



# Composite picosecond control of atomic state through a nanofiber interface

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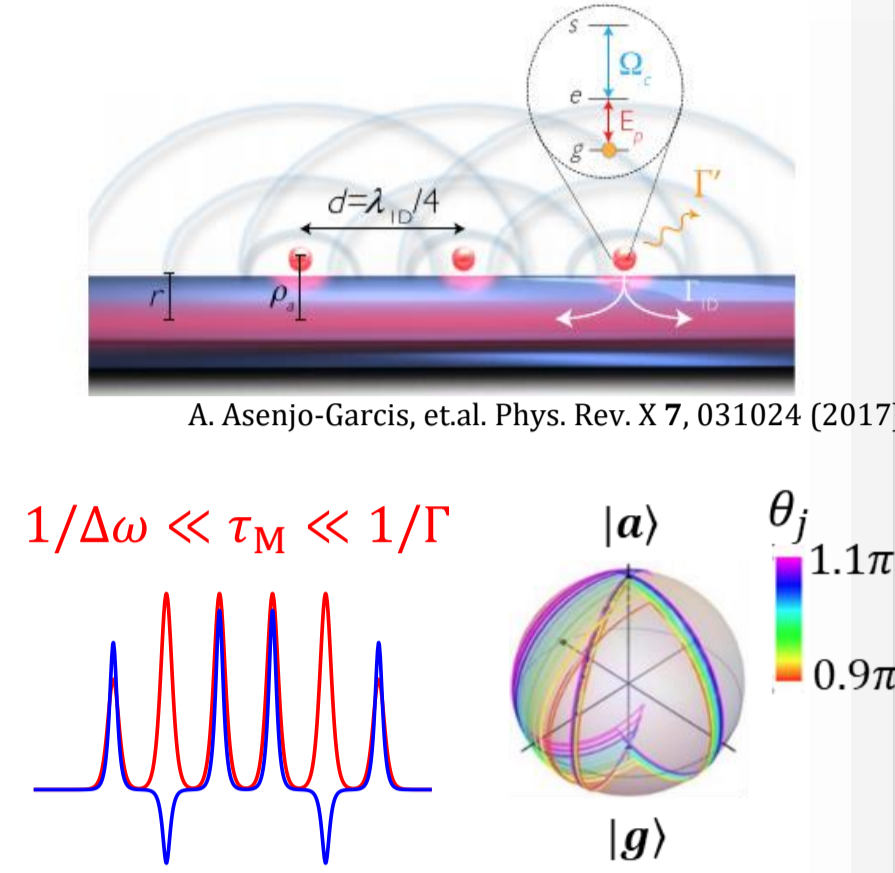
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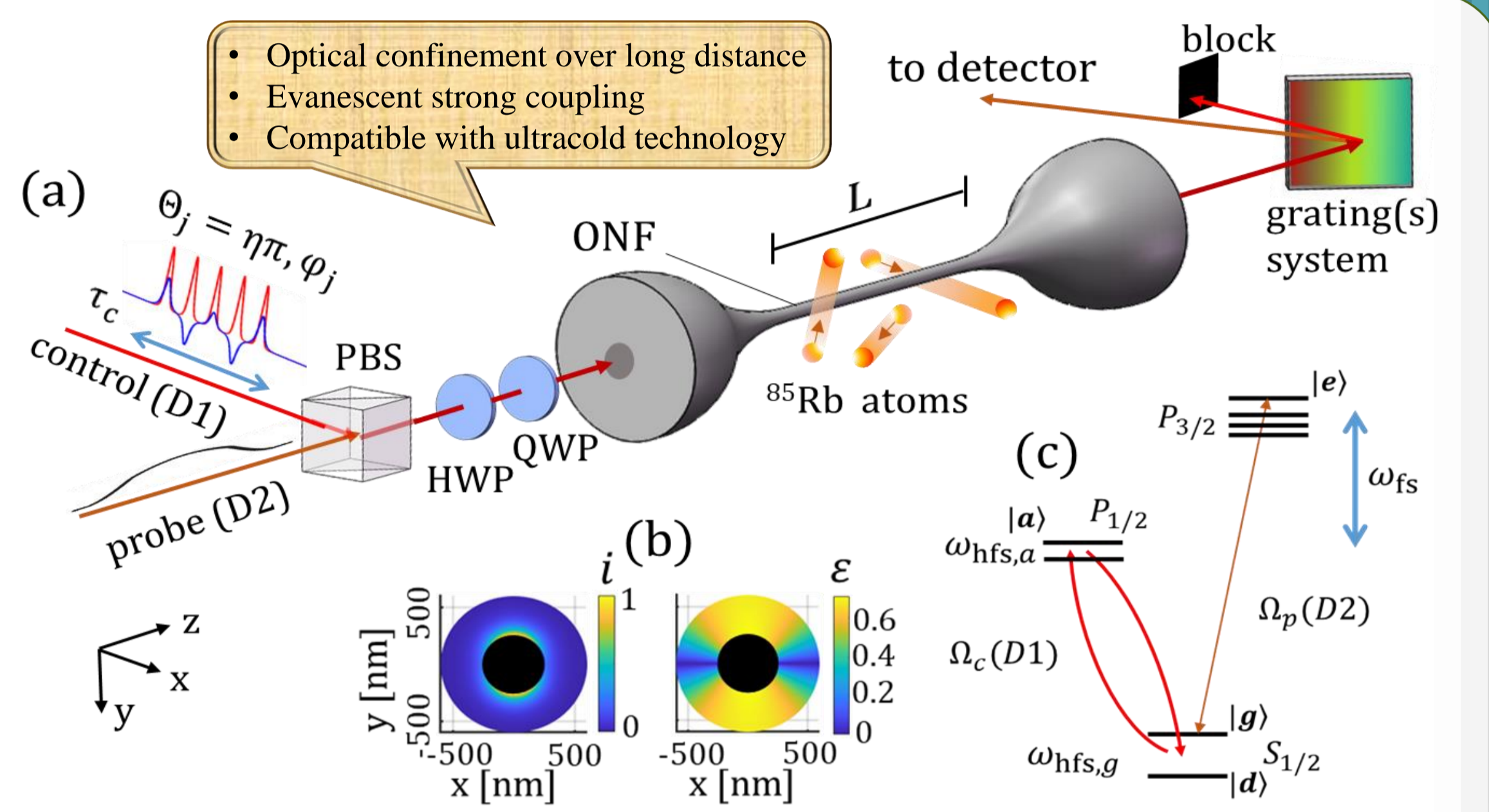
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## Introduction

