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# A Comparison of Stripe Modulations in $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$ and $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$

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We report combined soft and hard x-ray scattering studies of the electronic and lattice modulations associated with stripe order in  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$  and  $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$ . We find that the amplitude of both the electronic modulation of the hole density and the strain modulation of the lattice is significantly larger in  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$  than in  $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$  and is also better correlated. The in-plane correlation lengths are isotropic in each case; for  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$ ,  $\xi^{\text{hole}} = 255 \pm 5 \text{ \AA}$  whereas for  $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$ ,  $\xi^{\text{hole}} = 111 \pm 7 \text{ \AA}$ . We find that the modulations are temperature independent in  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$  in the low temperature tetragonal phase. In contrast, in  $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$ , the amplitude grows smoothly from zero, beginning 13 K below the LTT phase transition. We speculate that the reduced average tilt angle in  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$  results in reduced charge localization and incoherent pinning, leading to the longer correlation length and enhanced periodic modulation amplitude.

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## I. INTRODUCTION

One dimensional modulations of the lattice and of the charge and spin density appear to be both ubiquitous in the cuprates and intimately tied up with the physics of these materials. For example, superconducting phase coherence is suppressed in systems in which charge and spin stripes are static<sup>33-35</sup> while dynamic stripes may promote pairing<sup>36,37</sup>. Further, stripe order coexists with 2D superconducting correlations above the bulk superconducting phase<sup>38,39</sup> and indeed it has been argued that broken rotational symmetry may be a defining feature of the pseudogap phase - so-called “nematic” electronic order, a viewpoint for which there is some evidence<sup>40,41</sup>. Understanding these modulations is therefore a prerequisite to gaining a detailed knowledge of the physics of the cuprates.

In their full incarnation, these density waves - “stripes” - consist of three concomitant modulations; of the spin density, of the charge density and of the lattice itself. In a few cases, namely  $\text{La}_{2-x}\text{Ba}_x\text{CuO}_4$ <sup>35,39,42-44</sup> and  $(\text{La,R})_{2-x}\text{Sr}_x\text{CuO}_4$ , where R=Nd<sup>45</sup>, or Eu<sup>46,47,48</sup>, these modulations are stabilized by the symmetry of the low temperature crystal structure to form static structures. It is here, where the respective order parameters are well developed, that one can best characterize them. The basics of the static structures are known: they are comprised of hole-rich anti-phase antiferromagnetic domain walls running along the Cu-O bond direction. In real space the spin density then has twice the wavelength of the charge and lattice modulations. In a scattering experiment, these give rise to satellites at  $(0.5 \pm \delta, 0.5, 0)$  and  $(\pm 2\delta, 0, 0.5)$ , and symmetry related positions, respec-

tively. However, basic questions such as the relative amplitudes in the different compounds and the coupling between these modulations remain unanswered.

Here we report a combined soft and hard x-ray scattering study of the low temperature stripe order in  $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$  (LBCO) and  $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$  (LNSCO). In particular, soft x-ray scattering at the pre-edge of the O K-edge (529.3 eV) probes the spatial variation of the oxygen 2*p* partial density of states at the Fermi level, which is related to the mobile charge carrier density<sup>42,49,50</sup>, while hard x-ray (11.3 keV) scattering probes the associated lattice distortion. We find that the stripe correlation lengths are isotropic in the Cu-O plane in both systems, but LBCO is a factor of two better correlated than LNSCO. Further, by careful normalization to structural Bragg peaks we are able to quantify the relative size of the electronic modulation and the lattice distortion in these two cases. We find that the amplitude of the electronic modulation is  $\sim 10$  times larger in LBCO than in LNSCO, while the lattice distortion is only  $\sim 4$  times larger. We further find that the modulations are temperature independent in LBCO once the LTT phase is entered, suggesting that perhaps the intrinsic ordering temperature is significantly higher. In contrast, in LNSCO, the amplitude grows smoothly from zero, beginning 13 K below the LTT phase transition.

