Three-dimensional collective charge excitations in electron-doped copper oxide superconductors

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High-temperature copper oxide superconductors consist of stacked CuO2 planes, with electronic band structures and magnetic excitations that are primarily two-dimensional1−2, but with superconducting coherence that is three-dimensional. This dichotomy highlights the importance of out-of-plane charge dynamics, which has been found to be incoherent in the normal state3−4 within the limited range of momenta accessible by optics. Here we use resonant inelastic X-ray scattering to charge dynamics across all three dimensions of the Brillouin zone. Polarization analysis of recently discovered collective excitations (modes) in electron-doped copper oxides5−7 reveals their charge origin, that is, without mixing with magnetic components5−7. The excitations disperse along both the in-plane and out-of-plane directions, revealing its three-dimensional nature. The periodicity of the out-of-plane dispersion corresponds to the distance between neighbouring CuO2 planes rather than to the crystallographic c-axis lattice constant, suggesting that the interplane Coulomb interaction is responsible for the coherent out-of-plane charge dynamics. The observed properties are hallmarks of the long-sought ‘acoustic plasmon’, which is a branch of distinct charge collective modes predicted for layered systems5−12 and argued to play a substantial part in mediating high-temperature superconductivity10−12.

The charge dynamics of systems with periodically stacked quasi-two-dimensional (2D) conducting planes are strongly affected in the presence of poorly screened interplane Coulomb interactions. In a simple layered electron gas with conducting planes separated by dielectric spacers8,9, the dispersion of plasmons (that is, the collective dynamical charge modes) changes from optical-like to acoustic-like as a function of out-of-plane momenta " (Fig. 1a) (such plasmons are referred to as ‘acoustic plasmons’ hereafter), a behaviour distinct from that in either pure 2D or isotropic 3D systems. For superconducting copper oxides, similar charge dynamics have been postulated because they consist of conducting CuO2 planes stacked along the c-axis with poor out-of-plane Coulomb screening10−12. Although plasmons have been observed in various spectroscopic studies at the Brillouin zone centre5,9,13 and also by transmission electron energy loss spectroscopy (EELS), which typically explores in-plane energy-momenta dispersions at qz = 0 (ref. 15), there is no information on its possible qz dependence. Experimental evidence of this previously undetected component and its characterization in energy and momentum can shed new light on long-standing hypotheses that connect out-of-plane charge dynamics to superconductivity. For instance, it has been proposed that 20% of the observed value of the high superconducting transition temperature To of the copper oxides can be attributed to the presence of acoustic plasmons10−12, where the large amount of energy stored in the interplane Coulomb interactions is related to the substantial energy savings associated with the high superconducting transition temperature16,17.

In this Letter, we focus our attention on the enigmatic ‘zone centre’ excitation previously discovered by Cu L-edge resonant inelastic X-ray scattering (RIXS) in the electron-doped copper oxides Nd1−xCeCuO4 (NCCO) (x = 0.15)9,10 and Sr1−yLa2−xCeCuO4 (LCCO) (x = 0.175), we resolve spectral features that are similar to those of NCCO at representative in-plane momentum transfers qx (red-shaded peak in Fig. 1b, c), suggesting the universality of this collective mode in electron-doped copper oxides. Speculation about its origin has included intra-band transitions6, collective modes of a quantum phase6, and plasmons8,11. Although the mode has been suspected to be of charge character, a definitive assessment has not been possible owing to the inability to distinguish between charge and magnetic excitations in previous measurements10,20.

We first identify the character of this excitation by determining the associated magnetic and charge contributions to the RIXS spectra. This can be uniquely achieved by resolving the polarization of both the incident and scattered photons20. Namely, magnetic excitations flip spins and necessarily change the angular momentum of the photons in the scattering process, that is, contribute to the crossed-polarization channel (ππ or σσ). Conversely, excitations preserve the angular momentum of the photon and contribute to the parallel polarization channel (ππ or σσ). Figure 1d, e shows polarization-resolved RIXS spectra for two different in-plane momenta. At qx = (0.045, 0) the features of the zone centre excitation are fully suppressed for crossed polarizations (ππ) and the spectrum contains only the parallel polarization (ππ) contribution. For larger momentum transfer qx = (0.095, 0) the mode disperses towards higher energy (about 0.8 eV) (Fig. 1e) and the well-studied paramagnon excitation7,20 emerges on a lower energy scale (about 0.3 eV). As expected, the paramagnon yields spectral weight in both polarizations owing to the mixture of single spin-flip excitation, double spin-flip and incoherent particle–hole charge excitations, whose spectral weight increases with increasing doping concentration21. Importantly, the zone centre excitations, which are separated in energy from the paramagnons, still appear only for parallel polarization geometries. Thus, we first conclude that the zone centre excitations are a branch of pure charge modes.