- Nanophotonic interfaces: strong interaction between confined photons and nearby atoms.
- We propose to achieve high precision nanophotonic control of atomic states by composite excitation with coherent arrays of picosecond pulses.
- This work: A proof-of-concept by controlling atomic states of mesoscopic vapor at a nanofiber interface.

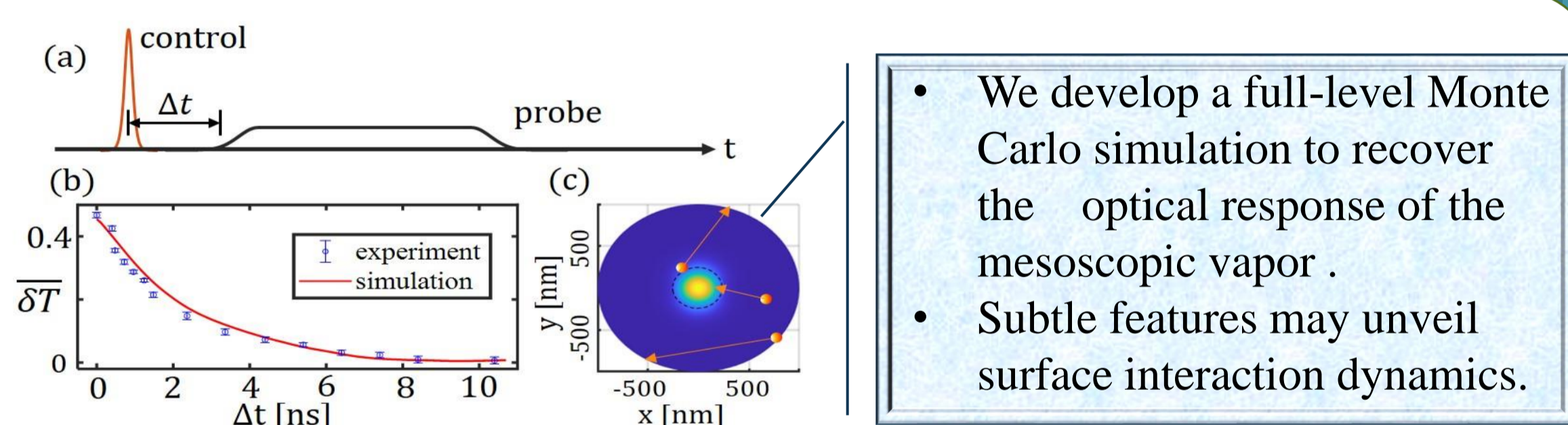


## Experimental Setup



- State-of-art atom-nanofiber interface. (L.Tong, Opt. Express, 12, 1025(2004), D.Su, New J. Phys. 21, 043053(2019)).
- Novel picosecond pulse shaping scheme for strong transition control [1].
- Composite picosecond D1 pulses to control transmission at D2.
- Controlling atoms at 300 m/s across the sub-micron evanescent field.
- Adjustable ellipticity distribution  $\varepsilon(r)$  through incident polarization control.