## II. EXPERIMENTAL

Samples of both LBCO and LNSCO were grown using the floating zone method, at Brookhaven National Laboratory, USA and the University of Tokyo, Japan respec-

tively. Both samples proved to be of high crystal quality, with mosaic spreads of  $\leq 0.02^\circ$  measured at 11.3 keV. Throughout this paper both systems are indexed with the high temperature tetragonal (HTT) ( $I_4/mmm$ ) unit cell.

Soft x-ray diffraction experiments were carried out on the X1A2 beamline at the NSLS, Brookhaven National Laboratory, USA using a six-circle in-vacuum diffractometer. The samples were cleaved *ex-situ* to reveal surfaces with a  $[001]_{\text{HTT}}$  surface normal and mounted such that the  $[001]_{\text{HTT}}$  and  $[100]_{\text{HTT}}$  directions lay in the scattering plane. Experiments were performed in a vertical scattering geometry with  $\sigma$ -incident x-rays, i.e. the electric field of the incident x-ray photons was always along the HTT  $b$ -axis, within the  $\text{CuO}_2$  planes. The samples were cooled in a He flow cryostat to a base temperature of 12 K. Scattered x-rays were detected with an in-vacuum CCD. During the measurements the beam footprint was smaller than the sample. Data were collected by performing a  $\theta - 2\theta$  scan, collecting a single CCD image at each datum. Each CCD image was then converted to reciprocal space coordinates and binned onto a regular three-dimensional grid<sup>51</sup>. Part of the soft x-ray experiments were performed at BESSY with a diffractometer described in Ref. 46.

Hard x-ray diffraction experiments on the same samples were conducted on beamline X22C at the NSLS. The incident photon energy was 11.3 keV and scattered x-rays were detected by means of a point-detector, with the resolution defined by slits. Experiments were conducted in a vertical scattering geometry, with the  $[001]_{\text{HTT}}$  and  $[100]_{\text{HTT}}$  directions lying in the scattering plane. For both LBCO and LNSCO the instrumental resolution was identical.

### III. RESULTS

#### A. Soft X-ray Scattering

We first discuss the characterization of the electronic modulation, beginning with LBCO. In Fig. 1 we show cuts through reciprocal space in the HK- (bottom) and HL- planes (top) for LBCO. These data were taken with  $E_i=529.3$  eV, an energy corresponding to the Mobile Carrier Peak (MCP) in the oxygen K-edge absorption<sup>49</sup>. At this energy the scattering from the charge stripes is maximized<sup>42</sup>. We extract a correlation length from these data by taking line cuts through the three dimensional grid and fitting to Lorentzian-squared lineshapes. We find that the stripe correlations within the  $a$ - $b$  plane are isotropic with  $\xi_H = 255 \pm 5 \text{ \AA}$ , where  $\xi = \frac{1}{\text{HWHM}}$ . Measurements at the Cu  $L_3$  edge revealed a similar correlation length. We note that because of the superior resolution of the present set up, this represents the highest precision measurement to date of this correlation length. It is consistent with previous work which found  $\xi \approx 200 \text{ \AA}$  at this energy<sup>42</sup>. As has been previously observed, the

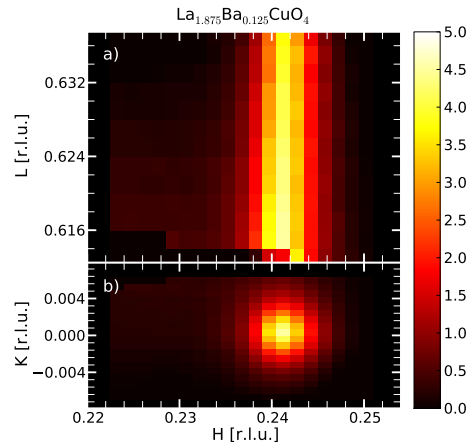


FIG. 1. Soft x-ray resonant scattering intensity in the vicinity of the  $(0.24, 0, L)_{\text{HTT}}$  stripe order peak for LBCO in the a)  $(H, 0, L)_{\text{HTT}}$  plane and b)  $(H, K, 0.624)_{\text{HTT}}$  plane in reciprocal space. Data were taken at  $T=15$  K, and  $E_i=529.3$  eV. At this energy, the scattered intensity reflects the modulation of the oxygen density of states in real space. The intensity map reveals that the stripe correlations are isotropic in the  $a$ - $b$  plane and are two dimensional.

stripes are weakly correlated in the  $c$ -axis direction, giving rise to the uniform streak of intensity along  $L$  in Fig. 1.

