


Stark Localization as a Resource for Weak-Field Sensing with Super-Heisenberg Precision

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Gradient fields can effectively suppress particle tunneling in a lattice and localize the wave function at all energy scales, a phenomenon known as Stark localization. Here, we show that Stark systems can be used as a probe for the precise measurement of gradient fields, particularly in the weak-field regime where most sensors do not operate optimally. In the extended phase, Stark probes achieve super-Heisenberg precision, which is well beyond most of the known quantum sensing schemes. In the localized phase, the precision drops in a universal way showing fast convergence to the thermodynamic limit. For single-particle probes, we show that quantum-enhanced sensitivity, with super-Heisenberg precision, can be achieved through a simple position measurement for all the eigenstates across the entire spectrum. For such probes, we have identified several critical exponents of the Stark localization transition and established their relationship. Thermal fluctuations, whose universal behavior is identified, reduce the precision from super-Heisenberg to Heisenberg, still outperforming classical sensors. Multiparticle interacting probes also achieve super-Heisenberg scaling in their extended phase, which shows even further enhancement near the transition point. Quantum-enhanced sensitivity is still achievable even when state preparation time is included in resource analysis.

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Introduction.—Cramér-Rao inequality lies at the foundation of estimation theory [1–3]. It bounds the precision for estimating an unknown parameter h , quantified by standard deviation δh , through $\delta h \geq 1/\sqrt{\mathcal{M}\mathcal{F}}$, where \mathcal{M} is the number of samples and \mathcal{F} is Fisher information. Generally, Fisher information scales with the probe size L as $\mathcal{F} \sim L^\beta$. While classical probes can at best achieve linear scaling, i.e., $\beta = 1$ (standard limit), quantum features may enhance the precision to super-linear scaling with $\beta > 1$ [4–6]. Originally, certain entangled states, known as GHZ states [7], have been used for achieving $\beta = 2$ (Heisenberg limit) [8–16]. However, those schemes are prone to decoherence and are fundamentally limited to Heisenberg scaling (i.e., $\beta = 2$). Alternatively, many-body probes can achieve quantum-enhanced sensitivity by harnessing a variety of quantum features. A class of such probes exploits various forms of criticality such as first-order [17–19], second-order [20–34], dissipative [35–41], time crystals [42], and topological [43–46] phase transitions. Other quantum many-body probes rely on quantum scars [47–50] and Floquet driving [51,52] as well as adaptive [53–62], continuous [41,63–66], and sequential [67,68] measurements. There are two key open problems in many-body sensors. First, although the precision of these probes is not fundamentally bounded (i.e., no restriction on β), it is very hard to find quantum probes whose precision goes beyond Heisenberg sensitivity (i.e.,

$\beta > 2$), with few exceptions [22,28,29,69–71]. Second, sensing weak fields with such probes is challenging as, for instance, critical points usually occur at finite field values, and the precision quickly drops away from that point.

The presence of a gradient field across a lattice makes the on-site energies off resonant. Consequently, the tunneling rate is suppressed, and the wave function of the particles localizes in space. This is known as Stark localization [72] and has been exploited for inducing single-particle [73–78] and many-body localization without disorder [78–95], probing the geometry of nanostructures [96], protecting coherence [97], investigating gauge theories [98–100], and creating quantum scars [48,101–105]. Stark localization has been experimentally observed in ion traps [106], optical lattices [102,107], and superconducting simulators [108]. In the limit of large one-dimensional systems, in the absence of disorder and nonlinearity, Stark localization takes place in infinitesimal fields for both single-particle [76] and multiparticle interacting [104] cases. One may wonder whether Stark systems can achieve quantum-enhanced precision sensing. The key fact is that Stark localization transition takes place at the zero-field limit. Quantum-enhanced sensitivity in such a transition can be a breakthrough in ultraprecise sensing of weak fields, a domain in which most many-body sensors fail. In addition, since Stark localization happens across the whole spectrum it provides more thermal robustness than conventional criticality which is limited to the ground state.

