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Quantum criticality in electron-doped BaFe_{2-x}Ni_xAs₂

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A quantum critical point is a point in a system's phase diagram at which an order is completely suppressed at absolute zero temperature (*T*). The presence of a quantum critical point manifests itself in the finite-*T* physical properties, and often gives rise to new states of matter. Superconductivity in the cuprates and in heavy fermion materials is believed by many to be mediated by fluctuations associated with a quantum critical point. In the recently discovered iron-pnictide superconductors, we report transport and NMR measurements on BaFe_{2-x}Ni_xAs₂ ($0 \le x \le 0.17$). We find two critical points at $x_{c1} = 0.10$ and $x_{c2} = 0.14$. The electrical resistivity follows $\rho = \rho_0 + AT^n$, with n = 1 around x_{c1} and another minimal n = 1.1 at x_{c2} . By NMR measurements, we identity x_{c1} to be a magnetic quantum critical point and suggest that x_{c2} is a new type of quantum critical point associated with a nematic structural phase transition. Our results suggest that the superconductivity in carrier-doped pnictides is closely linked to the quantum criticality.

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n cuprate high temperature superconductors¹ and heavy fermion² compounds, the superconductivity is accompanied by the normal-state properties deviated from a Landau-Fermi liquid. Such deviation has been ascribed to the quantum critical fluctuations associated with a quantum critical point $(OCP)^{3,4}$, whose relationship to the occurrence of superconductivity has been one of the central issues in condensed-matter physics in the last decades. In iron-pnictide high-temperature superconductors^{5,6}, searching for magnetic fluctuations has also become an important subject⁷. Quantum critical fluctuations of order parameters take place not only in spatial domain, but also in imaginary time domain^{8,9}. The correlation time τ_0 and correlation length ξ are scaled to each other, through a dynamical exponent z, $\tau_0 \propto \xi^z$. Several physical quantities, such as the electrical resistivity and spinlattice relaxation rate $(1/T_1)$, can be used to probe the quantum critical phenomena. For the quasiparticles scattering dominated by the quantum critical fluctuations, the resistivity scales as $\rho \propto T^n$. For a two-dimensional (2D) antiferromagnetic spin-density wave (SDW) QCP, the exponent n=1 is often observed^{1,2,10}. On the other hand, for a 2D order with q = (0,0) where z = 3, n = 4/3 at the QCP^{11,12}. Around a QCP, $1/T_1$ also shows a characteristic T-scaling¹¹.

BaFe_{2-x}Ni_xAs₂ is an electron-doped system¹³ where every Ni donates two electrons in contrast to Co doping that contributes only one electron¹⁴. Therefore, Ni doping suffers less from disorder which is usually harmful for a QCP to exist. In this work, we find two critical points at $x_{c1} = 0.10$ and $x_{c2} = 0.14$, respectively. By NMR measurements, we identify x_{c1} to be a magnetic QCP and suggest that x_{c2} is a QCP associated with the tetragonal-to-orthorhombic structural phase transition. The highest T_c is found around x_{c1} , which suggests that the superconductivity in the carrier-doped BaFe₂As₂ is more closely tied to the magnetic QCP, while the unusual quantum criticality associated with the structural transition deserves further investigation.

Results

Electrical resistivity measurements. Figure 1 shows the in-plane electrical resistivity data in BaFe_{2-x}Ni_xAs₂ for various *x*, which are fitted by the equation $\rho_{ab} = \rho_0 + AT^n$ (for data over the whole

temperature range, see Supplementary Fig. S1 and Supplementary Note 1). For a conventional metal described by the Fermi liquid theory, the exponent n = 2 is expected. However, we find n < 1.5



Figure 2 | The obtained phase diagram of BaFe_{2-x}**Ni**_x**As**₂. (a) The residual resistivity ρ_0 and the coefficient *A* as a function of Ni-doping *x*. (b) The obtained phase diagram of BaFe_{2-x}Ni_xAs₂. The *T*_N and θ are obtained from NMR spectra and $1/T_1T$, respectively. AF and SC denote the antiferromagnetically ordered and the superconducting states, respectively. The solid and dashed lines are the guides to the eyes. Colours in the normal state represent the evolution of the exponent *n* in the resistivity fitted by $\rho_{ab} = \rho_0 + AT^n$. The light yellow region at *x* = 0.10 shows that the resistivity is *T*-linear. The error bar for θ is the s.d. in the fitting of $1/T_1T$. The error bar for T_s represents the temperature interval in measuring the NMR spectra.