Figure 2 shows a comparison in the HK- plane, along with line cuts along the H direction, between LBCO and LNSCO. Immediately apparent is the far poorer correlation length in LNSCO, where we find that the correlation length is again isotropic but in this case  $\xi_H = 111 \pm 5 \text{ \AA}$ .

In both cases, the wavevector in the H direction, is 0.24 r.l.u. and is therefore incommensurate. These measurements were performed using an orientation matrix defined *in-situ* by the  $(002)_{\text{HTT}}$  and  $(101)_{\text{HTT}}$  Bragg reflections, and as such their values are reported with high confidence. Previous studies have found similar incommensurabilities at these dopings (see e.g. Refs. 42, 44, and 52). We will return to this point later.

We are also able to compare the peak intensities in the two systems. To do so, the intensities shown in Fig. 2 have been normalized to the integrated intensity of the respective  $(002)_{\text{HTT}}$  Bragg reflections (measured at 1 keV) and corrected for their different structure factors. The integrated intensity,  $V$ , of a reflection is proportional to the product of the peak intensity and the widths in three orthogonal directions,  $V \propto I_{\text{Peak}} \times \Gamma_H \times \Gamma_K \times \Gamma_L$ . As discussed,  $\Gamma_K = \Gamma_H$ , and we make the assumption that the correlation lengths along L are equal in the two materials. The ratio of integrated intensities is then  $V_{\text{AB}}^{\text{hole}} = (I_{\text{Peak A}} \times \Gamma_A^2) / (I_{\text{Peak B}} \times \Gamma_B^2)$ . Calculating from the data presented in Table I we find that the ratio of integrated intensities, corrected for all experimental factors is  $V_{\text{rel}}^{\text{hole}} = 97 \pm 10$ , i.e. the electronic charge stripe modulation in LBCO is  $\sim 9.8 \pm 0.5$  times larger in amplitude

TABLE I. Comparison of hole and lattice modulation wavevectors, correlation lengths and peak intensities for LBCO and LNSCO as measured with soft and hard x-rays respectively.

	Center	HWHM	Peak
	[r.l.u.]	[r.l.u.] $\times 10^{-3}$	Intensity
LBCO	529.3 eV	0.241	$7.82 \pm 0.01$
LNSCO	529.3 eV	0.238	$(1.54 \pm 0.005) \times 10^{-3}$
LBCO	11.3 keV	-	$0.225 \pm 0.002$
LNSCO	11.3 keV	-	$(4.17 \pm 0.18) \times 10^{-3}$

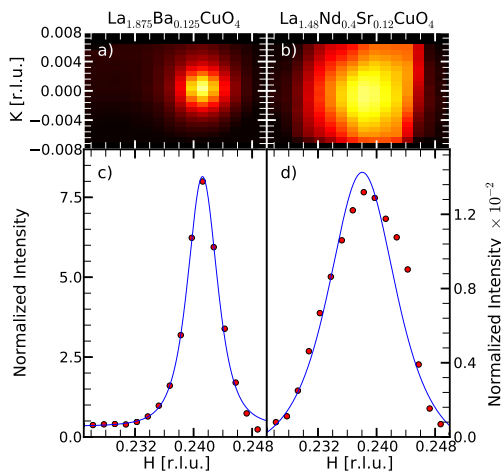


FIG. 2. Comparison of scattering from stripe modulations in LBCO and LNSCO. Data were taken at  $T=15$  K, and  $E_i=529.3$  eV. a) and c)  $(H, K, 0.624)_{\text{HTT}}$  plane and  $(H, 0, 0.624)_{\text{HTT}}$  scan (circles) for LBCO. b) and d) same for LNSCO. In c) and d) the solid line is the result of a fit to a Lorentzian squared lineshape. Scattered intensities are normalized to the respective  $(002)_{\text{HTT}}$  Bragg peaks and therefore may be compared directly to each other.

than in LNSCO.

