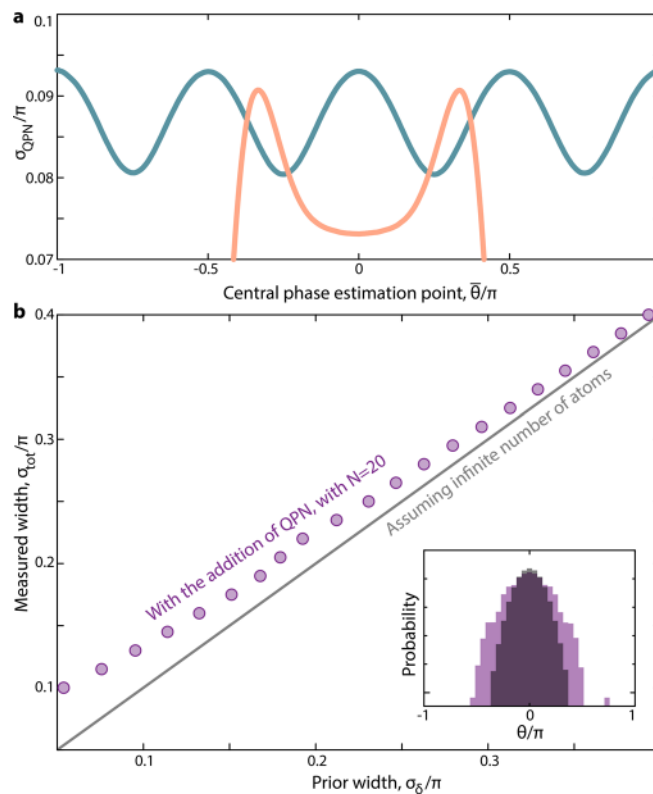
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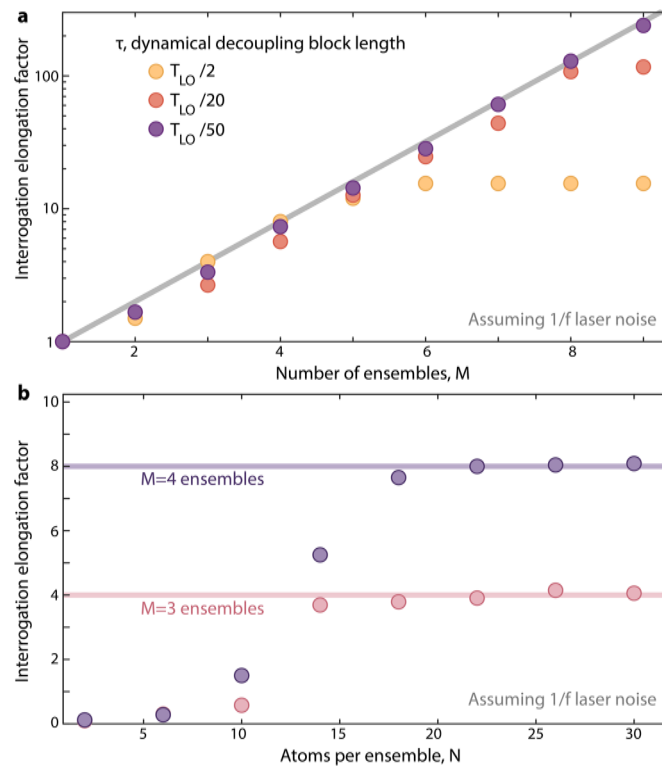
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Extended Data Fig. 1 | Quantum projection noise in a dual-quadrature measurement.

a. Added standard deviation due to quantum projection noise (QPN) for phase estimation around different average phases $\bar{\theta}$, plotted for $N = 20$ atoms utilized in a single-quadrature (orange) or dual-quadrature (green, 10 atoms per quadrature) measurement. The added QPN varies with the phase the measurement is taken at; thus as the prior width of the phase distribution grows over time, and a broader range of phases is sampled, the QPN will vary. **b.** To learn

the prior width from the measured width we sample random phases from a normal distribution, followed by sampling the observed phase from a binomial random process representing the projection uncertainty (inset). We use the sampled distributions for the dual-quadrature estimator to calculate the width including QPN for a range of prior laser widths. We then invert this function and interpolate if needed, to find the prior width for any measured width.



Extended Data Fig. 2 | Limits in multi-ensemble metrology. a, Asymptotic scaling of the extended interrogation time factor as a function of the number of ensembles M employed. We numerically calculate the maximal interrogation time at a fixed phase slip probability for different dynamical decoupling block lengths τ , in terms of the local-oscillator coherence time T_{LO} , assuming $1/f$ frequency noise and infinite atom number. The addition of more ensembles enables exponential scaling of the maximal interrogation time (solid line marks 2^{M-1}) up to a saturation point set by the effective decoupling bandwidth.

The latter can be extended by reducing the block length while maintaining a sufficiently high Rabi frequency with respect to the fast noise frequency. **b**, For the optimal decoupling sequence found in **a**, we plot the extended interrogation time as a function of the number of atoms per ensemble N . For a small number of atoms per ensemble, quantum projection noise results in an enhanced rate of false positive indication of a phase slip, negating any advantage. We find that $N \approx 20$ atoms per ensemble suffice for efficient operation.