A second insight can be obtained from a comprehensive mapping of the energy–momentum dispersion in all three dimensions of reciprocal space, in contrast to previous RIXS experiments that explored the projected in-plane momentum without focusing on the qz dependence6−8.

Figure 2a, b shows the RIXS intensity maps as a function of momentum transfer along the hh- and h-directions (that is, along (0, 0, l)−(h, h, l) and (0, 0, l)−(h, 0, l), respectively) at l = 1 and l = 1.65. We denote
Plasmons in a layered electron gas and dispersive charge excitations in electron-doped copper oxides.

A. Plasmon dispersion in a layered electron gas as a function of in- and out-of-plane momentum transfers \( q_{\parallel} \) and \( q_z \), respectively. Different branches correspond to specific out-of-plane momentum transfers \( q_z \) and are a result of the interplane Coulomb interaction between the periodically stacked planes with distance \( d \) (see inset). Their spectrum varies from a single optical branch (light blue line) with the characteristic plasma frequency \( \omega_p \) for \( q_z = 0 \), to a range of acoustic branches (light red lines) with the lowest-energy branch for \( q_z = \pi/d, 3\pi/d \). The electron–hole pair excitation continuum is illustrated by the orange shaded area. B, C, RIXS spectra of NCCO (\( x = 0.15 \)) and LCCO (\( x = 0.175 \)) at in-plane momentum transfers \( q_{\parallel} = (0.045, 0) \) and (0.095, 0) for incident photon energies tuned to the Cu L_{3,2}-edge at temperature \( T \approx 20 \) K. The spectral peak assigned to the dispersive zone centre excitation is highlighted by the red-shaded peak profile. The additional peak at about 0.3 eV in \( \sigma \) is identified as the paramagnon, justified by the polarization-resolved RIXS spectra shown in D. E. The NCCO and LCCO spectra are offset in the vertical direction for clarity. D, E. Polarization-resolved RIXS spectra of LCCO at \( q_{\parallel} = (0.045, 0) \) and (0.095, 0). Charge excitations are detected in the parallel polarization channel (\( \sigma \sigma \), red line) while magnetic excitations are detected in the spin-flip crossed-polarization channel (\( \sigma \pi \), blue line). The reference spectrum (open symbols) is taken with \( \pi \) polarized incident photons in the absence of polarization analysis, corresponding thus to the sum of \( \sigma \sigma \) and \( \sigma \pi \). Red and blue markers indicate the relevant spectral weight maximum of the \( \sigma \sigma \) and \( \sigma \pi \) channel, respectively. a.u., arbitrary units.

remarkably, the zone centre excitation continues dispersing towards higher energy with further increasing \( l \), insensitive to the next high symmetry point at \( l = 1.5 \). In fact, the energy scale of the excitation continues to increase even at \( l = 1.8 \), the highest out-of-plane momentum transfer that was accessible in our experiment. We also observe the same peculiar behaviour in our momentum-resolved RIXS measurements on NCCO (see Extended Data Fig. 3), suggesting a universal origin of the branch of charge modes from the three-dimensional (3D) nature of the Coulomb interaction in an otherwise layered quasi-2D material.

These results can be rationalized by doubling the out-of-plane Brillouin zone size, implying that the crystallographic unit cell with lattice constant \( c \) does not set the periodicity of the zone centre excitation, but rather \( d = c/2 \): the nearest-neighbour CuO_2 plane spacing (Fig. 3b). Hence, a new index \( I^* \) in units of \( 2\pi/d \) appropriately describes the Brillouin zone that is ‘felt’ by the zone centre excitation and establishes the proper periodicity of the dispersion. An obvious mechanism that could induce such a Brillouin zone ‘reconstruction’ in a quasi-2D system is the interplanar Coulomb interaction.