### B. Hard X-ray Scattering

We next measure the lattice modulation that is concomitant with the electronic modulation using hard x-rays. In Fig. 3, we show scans performed with  $E_i=11.3$  keV through the  $(0.24, 0, 11.5)$  charge stripe wavevector. We have again normalized the scattered intensity by a nearby Bragg peak, in this case the  $(1, 0, 12)_{\text{HTT}}$ . Due to the poorer experimental resolution of the hard x-ray measurements, it is necessary to deconvolve the experimental resolution. As an upper bound on this resolution, we took the sharpest of the LBCO and LNSCO  $(1, 0, 12)_{\text{HTT}}$  reflections. After deconvolving the instrumental resolution we find  $\xi_{\text{LBCO}}^{\text{lattice}} = 178 \pm 50$  Å

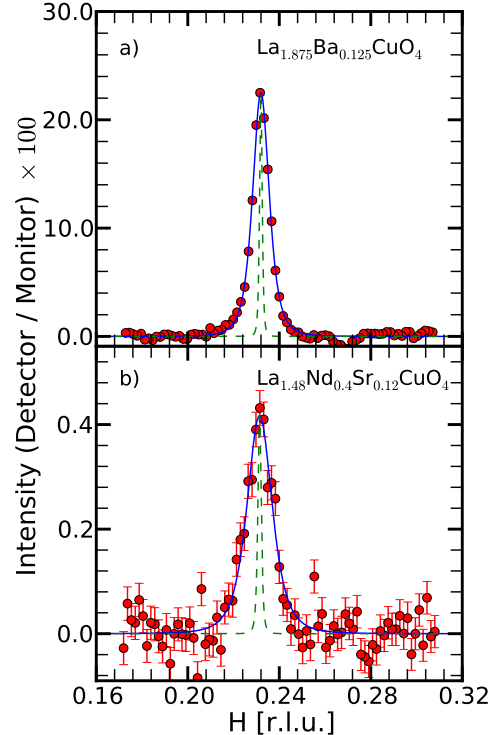


FIG. 3. Comparison of stripe order peaks in a) LBCO and b) LNSCO as measured by hard x-rays,  $E_i=11.3$  keV at  $T=12$  K. Solid lines are the results of fits to Lorentzian squared lineshapes. Dashed lines represent the width of the  $(1, 0, 12)_{\text{HTT}}$  nearby Bragg reflection in each case, demonstrating the two samples are of similar quality. Scattered intensities are normalized to the  $(1, 0, 12)_{\text{HTT}}$  Bragg peak and therefore may be compared directly to each other.

and  $\xi_{\text{LNSCO}}^{\text{lattice}} = 107 \pm 20$  Å. Comparing the widths of the stripe peaks as measured with soft and hard x-rays, we find that the widths are equal within experimental error for the two x-ray energies in both LBCO and LNSCO.

From the hard x-ray deconvolved width and the peak intensity, we can calculate the ratio of the lattice modulations for the two systems following the same arguments as before. We find that the ratio of the integrated intensities of the lattice modulations is  $V_{\text{rel}}^{\text{lattice}} = 20 \pm 6$ , i.e. the lattice modulation amplitude is  $\sim 4.4 \pm 0.7$  times larger in LBCO than in LNSCO.

### C. Temperature Dependence

We next present the temperature dependence of the respective modulations, and of the LTT phase, for the two systems. This is shown in figure 4. For soft x-rays, the LTT phase is measured through the forbidden  $(001)_{\text{HTT}}$  reflection, observable only in the LTT phase at