Figure 1 | The in-plane electrical resistivity. The in-plane electrical resistivity ρ_{ab} for BaFe_{2-x}Ni_xAs₂. The data in the normal state are fitted by the equation $\rho_{ab} = \rho_0 + AT^n$ (red curve).

for $0.09 \le x \le 0.14$. This is a notable feature of non-Fermi liquid behaviour. Most remarkably, a *T*-linear behaviour (n = 1) is observed for $x_{c1} = 0.10$ and persists up to T = 100 K. Another minimal n = 1.1 is found for $x_{c2} = 0.14$, which is in good agreement with a previous transport measurement¹⁵. An equally interesting feature is that both the residual resistivity ρ_0 and the coefficient *A* show a maximum at x_{c1} and x_{c2} as seen in Fig. 2a. The evolution of the exponent *n* with Ni content is shown in Fig. 2b.

NMR measurements. We use NMR to investigate the nature of x_{c1} and x_{c2} . We identify x_{c1} to be a SDW QCP and suggest that x_{c2} is a tetragonal-to-orthorhombic structural phase transition QCP. Figure 3a,b display the frequency-swept spectra of ⁷⁵As NMR for x = 0.05 and 0.07, respectively. The very narrow central transition peak above the antiferromagnetic transition temperature $T_{\rm N}$ testifies a good quality of the samples. When a magnetic order sets in, the spectra will split into two pairs as labelled in Fig. 3a, due to the development of an internal magnetic field H_{int} . For x = 0.05, we observed two split broad peaks below $T_N = 74$ K. This is due to a distribution of H_{int} , which results in a broadening of each pair of the peaks. Upon further doping, H_{int} is reduced and its distribution becomes larger, so that only one broad peak can be seen below $T_{\rm N} = 48$ K for x = 0.07. Similar broadening of the spectra was also observed previously in a very lightly-doped sample BaFe_{1.934}Ni_{0.066}As₂ (ref. 16). For both x, the spectra can be reproduced by assuming a Gaussian distribution of H_{int} as shown

by the red curves. We obtain the averaged internal field $\langle H_{\text{int}} \rangle = 0.75 \text{ T}$ at T = 15 K for x = 0.05, and $\langle H_{\text{int}} \rangle = 0.39 \text{ T}$ at T = 25 K for x = 0.07. By using a hyperfine coupling constant of 1.88 T per μ_{B} obtained in the undoped parent compound¹⁷, the averaged ordered magnetic moment $\langle S \rangle$ is deduced. As seen in Fig. 4, the ordered magnetic moment develops continuously below T_{N} , being consistent with the second-order nature of the phase transition. It saturates to $0.43 \,\mu_{\text{B}}$ for x = 0.05, and $0.24 \,\mu_{\text{B}}$ for x = 0.07, respectively. The ordered moment is smaller than that in the hole-doped Ba_{1-x}K_xFe₂As₂ (ref. 18), which is probably due to the fact that Ni goes directly into the Fe site and is more effective in suppressing the magnetic order. Upon further doping, at x = 0.09 and x = 0.10, however, no antiferromagnetic transition was found, as demonstrated in Fig. 3c, which shows no splitting or broadening ascribable to a magnetic ordering.

The onset of the magnetic order in the underdoped regime is also clearly seen in the spin-lattice relaxation. Figure 3d shows $1/T_1$ for x = 0.07, which was measured at the position indicated by the arrow in Fig. 3b, in order to avoid any influence from possible remnant paramagnetic phase, if any. As seen in Fig. 3d, a clear peak is found at $T_N = 48$ K due to a critical slowing down of the magnetic moments. Below T_N , $1/T_1$ decreases down to T_c . Most remarkably, $1/T_1$ shows a further rapid decrease below T_c , exhibiting a T^3 behaviour down to T = 5 K. Such a significant decrease of $1/T_1$ just below T_c is due to the superconducting gap opening in the antiferromagnetically ordered state. This is clear and direct evidence for a microscopic coexistence of