In this Letter, we show that Stark probes can achieve strong super-Heisenberg scaling in a region stretched all the way in the extended phase to the transition point, for both single-particle and multiparticle interacting cases, allowing for ultraprecision weak-field sensing. In single-particle probes, one can achieve $\beta = 5.98$ for the ground state and $\beta = 4.11$ for the midspectrum eigenstates, through a simple position measurement. Moreover, we determine several critical exponents and their universal relationship and show that thermal fluctuations reduce the precision from super-Heisenberg to Heisenberg scaling, still outperforming classical sensors. In addition, interacting many-body ground states achieve super-Heisenberg scaling, with $\beta = 4.26$, which is close to the midspectrum single-particle case. This is because in the half-filling regime, the many-body ground state can be approximated by filling single-particle eigenstates up to midspectrum.

Ultimate precision limit.—Let us consider a quantum probe whose density matrix $\rho(h)$ depends on an under scrutiny parameter h . To do so, one has to perform a measurement, described by a set of projective operators $\{\Pi_i\}$, on the probe. The result is described by a classical probability distribution in which each outcome appears with the probability $p_i(h) = \text{Tr}[\Pi_i \rho(h)]$. For this measurement setup, \mathcal{F} in the Cramér-Rao inequality is called Classical Fisher Information (CFI), $\mathcal{F}_C(h) = \sum_i p_i(h) [\partial_h \ln p_i(h)]^2$ [109]. Optimizing the CFI over all possible measurement setups leads to Quantum Fisher Information (QFI), namely $\mathcal{F}_Q(h) = \max_{\{\Pi_i\}} \mathcal{F}_C(h)$, determining the ultimate precision limit achievable by a quantum probe [110]. A closed form for the QFI is obtained through fidelity susceptibility as $\chi(h) = 2[1 - \tilde{F}(h)]/\Delta h^2$, where $\Delta h \rightarrow 0$ [111]. Here, $\tilde{F}(h) = \text{Tr} \sqrt{\rho(h)^{1/2} \rho(h + \Delta h) \rho(h)^{1/2}}$ is the fidelity between $\rho(h)$ and $\rho(h + \Delta h)$. The QFI then takes the form $\mathcal{F}_Q(h) = 4\chi(h)$.

Single-particle probe.—We consider a one-dimensional probe with L sites in which one particle can tunnel to its neighbors with rate J . The probe is affected by a gradient field h which we would like to estimate. The Hamiltonian is

$$H(h) = J \sum_{i=1}^{L-1} (|i\rangle\langle i+1| + |i+1\rangle\langle i|) + h \sum_{i=1}^L i|i\rangle\langle i|. \quad (1)$$

As h/J increases, the system goes through a phase transition from an extended to a Stark localized phase [76,78,79,83,84]. The transition is dramatic and affects the entire spectrum of the system which makes it very distinct from the conventional quantum phase transition as a ground state feature [87].

For our system in Eq. (1), in the thermodynamic limit (i.e., $L \rightarrow \infty$), the localization transition for all energy levels takes place at $h = h_c = 0$ [76]. One can renormalize the energy of the system as $\varepsilon = (E - E_{\min})/(E_{\max} - E_{\min})$, with E_{\min} and E_{\max} as extremal eigenenergies of the