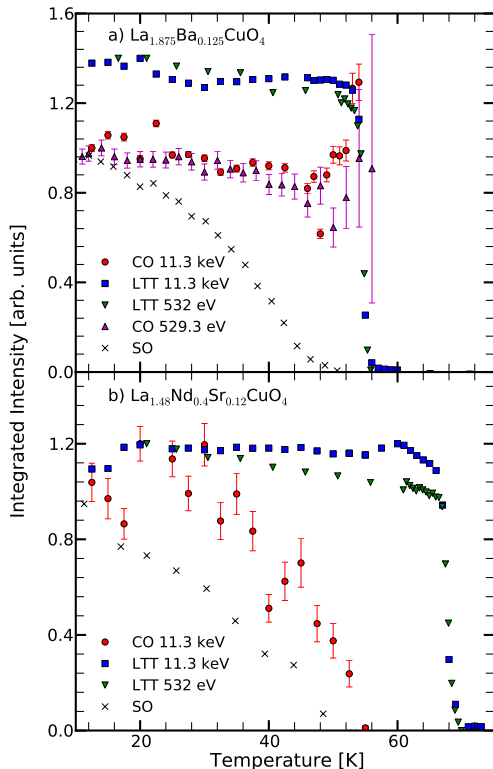


FIG. 4. Temperature dependence of the lattice modulation (red circles), electronic modulation (magenta up triangles), spin modulation (black crosses) and the LTT phase as measured by soft (green down triangles) and hard x-rays (blue squares) in both LBCO (a, top panel) and LNSCO (b, bottom panel). The data presented for the spin modulation were taken from Ref. 44 and Ref. 52 for LBCO and LNSCO, respectively.

$E_i = 532 \text{ eV}$ <sup>47</sup>. For hard x-rays, the  $(1, 0, 12)_{\text{HTT}}$  reflection, observable only in the LTT phase is used. In the case of the superlattice modulations, the integrated intensity of the reflection is taken to be  $\propto I \times \Gamma^2$ . This is consistent with two dimensional stripes, where the  $L$ -axis dependence of the scattering is dominated by the structure factor due to the very short correlation length<sup>53</sup>.

There are several important things to note from this plot. First, for LBCO it is clear that an observable stripe peak, both the strain modulation and the charge density modulation, is only visible within the LTT phase. In addition, the integrated intensity is constant, within experimental error, for both the electronic and structural modulations. This demonstrates that the amplitudes of both these modulations are constant up to the LTT transition temperature, that is the order parameter is not changing, only the correlation length. This suggests that had the LTT transition occurred at a higher tempera-

ture in LBCO then it is likely that both the electronic and structural modulations would also have persisted to higher temperatures.

In contrast, for LNSCO (Fig. 4b) the amplitude decays upon warming and is unobservable above  $T_{\text{CO}}=55 \text{ K}$ , well below the LTT transition at  $T_{\text{LTT}}=68 \text{ K}$ . This behavior is reminiscent of recent results in  $\text{La}_{1.8}\text{Eu}_{0.2-x}\text{Sr}_x\text{CuO}_4$  which has an even larger difference between  $T_{\text{LTT}}$  and  $T_{\text{CO}}$  of  $\sim 55 \text{ K}$ <sup>46,47</sup>.

Finally, it is worth noting that the temperature dependence of the LTT-peaks are identical in both samples when measured at 11.3 keV and 530 eV, demonstrating that the soft x-rays are probing the same bulk-like behavior as the hard x-rays and that the samples are homogeneous.

## IV. DISCUSSION

### A. Correlation Lengths

The result that the charge stripes display isotropic correlation lengths is perhaps surprising given that the stripes in a single  $\text{CuO}_2$  layer are highly anisotropic. While four-fold rotational symmetry is recovered when averaging over the whole crystal, since successive sheets along the  $c$ -axis are rotated by  $90^\circ$  with respect to each other, these experiments only observe charge stripes with a propagation wavevector along one direction in reciprocal space - by virtue of the diffraction condition. There is therefore no *a priori* reason to expect these correlations to be isotropic. One possible explanation is that nearest-neighbor  $c$ -axis correlations couple the charge stripe correlations on alternating layers. It has been suggested, for example, that such inter-layer coupling could occur between the 1D metallic behavior of a stripe and the  $\vec{q}$ -vector in the neighboring stripe<sup>37</sup>. Unfortunately, from our diffraction measurements it is not possible to say anything about nearest-neighbor  $c$ -axis coupling due to the  $90^\circ$  rotation of the stripes. A second possibility is that the isotropy is driven by poorly screened (long range) Coulomb interactions, as has been suggested in the context of a ladder compound<sup>54</sup>. Finally, and most prosaically, it is possible that the charge stripe correlations are limited by intrinsic crystallographic disorder, for example due to defects arising from randomly distributed dopant ions<sup>55</sup>. As such order is presumably random, one would expect it to be isotropic, and that it would therefore cause an isotropic correlation length. We will return to the role of crystallographic order again shortly in the discussion of modulation amplitudes.