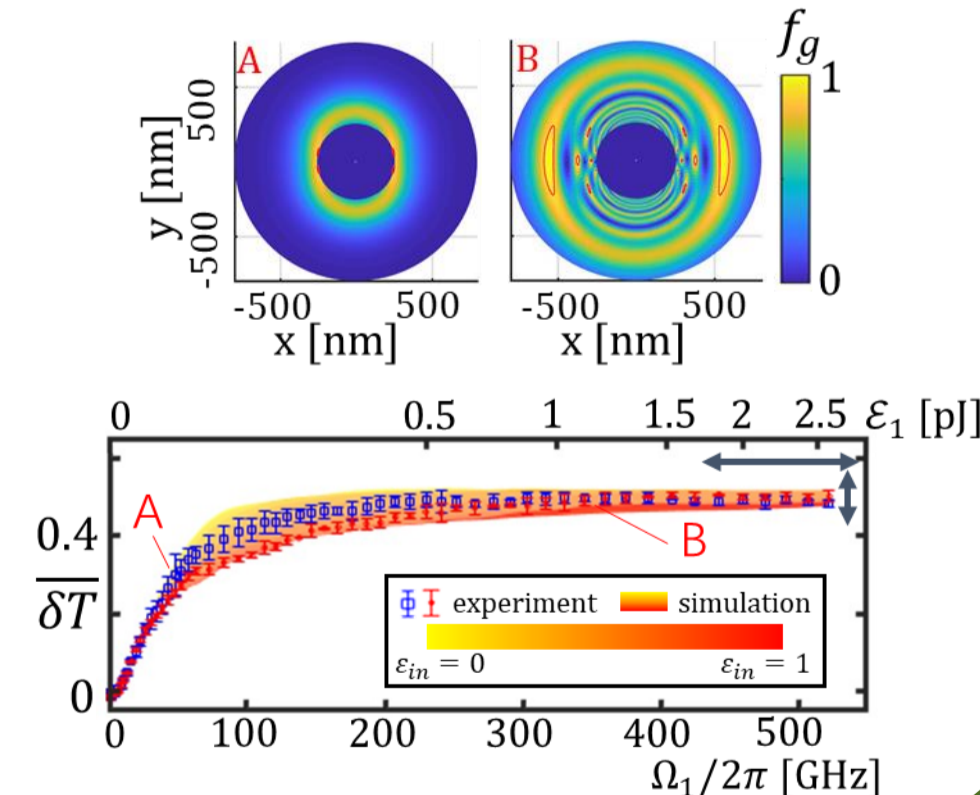
## One pulse benchmark



- We develop a full-level Monte Carlo simulation to recover the optical response of the mesoscopic vapor.
- Subtle features may unveil surface interaction dynamics.

- For N=1 picosecond control pulse, we delay the probe  $\Delta t$  to study the transient optical response of the thermal atoms (Fig. (a)). The result is shown in Fig. (b) with scatter plots. Here  $\delta T$  represents the control induced transient change of probe transmission, which reflects the D1 population inversion efficiency.

- We scan the control pulse energy  $\varepsilon_1$  with linear or circular incident polarization at  $\Delta t=0$  delay.
- Transient change of absorption  $\delta T$  up to ~45% is observed merely @ 1pJ.
- Incident helicities of control-probe matters.