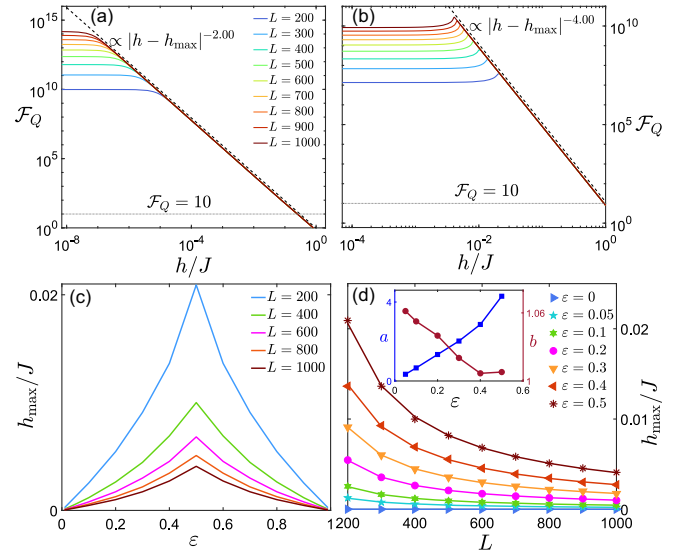


FIG. 1. The QFI versus h/J when our probe with sizes L is prepared in (a) the ground state and (b) the midspectrum eigenstate. The dashed lines in both panels are the best fit of \mathcal{F}_Q in the localized phase, showing the behavior in the thermodynamic limit. (c) The transition point h_{\max}/J versus energy ε , in various system sizes, indicating the emergence of mobility edges in finite systems. (d) The scaling of h_{\max}/J with L for various ε . Markers and solid lines represent numerical results and fitting function $h_{\max} = aL^{-b}$, respectively. Inset: a and b for different ε .

Hamiltonian $H(h)$ to fit the whole spectrum of the system within $0 \leq \varepsilon \leq 1$. For any given ε , the QFI with respect to h has been calculated for the closest eigenstate. In Figs. 1(a) and 1(b), we plot \mathcal{F}_Q as a function of h/J for various L , when our probe is in the ground state ($\varepsilon = 0$) and a midspectrum eigenstate ($\varepsilon = 0.5$), respectively. The QFI takes its maximum value at $h = h_{\max}$, which is expected to become the critical point, i.e., $h_{\max} = h_c$, in the thermodynamic limit. For the ground state [see Fig. 1(a)], the maximum takes place at vanishingly small fields, namely $h_{\max} \rightarrow 0$. For the midspectrum [see Fig. 1(b)], however, a clear peak for the QFI can be observed at a nonzero h_{\max} . Regardless of the energy levels, several features can be observed. First, by increasing L , the peak of the QFI, namely $\mathcal{F}_Q(h_{\max})$, dramatically enhances showing divergence in the thermodynamic limit. Second, the position of the peak gradually moves toward zero suggesting that in the thermodynamic limit one has $h_{\max} \rightarrow h_c = 0$. Third, despite the decay of the QFI in the localized regime, its value remains high for a large interval of h , e.g., to have $\mathcal{F}_Q \geq 10$, the gradient field can be within the range $0 \leq h/J \leq 0.25$ ($0 \leq h/J \leq 1$) for the ground (midspectrum) state. Fourth, in the localized regime, after a certain threshold the QFI becomes size independent showing divergence to the thermodynamic limit, represented by dashed lines in Figs. 1(a) and 1(b). Finite-size effects are evident in the initial plateaus of the QFI, representing the extended phase of the system. Interestingly, the dashed

lines suggest an algebraic behavior of the QFI, namely $\mathcal{F}_Q(h) \propto |h - h_{\max}|^{-\alpha}$, in the localized phase which can be perfectly fitted by $\alpha = 2.00$ for the ground state and $\alpha = 4.00$ for the midspectrum eigenstates. The fast convergence to the thermodynamic limit in the localized phase is discussed in more detail in the Supplemental Material [112].