Figure 3 | NMR spectra with doping and temperature dependence of 1/T₁ at optimal doping. ⁷⁵As-NMR spectra obtained by sweeping the frequency at a fixed external field $H_0 = 11.998$ Tapplied along the *c* axis. (**a**) The spectra above and below $T_N = 74$ K for x = 0.05. Solid, dashed and dotted lines indicate, respectively, the position of the central transition, left- and right-satellites, when an internal magnetic field develops. (**b**) The spectra above and below $T_N = 48$ K for x = 0.07. The red curve represents the simulations by assuming a Gaussian distribution of the internal magnetic field. The black arrow indicates the position at which T_1 is measured below T_N . (**c**) The central transition line at and below T_c for x = 0.10. The spectrum shift at T = 1.89 K is due to a reduction of the Knight shift in the superconducting state. (**d**) The temperature dependence of $1/T_1$ for x = 0.07. The solid and dashed lines show the T^3 - and T-variation, respectively. The solid and dashed arrows indicate T_c and T_N , respectively. The error bar in $1/T_1$ is the s.d. in fitting the nuclear magnetization recovery curve.



Figure 4 | The temperature dependence of the internal magnetic field and the ordered moment for x = 0.07. Temperature dependence of the averaged internal magnetic field $\langle H_{int} \rangle$ (left vertical axis) and the averaged ordered moment $\langle S \rangle$ (right vertical axis) for x = 0.07. The solid and dashed arrows indicate T_c and T_N , respectively. The error bar represents the absolute maxima (minima) that can fit a spectrum.

superconductivity and antiferromagnetism, as the nuclei being measured experience an internal magnetic field yet the relaxation rate is suppressed rapidly below T_c . For x = 0.10, as mentioned already, no antiferromagnetic transition was found. Namely $T_N = 0$. This is consistent with the extrapolation of the T_N versus x relation that gives a critical point that coincides with $x_{c1} = 0.10$, which is further supported by the spin dynamics as elaborated below.

Figure 5a shows the quantity $1/T_1T$ for $0.05 \le x \le 0.14$. The $1/T_1T$ decreases with increasing T down to around 150 K, but starts to increase towards $T_{\rm N}$ or $T_{\rm c}$. The increase at low T is due to the antiferromagnetic spin fluctuation $\left(\left(\frac{1}{T_1T}\right)_{AF}\right)$, and the decrease at high T is due to an intraband effect $\left(\left(\frac{1}{T_1T} \right)_{intra} \right)$. Namely, $\frac{1}{T_1T} = \left(\frac{1}{T_1T}\right)_{AF} + \left(\frac{1}{T_1T}\right)_{intra}$. A similar behaviour was also seen in Ba(Fe_{1-x}Co_x)₂As₂ (ref. 19). We analyse the $(1/T_1T)_{AF}$ part by the self-consistent renormalization theory for a 2D itinerant electron system near a QCP¹¹, which predicts that $1/T_1T$ is proportional to the staggered magnetic susceptibility $\chi''(Q)$. As $\chi''(Q)$ follows a Curie–Weiss law¹¹, one has $\left(\frac{1}{T_1T}\right)_{AF} = \frac{a}{T+\theta}$. The intraband contribution is due to the density of state at the Fermi level, which is related to the spin Knight shift (K_s) through the Korringa relation $K_s^2 T_1 T = \text{constant}^{20}$. The Knight shift was found to follow a T-dependence of $K = K_0 + K_s \exp(-E_g/k_BT)$ as seen in Fig. 5b, where K_0 is T-independent, while the second is due to the band that sinks below the Fermi level^{21,22}. Correspondingly, we can write $\left(\frac{1}{T_1T}\right)_{\text{intra}}$ $= b + c \times$ $\exp(-2E_g/k_BT)$. The resulting θ is plotted in Fig. 2b. Note that θ is almost zero at x = 0.10, which yields a constant $1/T_1$ above T_c as seen in Fig. 6. The result of $\theta = 0$ means that the staggered magnetic susceptibility diverges at T = 0, indicating that x = 0.10is a magnetic QCP. Therefore, the exponent n=1 in the resistivity is due to the magnetic QCP (Generally speaking, scatterings due to the magnetic hot spots give a T-linear resistivity¹¹, but those by other parts of the Fermi surface will not^{23,24}. In real materials, however, there always exist some extent of impurity scatterings that can connect the magnetic hot spots