## Acknowledgement & Reference

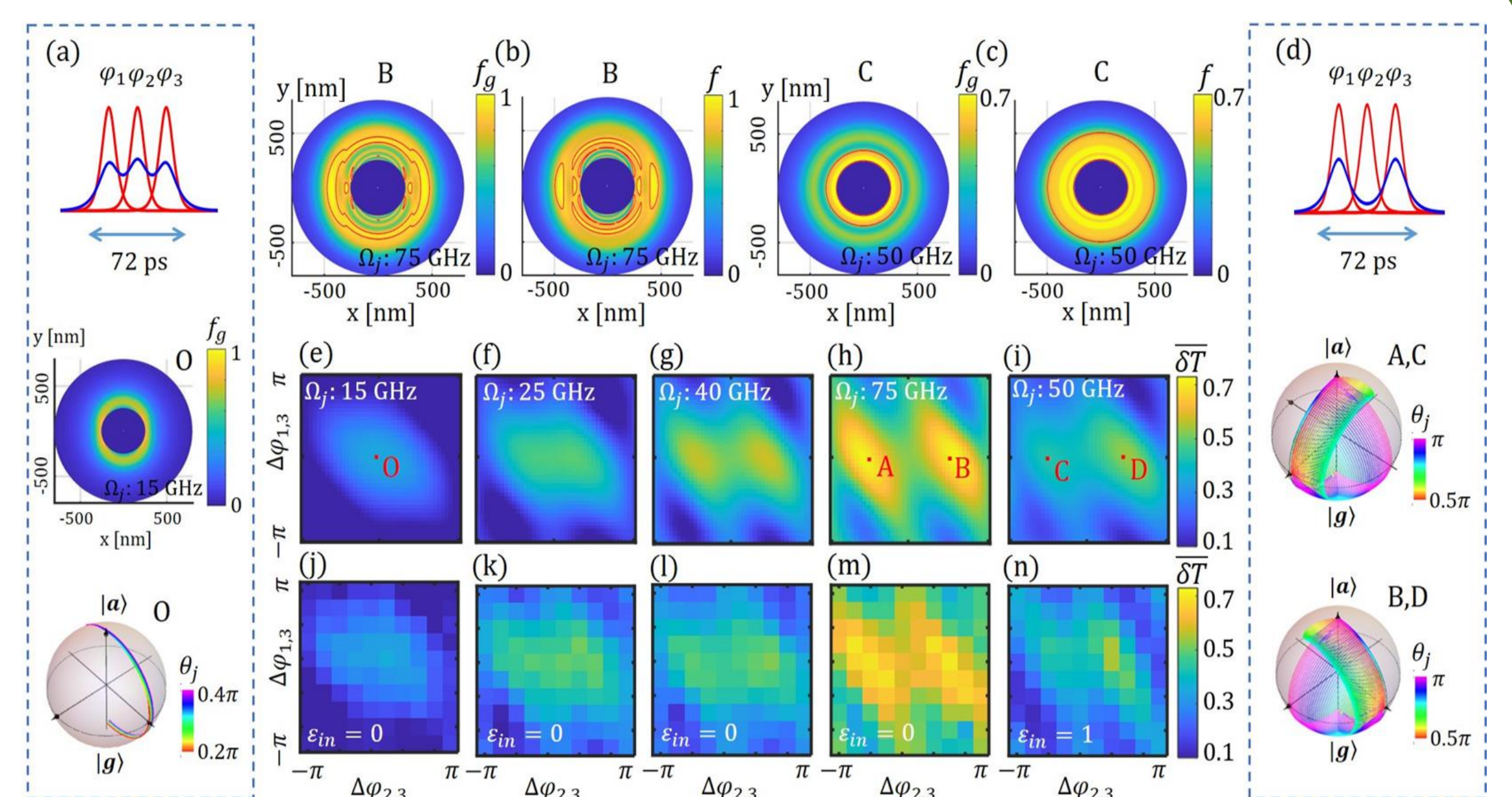
- National Key Research Program of China under Grant No. 2016YFA0302000 and No. 2017YFA0304204, National Natural Science Foundation of China under Grant No. 12074083, 