To see how single-particle Stark localization transition depends on energy, in Fig. 1(c) we plot h_{\max} as a function of ε for various system sizes. The figure clearly indicates an energy-dependent transition which is known as mobility edge. In fact, midspectrum eigenstates are harder to localize than the eigenstates at the edges of the spectrum, a common feature that has also been observed in Anderson [113,114] and many-body [115] localization. In the thermodynamic limit, mobility edge disappears, and the transition takes place at $h_{\max} = 0$ across the whole spectrum. To see how h_{\max} decreases with an increasing system size at different energy scales, in Fig. 1(d), we plot h_{\max} versus L for energies starting from the ground state ($\varepsilon = 0$) to midspectrum ($\varepsilon = 0.5$). Our numerical simulation (denoted by markers) is well described by the fitting function $h_{\max}(L) = aL^{-b}$ (solid lines). The exponents a and b are plotted as a function of ε in the inset of Fig. 1(d). While the exponent b shows a stable behavior around $b \simeq 1$, the exponent a changes dramatically from $a \simeq 0$ (for the ground state) to $a \simeq 4$ (for the midspectrum).

Super-Heisenberg sensitivity.—In the extended phase, the QFI heavily depends on size L . To see how the QFI scales with the probe size, in Figs. 2(a) and 2(b), we plot the QFI at the transition point, i.e., $\mathcal{F}_Q(h_{\max})$, as a function of L for the ground ($\varepsilon = 0$) and the midspectrum ($\varepsilon = 0.5$) states, respectively. The QFI is shown by markers, and red lines are fitting functions $\mathcal{F}_Q(h_{\max}) \propto L^\beta$ with $\beta = 5.98$ for the ground state and $\beta = 4.11$ for the midspectrum states. This shows strong super-Heisenberg scaling in which β , by our knowledge, exceeds all other known

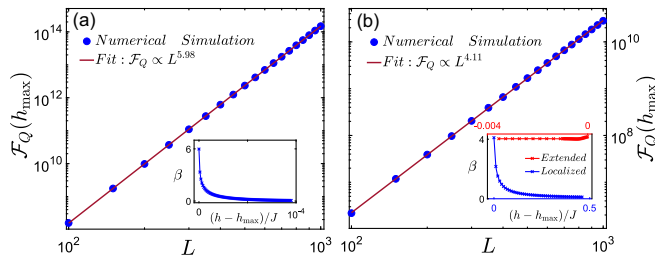


FIG. 2. The maximum of QFI (markers) as a function of L for (a) the ground state and (b) the midspectrum eigenstate. The solid red lines are fitting of the form $\mathcal{F}_Q(h_{\max}) \propto L^\beta$ with $\beta = 5.98$ for $\varepsilon = 0$ and $\beta = 4.11$ for $\varepsilon = 0.5$. The inset of each panel represents the behavior of β away from criticality. In the midspectrum (right panel), β remains almost steady in the extended phase and drops in the localized phase.

many-body probes with local interaction and bounded spectrum. We highlight these as our main results. By entering the localized regime β goes down and eventually vanishes. The smooth decay of β is depicted in the insets of Figs. 2(a) and 2(b).

Finite-size scaling.—On one side, Figs. 1(a) and 1(b) suggest that in the thermodynamic limit (i.e., $L \rightarrow \infty$) QFI behaves as $\mathcal{F}_Q \propto |h - h_{\max}|^{-\alpha}$. On the other hand, the finite-size analysis at the transition point $h = h_{\max}$ [see Figs. 2(a) and 2(b)] indicates $\mathcal{F}_Q(h = h_{\max}) \propto L^\beta$. These two behaviors suggest

$$\mathcal{F}_Q(h) \propto \frac{1}{L^{-\beta} + A|h - h_{\max}|^\alpha}, \quad (2)$$

where A is a constant. In the localized regime, where $|h - h_{\max}|^\alpha \gg L^{-\beta}$, the dependence on L becomes negligible. Thanks to the large value of β , the convergence to this limit is very rapid in our Stark probe; see the Supplemental Material [112] for more details. The algebraic behavior of QFI hints that this transition might be of the second-order type. Any second-order phase transition is accompanied by a diverging length scale as $\xi \sim |h - h_c|^{-\nu}$, where the exponent ν controls the speed of divergence. In the case of localization transition, ξ is indeed the localization length. To verify the nature of the transition and determine the critical exponents, we consider a conventional second-order finite-size scaling analysis. This implies that the QFI follows the Ansatz