Interestingly, the data shown in Figs. 2, 3a indicate that the plasmon peak broadens when approaching the equivalent zone centre (0, 0, \( I^* = 1 \)) along the \( I^*-\)direction (see also Extended Data Figs. 1c, 3c). In fact, at momentum transfer (0.025, 0, \( I^* = 0.925 \)), that is, close to the equivalent zone centre (0, 0, \( I^* = 1 \)), the plasmon linewidth of approximately 0.5 eV (Fig. 3a) is similar to previous transmission EELS reports and optical conductivity measurements at (0, 0, 0). The increasing incoherence of the mode near the zone centre is consistent with the Cu L_{3,2}-edge at temperature \( T \approx 20 \) K. The spectral peak assigned to the dispersive zone centre excitation is highlighted by the red-shaded peak profile. The additional peak at about 0.3 eV in \( \sigma \) is identified as the paramagnon, justified by the polarization-resolved RIXS spectra shown in D. E. The NCCO and LCCO spectra are offset in the vertical direction for clarity. D, E. Polarization-resolved RIXS spectra of LCCO at \( q_{\parallel} = (0.045, 0) \) and (0.095, 0). Charge excitations are detected in the parallel polarization channel (\( \sigma \sigma \), red line) while magnetic excitations are detected in the spin-flip crossed-polarization channel (\( \sigma \pi \), blue line). The reference spectrum (open symbols) is taken with \( \pi \) polarized incident photons in the absence of polarization analysis, corresponding thus to the sum of \( \sigma \sigma \) and \( \sigma \pi \). Red and blue markers indicate the relevant spectral weight maximum of the \( \sigma \sigma \) and \( \sigma \pi \) channel, respectively. a.u., arbitrary units.
with incoherent charge dynamics inferred from c-axis optical conductivity. However, such linewidth evolution appears to be different from the Landau quasi-particle picture in which the plasmon peak should be sharpest at the zone centre where the particle–hole (Landau) continuum is minimal and well separated from the plasmon. Other mechanisms, such as the presence of non-Fermi-liquid-producing interactions, polar interlayer electron–phonon coupling and Umklapp mechanisms, such as the presence of non-Fermi-liquid-producing interactions, polar interlayer electron–phonon coupling and Umklapp mechanisms, may reconcile these observations.

We now investigate the acoustic plasmon bands in LCCO as a function of the carrier density, which is nominally the Ce concentration. We have also verified the systematics of doping concentration among different samples using an internal spectral reference—the dd excitation of RIXS spectra (Methods and Extended Data Fig. 5). As shown in Fig. 4a, b, the plasmon bands exhibit a detectable doping dependence. In a naive picture of the free electron model, the plasmon energy is expected to increase proportionally to \( \sqrt{x/m^*} \), where \( m^* \) is the effective electron mass. Consistent with such an expectation the mode energies of \( x = 0.11 \) to about 0.15 increase linearly with \( \sqrt{x} \) (Fig. 4c), further substantiating the attribution of the zone centre excitation to a plasmon. However, for higher dopings, the rate of increase slows down and appears to hit a plateau for \( x = 0.17 \) and 0.18. This observation suggests a possible variation in the band dispersion or Fermi surface at approximately \( x \approx 0.15 \). Recent Hall-effect measurements on LCCO have indicated Fermi-surface reconstruction due to antiferromagnetic correlations ending at around \( x \approx 0.14 \) (ref. 26), corroborating our observation.

We note that our observation appears to be distinct from a recent high-resolution EELS measurement, which reported featureless in-plane charge excitations in a hole-doped copper oxide. This suggests a distinct behaviour of the charge degrees of freedom between electron- and hole-doped copper oxides. However, given the similarities of the layered 2D \( \text{CuO}_2 \) plane structure between electron- and hole-doped materials, we should expect a similar three-dimensionality in the charge dynamics of the hole-doped copper oxides. Although previous RIXS studies on hole-doped compounds primarily focused on magnetic, orbital and other high-energy excitations, few investigated the region of energy–momentum space necessary to identify and characterize the acoustic plasmon. More detailed RIXS measurements on hole-doped compounds, in both single and multi-layer systems with higher \( T_c \), will be able to clarify these issues.