61875110, 62105191, 62035013, 62075192.
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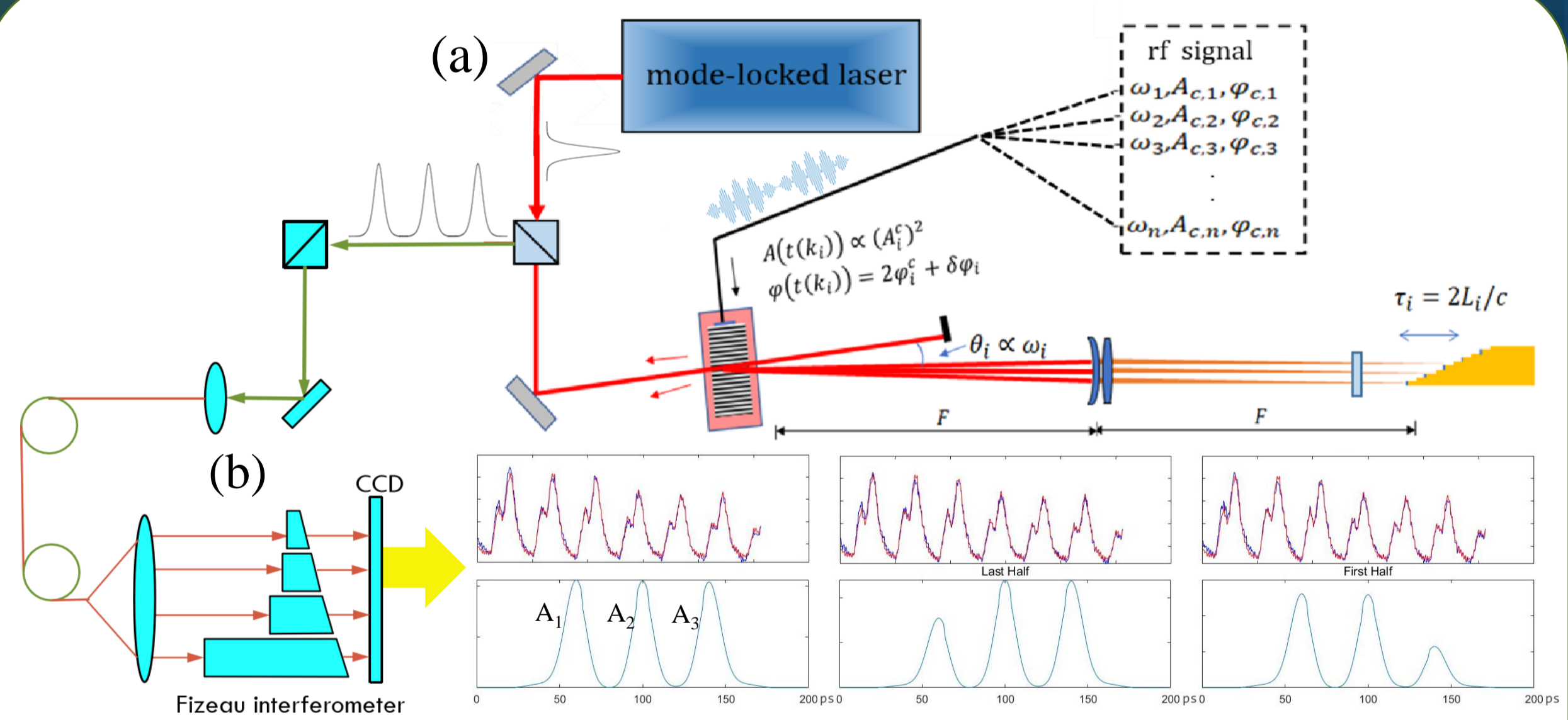
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## Composite control with three-pulse seq.