### B. Modulation Amplitude

The result that the electronic charge stripe modulation in LBCO is 10 times larger in amplitude than in LNSCO is surprising. Taken at face value it implies a very large

difference in the value of the charge order parameter in these two systems and appears to contradict expectations based on measurements of the ordered magnetic moments in the two systems. For example,  $\mu$ SR measurements show similar ordered moments of  $0.3 \mu_B$  with similarly large magnetically-ordered volume fractions in LBCO and LNSCO<sup>56</sup>. Since the magnetic order is constrained by the charge order, it would seem counterintuitive for the charge order parameters to be so different when the magnetic order parameters are so similar. Further, a recent estimate of the hole concentration per Cu site in the charge carrier stripes in  $\text{La}_{1.675}\text{Eu}_{0.2}\text{Sr}_{0.125}\text{CuO}_4$  (LESCO) resulted in a value considerably larger than 0.2 holes<sup>46</sup>. Based on the similarity of LESCO and LNSCO, in terms of correlation length and temperature dependence, one might expect LESCO and LNSCO to have similar hole modulations. Such an expectation is then, at first glance, inconsistent with the present measurement, which suggests LBCO is 10 times larger, which would be an unphysical result. A previous estimate of the integrated hole weight in LBCO, for example, suggested it is 0.59 holes<sup>42</sup>. However, the LESCO estimate came from an analysis of the resonant lineshape, which reflects the amplitude of the modulation in the modulated regions. There are a number of reasons why a measurement based on the diffraction signal could lead to a different result. Firstly, the ordered volume of sample could be different for LBCO and LNSCO, with LNSCO having an ordered volume up to 100 times smaller than that of LBCO. This seems extremely unlikely. A second possibility is that these diffraction measurements do not fully capture all the intensity of the stripe correlations and that in LNSCO there is significant intensity outside the detected region in reciprocal space - i.e. in diffuse tails. Missing this intensity would result in a much reduced apparent hole amplitude as measured in LNSCO compared with LBCO. This effect would not be reflected in the resonant lineshape analysis. Such an explanation is perhaps consistent with the fact that the charge stripes are more poorly correlated in LNSCO as compared to LBCO. However, at this point, it is not possible to know which, if any, of these explanations is the correct one.

We now address the differences between LBCO and LNSCO, searching for hints as to the origin of the differences in stripe amplitude. First, the average structures of LBCO and LNSCO are remarkably similar<sup>58,59</sup>. Secondly, the electron-phonon coupling for the Cu-O bond stretching mode appears to be the same<sup>60</sup>. Finally, the average A-site cation radius is again very similar: 1.232 Å and 1.211 Å for LBCO and LNSCO, respectively<sup>61</sup>. What then is the origin of the difference in the two systems? Local pinning of stripe domains (as occurs with Zn doping<sup>62</sup>, for example) can cause neighboring stripe domains to be out of phase with one another, and destructive interference may limit correlation lengths and average amplitudes. The orthorhombic strains (associated with the octahedral tilts) at temperatures just above the LTT transition are twice as large in LNSCO<sup>57</sup>

as in LBCO<sup>44</sup> and are a candidate for the origin of this pinning. Indeed in the ordered phase the LTT tilt angle is larger in LNSCO than in LBCO ( $4.4^\circ$  compared to  $3.5^\circ$ , see Ref. 58) and it has previously been shown that the electronic properties are sensitive to this degree of tilt, with a critical value of  $\Phi_c = 3.6^\circ$  below which the electronic properties were unaffected<sup>63</sup>. By analogy with perovskite manganites<sup>64</sup>, we may expect that the larger modulations of the Cu-O-Cu bond angles resulting from this tilt will be associated with a greater tendency to localize charge and perturb the stripe modulations, especially in the vicinity of divalent dopant ions. Thus, the reduced average tilt angle in LBCO might explain the longer correlation lengths and enhanced modulation amplitudes of the periodic stripes.