Figure 5 | The quantity 1/T₁T and the Knight shift. (a) Temperature dependence of $1/T_1T$ and (b) the Knight shift *K* for various *x*. The solid line is a fitting of $\frac{1}{T_1T} = \frac{a}{T+\theta} + b + c \times \exp(-2E_g/k_BT)$. The dashed curve is a fit to $K = K_0 + K_s \times \exp(-E_g/k_BT)$. The obtained parameters are $E_g/k_B = 620 \pm 40$, 510 ± 40 , 370 ± 30 , 365 ± 30 , 350 ± 30 and 330 ± 30 K for x = 0.05, 0.07, 0.09, 0.10, 0.12 and 0.14, respectively. The solid and dashed arrows indicate T_c and T_N , respectively. The error bar for $1/T_1T$ is the s.d. in fitting the nuclear magnetization recovery curve. The error bar for *K* was estimated by assuming that the spectrum-peak uncertainty equals the point (frequency) interval in measuring the NMR spectra.

and other parts of the Fermi surface as to restore the n = 1 behaviour at a magnetic QCP).

Next, we use ⁷⁵As NMR to study the structural phase transition. A tetragonal-to-orthorhombic structural transition was found in the parent compound²⁵, but no direct evidence for such structural transition was obtained in the doped BaFe_{2-x}Ni_xAs₂ thus far. In BaFe_{2-x}Co_xAs₂, a structural transition was detected in the low-doping region²⁶, but it is unclear how the transition temperature T_s would evolve as doping level increases. The ⁷⁵As nucleus has a nuclear quadrupole moment that couples to the electric field-gradient (EFG) V_{xx} ($\alpha = x,y,z$). Therefore, the ⁷⁵As-NMR spectrum is sensitive to a structural phase transition, as below T_s the EFG will change appreciably. Such change was indeed confirmed in the parent compounds BaFe₂As₂ (ref. 17) and LaFeAsO (ref. 27). When a magnetic field H_0 is applied in the *ab* plane, the NMR resonance frequency *f* is expressed by

$$f_{m \leftrightarrow m-1}(\phi, \eta) = f_0 + \frac{1}{2} \nu_Q \left(m - \frac{1}{2} \right) \times (\eta \cdot \cos 2\phi - 1)$$
 (1)



Figure 6 | The 1/T₁ for all the superconducting samples. The temperature dependence of the spin-lattice relaxation rate $1/T_1$ for various *x* of BaFe_{2-x}Ni_xAs₂. The solid and dashed arrows indicate T_c and T_{N_r} respectively. The error bar is the s.d. in fitting the nuclear magnetization recovery curve.

where m = 3/2, 1/2 and -1/2, φ is the angle between H_0 and the *a* axis, v_Q is the nuclear quadrupole resonance frequency, which is proportional to the EFG, and $\eta \equiv \frac{|V_{xx} - V_{yy}|}{|V_{zz}|}$. For a tetragonal crystal structure, $\eta = 0$. For an orthorhombic structure, however, the *a* axis and *b* axis are not identical, which results in an asymmetric EFG so that $\eta > 0$. Therefore, for a twined single crystal, the field configurations of $H_0||$ *a* axis ($\varphi = 0^\circ$) and $H_0||$ *b* axis ($\varphi = 90^\circ$) will give a different $f_{m \leftrightarrow m-1}(\varphi, \eta)$, leading to a splitting of a pair of the satellite peaks into two. The above argument also applies to the case of electronic nematic phase transition such as orbital ordering, as the EFG is also sensitive to a change in the occupation of the on-site electronic orbits.