- We apply a phase-coherent N=3 pulse sequence to control the atoms in the evanescent field. Figs. (j-n) show the experimental measured  $\delta T$  at different incident polarization and power (associated with  $\Omega_j$ ).
- The experimental observations are corroborated with a full-level simulation (Figs. (e-i)).
- The broken sign symmetry in Fig.(i)(n) is associated with substantial  $|g\rangle \leftrightarrow |d\rangle$  Raman transfer, as unveiled by comparing the  $\rho_{gg}$ -depletion efficiency  $f_g(r)$  with the expected inversion efficiency  $f(r)$  in Fig. (c) according to the simulations.
- Behind the ~70% enhancement to the probe transmission is local population inversion efficiency of >90% (red curve in Fig.(b)) over a 100 nm-sized area. The control efficiency is improvable with larger N-sequences for high-fidelity manipulation of confined electric dipoles.

## Pico-array generation and measurement



Use monochromatic Fizeau interferogram to fit the pulse interferogram with sech pulse model and cosinoidal envelope correction in the frequency domain.

- (a) Picosecond pulse array is generated with a direct time-domain pulse shaping method [1].
- (b) The array is too fast to be detected in the time domain directly. Instead, we develop a model-based method to infer the array waveform through spectrum measurement, using a commercial Fizeau interferometer (Moglabs FZW600).
- (c) The Fizeau-inferred  $\phi_j$  agrees quite well with rf-controlled  $\phi_j^c$ . The deviations are likely due to nonlinear AOM transduction [1].

## Summary and outlook

- We demonstrate a composite picosecond scheme to achieve error-resilient control of a strong transition within the proximity of an optical nanofiber. A phase-coherent 3-pulse excitation efficiently invert the D1 transition of a <sup>85</sup>Rb vapor across the evanescent field, leading to an enhancement of fiber transmission of a nanosecond D2 probe by up to ~70%.
- Full-level numerical simulation suggests the composite scheme supports high fidelity control of alkaline atoms (f~99% level) through the ONF interface, robust against the near-field inhomogeneity associated with the rapidly varying intensity and polarization distributions. In future work, cyclic inversions driven by oppositely propagating guided pulses can shift the dipole spinors in k-space to control the collective interaction [4], thereby support efficient access to many-body physics in the subradiant manifold of the ONF-atom system featuring infinite-range 1D interaction [5,6].