### C. Incommensurability

Here, we address the incommensurability, that is the deviation from the ideal value of  $\delta = 0.25$  at this doping. In each of these systems, we find  $\delta = 0.24$ . In LNSCO this deviation was previously attributed to a mixture of two different commensurate stripe spacings<sup>57</sup>. A random ordering of these components gave a good match to the spin and lattice incommensurabilities and correlation lengths, as determined with neutrons. However, the considerably longer charge and lattice correlation lengths in LBCO compared to LNSCO together with the same incommensurability in the two systems, is not compatible with this random-ordering model. Within a model of two integral stripe spacings, correlated ordering of the two components would be necessary to explain these results. This is beyond the scope of the present work.

## V. CONCLUSIONS

In summary, our comparison of the charge and lattice modulations in LBCO and LNSCO shows that the in-plane correlations of the charge stripes in these two systems are isotropic, and a factor of two larger in LBCO. These two systems show different temperature dependencies, with the integrated intensity of the charge modulation in LBCO constant up to the LTT transition, while in LNSCO it decreases on warming, disappearing at  $\sim 55$  K, well below the LTT transition in this material. Comparing the amplitudes of the lattice modulation we find that the lattice modulation in LBCO is 4 times stronger than in LNSCO. Performing the same comparison on the electronic modulation suggests that the electronic modulation, as measured by diffraction is 10 times stronger in LBCO when compared with LNSCO. This result, based on measurement of the integrated intensity of the stripe modulation, appears at odds with measurements performed by resonant lineshape analysis. It is clear that further work is required to solve this puzzle, and we hope that this work encourages such measurements.