As shown in Fig. 7a,b, only one pair of satellite peaks is observed at high temperature for both x = 0.05 and 0.07. Below certain temperature, T_{s} , however, we observed a splitting of the satellite peaks. This is strong microscopic evidence for a structural transition occurring in the underdoped samples, where the NMR spectra split owing to the formation of the twinned orthorhombic domains. In fact, each satellite peak can be well reproduced by assuming two split peaks. The obtained $T_s = 90$ K for x = 0.05 and $T_{\rm s} = 70 \,\text{K}$ for x = 0.07 agrees well with the resistivity data where an upturn at T_s is observed, see Fig. 8. This feature in the spectra persists to the higher dopings x = 0.10 and 0.12. For the optimal doping x = 0.10, the spectrum is nearly unchanged for $40 \text{ K} \le T \le 60 \text{ K}$, but suddenly changes at 20 K. This indicates that a structural transition takes place below $T_s = \sim 40$ K for this doping composition. For x = 0.12, the two satellite peaks do not change at high temperatures, but shift to the opposite direction below T = 10 K. Each broadened peak at T = 6 K can be well fitted by a superposition of two peaks. This is the first observation that a structural phase transition takes place below T_c . For x = 0.14, however, no broadening of the spectra was found down to T = 4.5 K; the spectrum shift to the same direction is due to a reduction of the Knight shift in the superconducting state. This indicates that $T_s = 0$ around $x_{c2} = 0.14$. The T_s results obtained by NMR are summarized in the phase diagram as shown in Fig. 2b.

Therefore, the minimal n = 1.1 of the resistivity exponent around x_{c2} implies quantum critical fluctuations associated with the structural QCP. We hope that our work will stimulate more experimental measurements such as elastic constant, which is also sensitive to such critical fluctuations²⁸. It should be emphasized



Figure 7 | The satellite peaks corresponding to the $\frac{1}{2} \leftarrow \rightarrow \frac{3}{2}$ and $-\frac{3}{2} \leftarrow -\frac{1}{2}$ **transitions. (a-d)** The satellite peaks with the external field H_0 applied along the *a* axis or *b* axis above and below T_s for x = 0.05, 0.07, 0.10 and 0.12. The peaks split below T_s due to a change in EFG. All the spectra are fitted by a single Gaussian function above T_s , but by two Gaussian functions below T_s . (e) The satellite peaks for x = 0.14, which only shift to the same direction due to a reduction of the Knight shift as in Fig. 3c.

here that the exponent n = 1.1 is smaller than n = 4/3 expected for the fluctuations dominated by q = (0,0) (ref. 12). Our result suggests that the fluctuation associated with x_{c2} is local (namely, all wave vectors contribute to the quasiparticles scattering), which would lead to an n = 1 (ref. 29).

Discussion

The mechanism for the superconductivity in iron-pnictides has been discussed in relation to magnetic fluctuations^{30–32} as well as structural/orbital fluctuations³³. In this context, it is worthwhile



Figure 8 | The enlarged part of the electrical resistivity and its derivative for the underdoped samples. (a) Enlarged part of the in-plane electrical resistivity ρ_{ab} (right vertical axis) and its derivative (left vertical axis) for x = 0.05. (b) Data for x = 0.07. The solid and dashed arrows indicate the structural phase transition temperature T_s and the antiferromagnetic transition temperature T_N determined by NMR, respectively.

noting that T_c is the highest around x_{c1} . This suggests that the superconductivity in $BaFe_{2-x}Ni_xAs_2$ is more closely related to the antiferromagnetic QCP rather than the structural QCP. Also, previous results suggestive of a magnetic QCP was reported for BaFe₂(As_{1 - x}P_x)₂ (refs 34,35), but isovalent P substitution for As does not add carriers there. In the present case, Ni directly donates electrons into the system (chemical doping), so the tuning parameter is totally different here. Thus, our work demonstrates that $BaFe_{2-x}Ni_xAs_2$ is a new material that provides a unique opportunity to study the issues of quantum criticality. In particular, our result suggests that x_{c2} is a new type of QCP at which the exponent n cannot be explained by existing theories. Although the importance of the tetragonal-toorthorhombic structural transition, which is often associated with electronic nematicity^{36,37}, has been pointed out in the pnictides^{33,38,39}, the physics of a QCP associated with it is a much less-explored frontier, which deserves more investigations in the future.