$$\mathcal{F}_Q(h) = L^{\alpha/\nu} g[L^{1/\nu}(h - h_c)], \quad (3)$$

where $g(\cdot)$ is an arbitrary function. To verify the above ansatz one can plot $L^{-\alpha/\nu} \mathcal{F}_Q(h)$, versus $L^{1/\nu}(h - h_c)$ for different L . By tuning the parameters (h_c, α, ν) one tries to collapse all the curves of different system sizes on a single one. In Figs. 3(a) and 3(b), we plot the best achievable data collapse, using the PYTHON package PYFSSA [116,117] for $L = 200$ to $L = 1000$ for $\varepsilon = 0$ and $\varepsilon = 0.5$, respectively. Our careful finite-size scaling analysis results in $(h_c, \alpha, \nu) = (1.02 \times 10^{-9}, 2.00, 0.33)$ and $(h_c, \alpha, \nu) = (1.03 \times 10^{-5}, 4.00, 1.00)$, for

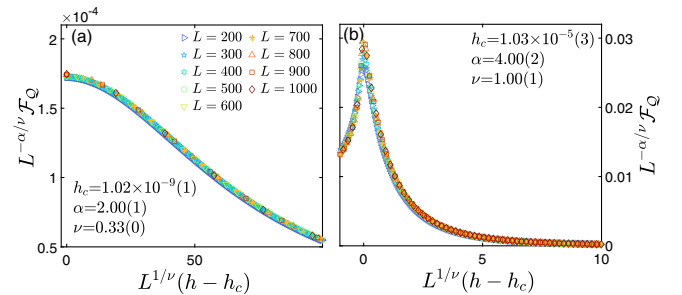


FIG. 3. Finite-size scaling analysis using Eq. (3) for (a) the ground state and (b) the midspectrum eigenstate. The optimal data collapse is obtained for the reported (h_c, α, ν) .

the ground and the midspectrum states, respectively. The values obtained for α are fully consistent with the values extracted by independent data fitting in Figs. 1(a) and 1(b).

The two Ansätze [Eqs. (2) and (3)] describe the QFI and thus cannot be independent. Indeed, by factorizing L^β from Eq. (2) one finds that these two will be the same if and only if

$$\beta = \frac{\alpha}{\nu}. \quad (4)$$

This means that α , β , and ν are not independent exponents, and two of them can describe the Stark transition. In the Supplemental Material [112], by providing the values of α , ν , and β we show the validity of Eq. (4) across the whole spectrum.

Optimal measurement.—Saturating the quantum Cramér-Rao bound generally demands complex optimal measurements which may even depend on the unknown parameter h . Therefore, finding an experimentally feasible set of measurements with precision close to the Cramér-Rao bound is highly desirable. Interestingly, in our probe, a simple position measurement described by local projective operators $\{\Pi_i = |i\rangle\langle i|\}_{i=1}^L$ saturates the QFI across the whole spectrum (see the Supplemental Material [112]).

Thermal probes.—Apart from the ground state, accessing individual eigenstates is very difficult. In practice, our probe might be described by a thermal state $\rho(h, T) = e^{-H/KT} / \text{Tr}[e^{-H/KT}]$, where T is temperature and K is the Boltzmann constant. Note that the QFI is still computed with respect to h and T is only a parameter. We consider two different scenarios with h in the extended phase, namely $h = 10^{-8}J$, and in the localized phase, namely $h = 0.05J$. The QFI, versus h , for both cases is plotted as a function of T in Figs. 4(a) and 4(b), respectively. The QFI starts with a plateau, whose width is $KT \simeq \Delta E$, where $\Delta E = E_2 - E_1$ is the energy gap. In this regime, the thermal state is described by the ground state. That is why in the extended phase [Fig. 4(a)], the plateaus are separated for each L while in the localized phase [Fig. 4(b)], they collapse. Interesting behavior can be