Our observation of 3D plasmon modes indicates that the copper oxides, at the very least, should be modelled as layered 2D systems when describing their charge dynamics, a fact that is often overlooked. This change of perspective has important implications. First, in a 2D doped Mott insulator, the in-plane charge fluctuations are strongly suppressed by the Coulomb interaction. Thus the spin dynamics become the most prominent low-energy excitations, thought to be most relevant to high-\( T_c \) superconductivity. Our results challenge this view by demonstrating that the low energy charge fluctuations can be quite active owing to the layered structure of the copper oxide. Early theories suggested that acoustic plasmons may be able to mediate pairing, or perhaps more importantly, enhance \( T_c \) for Cooper pairs bound by other interactions. Second, the Coulomb energy stored between \( \text{CuO}_2 \) planes can be important and may play a part in the energy savings associated with the superconducting transition. Third, the energy of the plasmon extrapolates to approximately zero at the projected zone centre. This implies a negligible single electron hopping between adjacent \( \text{CuO}_2 \) planes, leaving the Coulomb interaction as the sole donor of energy.
source of interplanar coupling. Such restriction of the charge to the 2D planes may enhance the effects of quantum confinement at the heart of topological theories for superconductivity in copper oxides. However, important questions remain about the impact of the interplanar Coulomb interaction on the electronic structure, the pseudogap and charge- and spin-density-wave orders. In a broader context, our result fits within the general framework that the Coulomb interaction affects charge dynamics in nanoscale heterostructures of 2D quantum materials, including transitional metal chalcogenides and graphene. Manipulating the Coulomb interaction could enable fine-tuning of desired properties of artificial quantum materials, such as plasmonics in nanostructures, with a range of applications from sensors to photonic and electronic devices for communications (for example, see the Plasmonics Focus Issue introduced in ref. 32).

Fig. 3 | Out-of-plane plasmon dispersion. a, RIXS intensity map of LCCO ($x = 0.175$) for momentum transfer along the out-of-plane direction at $l^* = 0.25$. The out-of-plane momentum is indicated by the indices $l$ (top scale, units of $2\pi/c$) corresponding to the crystallographic $c$-axis, and $l^*$ (bottom scale, units of $2\pi/x$) corresponding to the CuO2 plane spacing. White symbols indicate fitted peak positions of the zone centre excitation. The black vertical line highlights the high symmetry plane spacing. White symbols indicate fitted peak positions of the zone centre mode for momentum transfer along the out-of-plane $h$-axis. The symbol style refers to the dopings as indicated in Methods. The energy is expressed in units of $t/d$, the hopping integral between the CuO2 planes in red. La/Ce Cu O3, LCCO with the crystallographic unit cell indicated by black lines and the CuO2 planes in red. The energy loss is 0.035 to 0.070, constrained by assuming that the mode energy is 0 at $x = 0.11$ to 0.15, constrained by assuming that the mode energy is 0 at $x = 0.11$ to 0.15. Constrained by assuming that the mode energy is 0 at $x = 0.11$ to 0.15.

Fig. 4 | Doping dependence of the plasmon. a, Energy dispersion of the LCCO zone centre mode for momentum transfer along the $h$-direction at $l^* = 0.5$. Red symbols are the fitted peak positions for Ce doping concentrations $x = 0.11, 0.13, 0.15, 0.17$ and 0.18. The bottom inset shows the temperature versus doping phase diagram of LCCO thin films adapted from ref. 29 including a superconducting dome (orange shading, SC) and a small Fermi surface region (blue shading) undergoing a crossover to a reconstructed Fermi surface (blue shading, FSR) that ends at $x = 0.14$. b, Representative raw RIXS spectra (open symbols) for momentum transfer $h = 0.035$ and $l^* = 0.5$ for different dopings. The anti-symmetrized Lorentzian fit profiles are shaded in red with vertical black markers indicating the peak positions and the horizontal bars indicating the error bars. Spectra are offset in the vertical direction for clarity. c, Mode energies versus the square root of $x$ for in-plane momenta from $h = 0.035$ to $h = 0.095$ at $l^* = 0.5$. The symbol style refers to the dopings as indicated in a. Dashed lines are linear fits of the mode energy versus $\sqrt{x}$ for dopings $x = 0.11$ to 0.15, constrained by assuming that the mode energy is 0 at $x = 0$. We note that the rate of increase deviates from linear behaviour and forms a plateau for higher doping, as mentioned in the text. Error bars in this figure are estimated from the uncertainty in energy-loss reference-point determination ($\pm 0.01$ eV) together with the standard deviation of the fits.
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METHODS

The c-axis-oriented LCCO thin films with Ce concentrations of $x = 0.11, 0.13, 0.15, 0.17$ and $0.18$ were fabricated on (100) SrTiO$_3$ substrates by pulsed laser deposition using a KrF excimer laser. The annealing process was optimized for each $x$. The superconducting $T_c$ of the $x = 0.11$ and $0.13$ films is around $30 \, K$ and around $22 \, K$, respectively. The $x = 0.175$ and $0.18$ films did not show a superconducting transition. The NCCO single crystal with Ce concentration $x = 0.15$ was grown by the travelling-solvent floating-zone method in Os and annealed in Ar at $900^\circ \text{C}$ for 10 h. The $T_c$ is about $26 \, K$.