Methods

Sample preparation and characterization. The single crystal samples of $BaFe_{2-x}Ni_xAs_2$ used for the measurements were grown by the self-flux method⁴⁰. Here the Ni content *x* was determined by energy-dispersive x-ray spectroscopy. The T_c was determined by DC susceptibility measured by a superconducting quantum interference device with the applied field 50 Oe parallel to the *ab* plane. The T_c is 3.5, 14, 18.5, 18.2, 16.8 and 13.1 K for x = 0.05, 0.07, 0.09, 0.10, 0.12 and 0.14, respectively.

The resistivity measurements. Resistivity measurements were performed in quantum design physical properties measurement system by a standard dc four-probe method. Here we have used the same single crystals used in the NMR measurements. The electrical resistance was measured upon both warming and cooling processes in order to ensure no temperature effect from the electrodes on the samples. Both the warming and the cooling speed is 2 K min^{-1} .

Measurements of NMR spectra and T_1 **.** The NMR spectra were obtained by integrating the spin echo as a function of the RF frequency at a constant external magnetic field $H_0 = 11.998$ T. The nucleus ⁷⁵As has a nuclear spin I = 3/2 and the nuclear spin Hamiltonian can be expressed as

$$H = \gamma h I (H_0 + H_{\text{int}}) + \frac{h v_{\text{Q}}}{6} \left[3I_Z^2 - I(I+1) + \frac{1}{2} \eta \left(I_+^2 + I_-^2 \right) \right]$$
(2)

where $\gamma/2\pi = 7.9219 \text{ MHz T}^{-1}$ is the gyromagnetic ratio of ⁷⁵As, *h* is Planck constant, H_{int} is the internal magnetic field at the As nuclear spin site resulting from the hyperfine coupling to the neighbouring Fe electrons. The T_1 was determined by using the saturation-recovery method and the nuclear magnetization is fitted to $1 - \frac{M(t)}{M(\infty)} = 0.9 \exp^{-6t/T_1} + 0.1 \exp^{-t/T_1}$, where

M(t) is the nuclear magnetization at time t after the saturation pulse⁴¹. The curve is fitted very well and T_1 is of single component.

Simulation of the spectra. In order to reproduce the spectra, we assume that the distribution of the internal magnetic field is Gaussian as,

$$I(f) = \exp\left(-\frac{\left\{f - \left[\gamma(H_0 \pm \langle H_{\rm int} \rangle) + nv_c\right]\right\}^2}{2H_{\sigma}^2}\right)$$
(3)

where I(f) is the intensity of the spectra, H_{σ} is the distribution of the internal magnetic field $H_{\rm int}$ and v_c is the nuclear quadrupole resonance frequency tensor along the *c* axis. Then the spectrum is fitted by convoluting equation 3 with a Gaussian broadening function $p(ft) = A_n \frac{1}{\sqrt{2\pi\sigma_n}} \exp\left(-\frac{f^2}{2\sigma_n^2}\right)$ that already exists above $T_{\rm N}$. For n = 0 (central transition), $A_0 = 1$ and $2\sqrt{2\ln 2\sigma_0}$ is the full-width at half maximum (FWHM) of the central peak above $T_{\rm N}$. For $n = \pm 1$ (satellites), $A \pm_1 = 3/4$ and $2\sqrt{2\ln 2\sigma} \pm_1$ is the FWHM of satellite lines above $T_{\rm N}$. The red solid curve in Fig. 3a is a fit with $H_{\sigma} = 0.37$ T, $2\sqrt{2\ln 2\sigma_0} = 0.03$ MHz and $2\sqrt{2\ln 2\sigma} \pm_1 = 0.38$ MHz, which was taken from the spectrum at T = 100 K. The red solid curve in Fig. 3b is a fit with $2\sqrt{2\ln 2\sigma_0} = 0.04$ MHz and $2\sqrt{2\ln 2\sigma} \pm_1 = 0.5$ MHz, which was taken from the spectrum at T = 50 K.

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Author contributions

The single crystals were grown by D.L.S. and C.T.L. The NMR measurements were performed by R.Z., Z.L., J.Y. and G.-q.Z. The electrical resistivity was measured by R.Z., Z.L. and J.Y. G.-q.Z. coordinated the whole work and wrote the manuscript, which was supplemented by R.Z. All authors have discussed the results and the interpretation.

Additional information

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