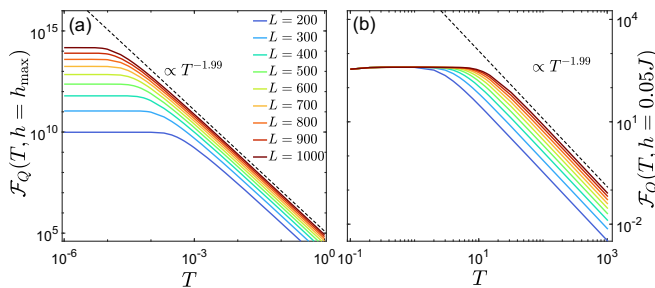


FIG. 4. The QFI of the thermal state as a function of temperature T for various system sizes, when the probe is operating at (a) the transition point $h = h_{\max} = 10^{-8}J$ and (b) the localized phase with $h = 0.05J$.

observed for $KT > \Delta E$, where the QFI decays as T increases. In this regime, as shown in Fig. 4, for both localized and extended probes, the QFI decays as $\mathcal{F}_Q \propto c(L)T^{-\mu}$, where $c(L) \propto L^\gamma$, with $\gamma = 2.0$ (see the Supplemental Material [112]) and $\mu = 1.99$ is a universal exponent. Therefore, in the limit of $KT > \Delta E$ the QFI is universally described as

$$\mathcal{F}_Q(h) \sim f(h)T^{-\mu}L^\gamma, \quad (5)$$

which shows Heisenberg scaling with respect to system size, outperforming classical sensors.

Many-body interacting probes.—In this section, we show that our probe can also operate in the case of multiparticle interacting systems. We consider a one-dimensional probe of size L with $N = L/2$ particles interacting via Hamiltonian

$$H(h) = J \sum_{i=1}^{L-1} (\sigma_i^x \sigma_{i+1}^x + \sigma_i^y \sigma_{i+1}^y + \sigma_i^z \sigma_{i+1}^z) + h \sum_{i=1}^L i \sigma_i^z, \quad (6)$$

where $\sigma_i^{x,y,z}$ are the Pauli operators and h , again, is the strength of the gradient field. To benefit from Matrix Product State (MPS) analysis for capturing large system sizes, we only focus on the ground state. In Fig. 5(a) we plot \mathcal{F}_Q as a function of h/J , in a half filling regime, for various L . We use exact diagonalization for systems up to $L = 20$ and MPS (using TeNPy Library [118]), with bond dimension $\chi = 1000$, for larger L 's. Surprisingly, the calculated QFI in the interacting many-body system behaves qualitatively similarly to the single-particle case in the midspectrum [see Fig. 1(b)]. While it shows a steady behavior in the extended phase, it fluctuates and eventually peaks at a nonzero h_{\max} . Clearly, $\mathcal{F}_Q(h_{\max})$ enhances by increasing L , signaling the divergence in the thermodynamic limit. In the localized phase, the QFI decays algebraically as $\mathcal{F}_Q \propto |h - h_{\max}|^{-\alpha}$ with $\alpha = 4.00$. Implementing careful finite-size scaling analysis results in

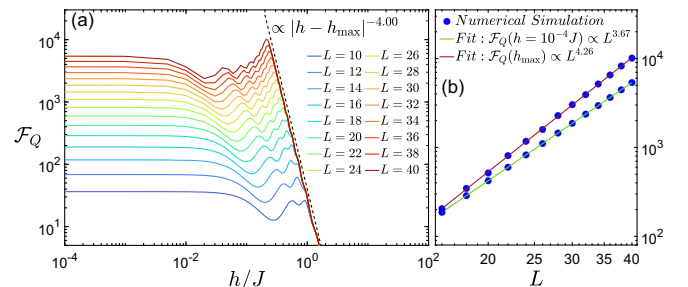


FIG. 5. (a) The QFI of a many-body interacting system with $N = L/2$ particles versus h/J when the probe of size L is in the ground state. The dashed line is the best fitting function of $|h - h_{\max}|$ in the localized phase illustrating the system's behavior in the thermodynamic limit. (b) The QFI at the extended and transition point, namely $h = h_{\max}$, as a function of L .