The RIXS measurements were performed at beamline ID32 of the ESRF using the high-resolution 'ERIXS' spectrometer. The scattering angle $2\theta$ can be changed in a continuous way from $50^\circ$ to $150^\circ$. The samples were mounted on the 6-axis in-vacuum Huber diffractometer/ manipulator and cooled to around $20 \, K$. The RIXS data were obtained with incident $e^-$ polarization (perpendicular to the scattering plane, high-throughput configuration). The incident photon energy was tuned to the maximum of the Cu $L_3$ absorption peak at about $931 \, eV$. The energy resolution was $\Delta E = 60 \, meV$ for the $x = 0.11, 0.15, 0.17$ and $0.18$ LCCO sample and $\Delta E = 68 \, meV$ for the $x = 0.13$ and $0.175$ LCCO and the $x = 0.15$ NCCO sample. The RIXS polarization-resolved measurements were conducted with a wider monochromator exit slit at a resolution of $\Delta E \approx 85 \, meV$ in order to partly compensate the reduced efficiency of the polarimeter with respect to the normal configuration. For each transferred momentum a non-resonant silver paint or carbon tape spectrum provided the exact position of the elastic (zero energy loss) line. For the polarimetric RIXS measurements of Fig. 1d, $e^-$ polarization incident on the sample was used and the graded multilayer served as analyser of the scattered photons, as explained in detail in ref. 38. We note that Cu L-edge RIXS is the so-called direct RIXS process, involving the resonant transition from the Cu 2p core level to the 3d$_{\uparrow \downarrow \uparrow \downarrow}$ orbital, which constitutes the electronic structure near the Fermi energy. The Cu L-edge RIXS is capable of probing a wide range of elementary excitations, including the orbital (that is, the $dd$-excitations), magnetic excitations, phonons and charged excitations.

In the theory calculations we consider the three-band Hubbard model with the following standard parameters in units of electronvolts: $U_{dd} = 8.5, U_{pp} = 4.1, t_{pd} = 1.13, t_{pp} = 0.49, \Delta_{pd} = 3.24$ (ref. 34). Determinant quantum Monte Carlo (DQMC) is used to solve the model on a fully periodic $16 \times 4$ cluster at a temperature of $T = 0.125 \, eV$. For each doping, 512 independently seeded Markov chains with 50,000 measurements each are run. The charge susceptibilities obtained by DQMC are analytically continued to real frequency using the maximum entropy method with model functions determined by the first moments of the data. The inverse dielectric function plotted in Fig. 3c is obtained via

$$\chi(q, \omega) = 1 + \frac{1}{\omega^2 - \text{Im}(\chi(q, \omega))}$$

where $\chi(q, \omega)$ is the real frequency charge susceptibility obtained from DQMC and the maximum entropy method. The long-range and 3D Coulomb interactions neglected in the three-band Hubbard model are captured by $V_D$. We use the layered electron gas form $17, 32$: $V_D = \frac{\alpha^2}{\delta^{1/2} \sqrt{2\pi d \epsilon_\infty}} \sinh(q_d d)$, where $d$ is the interplane spacing and $q_d$ are the in-plane and out-of-plane components of the momentum transfer, respectively. $\epsilon_\infty$, the sole free parameter of our calculation, is adjusted to give a roughly 1-eV mode for $q_d = 0$ and the smallest non-zero $q_d = 0.0625, 0$; its value is not varied with doping.

Fit of the plasmon dispersion in the layered electron gas model. We consider the energy--momentum dispersion of a plasmon mode in a layered electron gas model for momentum transfer $q_d$ along the $d$-direction at fixed out-of-plane momentum transfer values $q_z$, along the $l$-$m$-direction. Let $d$ be the spacing between adjacent CuO$_2$ planes and $q_d = \pm \pi/d$ and $\epsilon_\infty$ be the high-frequency dielectric constant due to the screening by the core electrons. Following ref. 38, the Coulomb potential of a layered electron gas is:

$$V_D = \alpha^2 \frac{\sinh(q_d d)}{q_d \cosh(q_d d) - \cos(q_d d)}$$

with

$$\alpha = \frac{e^2 d}{2\pi \delta^{1/2} \epsilon_\infty}$$

In an isotropic medium it is well-known that the 3D Coulomb potential is $e^2/\epsilon_\infty \cdot q^2$, while in a 2D plane the Coulomb potential is $e^2/2\epsilon_\infty \cdot q^2$. These are the two limits of the above form of the layered electron gas, with $V_D$ becoming $e^2/\epsilon_\infty \cdot q^2$ in the approximation of long wavelengths ($q_d = 0$ and $q_d = 0 < 1$) and $V_D = e^2/2\epsilon_\infty \cdot q^2$ for short wavelengths ($q_d > 1$, independent of $q_z$).
Extended Data Fig. 1 | Fits of the RIXS spectra. a, Fits of LCCO (x = 0.175) RIXS spectra at in-plane momentum transfer positions $q_h = (0.045, 0)$ and $(0.095, 0)$, representative of all fits performed in the scope of this work. The model uses a Gaussian for the elastic peak (green) and anti-symmetrized Lorentzians for all other contributions in the spectrum, convoluted with the energy resolution (here $\Delta E = 68$ meV) via Gaussian convolution. The anti-symmetrized Lorentizan is used to ensure zero mode intensity at zero energy loss, as explained in the supplementary information of ref. 40. The peak profiles of the zone centre excitation (plasmon) are shaded in red. b, Full-width at half-maximum (FWHM) of the zone centre excitation (plasmon) as extracted from the fits for momentum transfer along the $hh$- and $h*$- directions at $l^* = 0.5$, $l^* = 0.825$ and $l^* = 0.9$, corresponding to the fitted peak positions shown in Fig. 2c. Error bars are the standard deviation of the fits. c, FWHM of the zone centre excitation (plasmon) as extracted from the fits for momentum transfer along the out-of-plane direction at $h = 0.025$. The panel corresponds to the fitted peak positions shown in Fig. 3a.
Extended Data Fig. 2 | Raw data and fits of the RIXS spectra. 

a, b, Raw RIXS spectra (red) of LCCO (x = 0.175) together with the fits (solid black lines) for momentum transfer along the \( hh \)-direction (a) and \( h \)-direction (b) at different \( l^* \). The spectra are offset in the vertical direction for clarity.

c, Raw RIXS spectra together with the fits for momentum transfer along the \( l^* \)-direction at \( h = 0.025 \).
**Extended Data Fig. 3** | Three-dimensionality of the zone centre excitations in NCCO. a, b, RIXS intensity maps of NCCO (x = 0.15) for momentum transfer along the h-direction at \( l^* = 0.5 \) and \( l^* = 0.825 \). Red and grey symbols indicate least-squares-fit peak positions of the zone centre excitation and the paramagnon, respectively. The inset indicates the probe direction in reciprocal space. c, RIXS intensity map of NCCO (x = 0.15) for momentum transfer along the out-of-plane direction at \( h = 0.025 \). White symbols indicate fitted peak positions of the zone centre excitation. Error bars are estimated from the uncertainty in energy-loss reference-point determination (±0.01 eV) together with the standard deviation of the fits.
Extended Data Fig. 4 | Fits of the plasmon dispersion in the layered electron gas model. a, Fits (solid lines) of the mode energies of LCCO (x = 0.175) (red symbols) as a function of in-plane momentum transfer $q\parallel$ along the $h$-direction at $l^\ast = 0.5$, $l^\ast = 0.825$ and $l^\ast = 0.9$. The fit is global, that is, the three $l^\ast$ datasets are fitted simultaneously with the same fit parameter, as described in the Methods. Error bars of the data points are the same as those estimated in Fig. 2c.
Extended Data Fig. 5 | Verification of electron doping systematics via dd excitations in the RIXS spectra. a, dd excitations in RIXS spectra at momentum transfer (0.015, 0, 1) taken from samples with different Ce doping concentrations x. The energy positions of dd excitations shift to higher energy with increasing electron doping, which can be used as an internal reference to verify the doping concentrations. The inset shows a zoom-in of the leading-edge region of the dd excitations. b, The correlation between the Ce concentration x and the energy of the dd-leading edge (inflection point) and the 3d_{z^2-r^2} peak. All samples show good correlation except for the x = 0.175 sample, indicating a larger uncertainty of its doping concentration. Thus, the x = 0.175 data were not included in Fig. 4. The error bars are estimated from the standard deviation of the fit used to determine the energy of the dd-leading edge and the 3d_{z^2-r^2} peak.