$(h_c, \alpha, \nu) = (1.16 \times 10^{-5}, 4.03, 1.02)$, showing an agreement with the single-particle probes prepared in the midspectrum [see Fig. 1(b)].

To see the scaling of QFI, in Fig. 5(b) we plot $\mathcal{F}_Q(h_{\max})$ as a function of L which shows algebraic scaling as $\mathcal{F}_Q(h_{\max}) \propto L^\beta$, with $\beta = 3.67$ in the extended phase which is further enhanced at the transition point to $\beta = 4.26$. The exponent, in particular near the transition point, is also close to the one from the midspectrum single-particle probes. This interesting observation can be described in a hand-waving way. The ground state of a half filling system, at least in the *noninteracting* limit, can be constructed by filling single-particle eigenstates up to midspectrum. Although the presence of interaction makes this picture less precise, it shows that the overall sensing power of the probe is dominantly defined by midspectrum single-particle eigenstates. The clear distinction between the curves in Figs. 1(b) (i.e., midspectrum of single-particle probe) and 5(a) (i.e., ground state of many-body probe) is also indicated by a small deviation between their exponent β which can be assigned to the effect of interaction. Note that the gradual movement of h_{\max} toward zero suggests that, similar to the single-particle case, the Stark many-body localization transition also takes place at infinitesimal values of the field. This is consistent with the results of Ref. [104], found for Stark localization transition in the absence of disorder and nonlinearity, and shows that many-body Stark probes also serve for weak-field sensing with super-Heisenberg precision.

Resource analysis.—If probe size L is the only resource that we care about, then the QFI is the best figure of merit for resource efficiency. However, eigenstate preparation (e.g., ground state) is time-consuming which might be considered as another resource. For instance, using adiabatic evolution for ground state initialization near criticality requires a size-dependent time which scales as $t \sim L^z$. The critical exponent z quantifies the energy gap closing at the criticality $\Delta E \sim L^{-z}$ [28]. To include time as a resource, one may consider normalized QFI \mathcal{F}_Q/t as the figure of merit [28,30,68] which turns out to be $\mathcal{F}_Q/t \sim L^{\beta-z}$. As discussed in the Supplemental Material [112], we have estimated $z \simeq 2$, resulting in $\mathcal{F}_Q/t \sim L^4$, for the single-particle and $z \simeq 0.81$, resulting in $\mathcal{F}_Q/t \sim L^{3.45}$, for the many-body probes which both still show strong quantum-enhanced precision.

Conclusion.—We have shown that Stark probes are extremely precise in measuring weak gradient fields, achieving strong super-Heisenberg precision over a region which stretches all over the extended phase to the transition point. Quantum-enhanced sensitivity is still achievable even when preparation time is included in the resource analysis. We have considered both single-particle and many-body interacting systems. In the case of single particles, we have shown that super-Heisenberg precision can be achieved through a simple position measurement

across the whole spectrum. In this regime, we have determined the critical exponents and their relationship. We have also identified the universal behavior of the probe at thermal equilibrium, which shows that as temperature increases the scaling of precision reduces from super-Heisenberg to Heisenberg. For interacting many-body ground state, while a strong super-Heisenberg precision can still be achieved, the scaling exponents become close to the midspectrum single-particle case. This can be understood as the many-body ground state can be approximately constructed by filling single-particle eigenstates until midspectrum.

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Correction: Production errors in the second citation of Ref. [104] in the second paragraph and in the author list of Ref. [22] have been set right. Page numbers in Refs. [14,31] contained errors and have been fixed.