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- <sup>33</sup> J. M. Tranquada, J. D. Axe, N. Ichikawa, *et al.*, Phys Rev Lett, **78**, 338 (1997).
- <sup>34</sup> A. R. Moodenbaugh, Y. Xu, M. Suenaga, *et al.*, Phys. Rev. B, **38**, 4596 (1988).
- <sup>35</sup> M. Fujita, H. Goka, K. Yamada, *et al.*, Phys. Rev. B, **70**, 104517 (2004).
- <sup>36</sup> S. Kivelson, E. Fradkin, and V. Emery, Nature, **393**, 550 (1998).
- <sup>37</sup> V. J. Emery, S. A. Kivelson, and O. Zachar, Phys. Rev. B, **56**, 6120 (1997).
- <sup>38</sup> Q. Li, M. Hücker, G. D. Gu, *et al.*, Phys Rev Lett, **99**, 067001 (2007).
- <sup>39</sup> J. M. Tranquada, G. D. Gu, M. Hücker, *et al.*, Phys. Rev. B, **78**, 174529 (2008).
- <sup>40</sup> M. J. Lawler, K. Fujita, J. Lee, *et al.*, Nature, **466**, 347 (2010).
- <sup>41</sup> R. Daou, J. Chang, D. Leboeuf, *et al.*, Nature, **463**, 519 (2010).
- <sup>42</sup> P. Abbamonte, A. Rusydi, S. Smadici, *et al.*, Nature Phys., **1**, 155 (2005).
- <sup>43</sup> Y.-J. Kim, G. D. Gu, T. Gog, *et al.*, Phys. Rev. B, **77**, 064520 (2008).
- <sup>44</sup> M. Hücker, M. von Zimmermann, G. Gu, *et al.*, Phys. Rev. B, **83**, 104506 (2011).
- <sup>45</sup> J. Tranquada, B. Sternlieb, J. D. Axe, *et al.*, Nature, **375**, 561 (1995).
- <sup>46</sup> J. Fink, E. Schierle, E. Weschke, *et al.*, Phys. Rev. B, **79**, 100502 (2009).
- <sup>47</sup> J. Fink, V. Soltwisch, J. Geck, *et al.*, Phys. Rev. B, **83**, 092503 (2011).
- <sup>48</sup> We note that in Refs. 46 and 47 the correlation length was defined as  $\xi = \frac{2\pi}{\text{HWHM}}$  resulting in reported correlation lengths being larger by a factor of  $2\pi$  than presented in the present work. The definition in the present work is consistent with previous x-ray and neutron work.
- <sup>49</sup> C. T. Chen, F. Sette, Y. Ma, *et al.*, Phys Rev Lett, **66**, 104 (1991).
- <sup>50</sup> H. Romberg, M. Alexander, N. Nücker, *et al.*, Phys. Rev. B, **42**, 8768 (1990).
- <sup>51</sup> S. Partzsch and S. B. Wilkins, Unpublished.
- <sup>52</sup> J. M. Tranquada, J. D. Axe, N. Ichikawa, *et al.*, Phys. Rev. B, **54**, 7489 (1996).
- <sup>53</sup> M. von Zimmermann, A. Vigliante, T. Niemoller, *et al.*, Europhys. Lett., **41**, 629 (1998).
- <sup>54</sup> A. Rusydi, P. Abbamonte, H. Eisaki, *et al.*, Phys Rev Lett, **97**, 016403 (2006).
- <sup>55</sup> L. M. Rodriguez-Martinez and J. P. Attfield, Phys. Rev. B, **54**, R15622 (1996).
- <sup>56</sup> B. Nachumi, Y. Fudamoto, A. Keren, *et al.*, Phys. Rev. B, **58**, 8760 (1998).
- <sup>57</sup> J. M. Tranquada, N. Ichikawa, and S. Uchida, Phys. Rev. B, **59**, 14712 (1999).
- <sup>58</sup> J. D. Axe and M. K. Crawford, Journal of Low Temperature Physics, **95**, 271 (1994).
- <sup>59</sup> S. Katano, J. Fernandez-Baca, S. Funahashi, *et al.*, Physica C: Superconductivity, **214**, 64 (1993).
- <sup>60</sup> D. Reznik, L. Pintschovius, M. Ito, *et al.*, Nature, **440**, 1170 (2006).
- <sup>61</sup> R. D. Shannon, Acta Crystallographica Section A, **32**, 751 (1976).
- <sup>62</sup> H. Kimura, M. Kofu, Y. Matsumoto, *et al.*, Phys Rev Lett, **91**, 067002 (2003).
- <sup>63</sup> B. Buchner, M. Breuer, A. Freimuth, *et al.*, Phys Rev Lett, **73**, 1841 (1994).
- <sup>64</sup> P. G. Radaelli, M. Marezio, H. Y. Hwang, *et al.*, Phys. Rev. B, **54**, 8992 (1996).
- <sup>33</sup> J. M. Tranquada, J. D. Axe, N. Ichikawa, *et al.*, Phys Rev Lett, **78**, 338 (1997).
- <sup>34</sup> A. R. Moodenbaugh, Y. Xu, M. Suenaga, *et al.*, Phys. Rev. B, **38**, 4596 (1988).
- <sup>35</sup> M. Fujita, H. Goka, K. Yamada, *et al.*, Phys. Rev. B, **70**, 104517 (2004).
- <sup>36</sup> S. Kivelson, E. Fradkin, and V. Emery, Nature, **393**, 550 (1998).
- <sup>37</sup> V. J. Emery, S. A. Kivelson, and O. Zachar, Phys. Rev. B, **56**, 6120 (1997).
- <sup>38</sup> Q. Li, M. Hücker, G. D. Gu, *et al.*, Phys Rev Lett, **99**, 067001 (2007).
- <sup>39</sup> J. M. Tranquada, G. D. Gu, M. Hücker, *et al.*, Phys. Rev. B, **78**, 174529 (2008).
- <sup>40</sup> M. J. Lawler, K. Fujita, J. Lee, *et al.*, Nature, **466**, 347 (2010).
- <sup>41</sup> R. Daou, J. Chang, D. Leboeuf, *et al.*, Nature, **463**, 519 (2010).
- <sup>42</sup> P. Abbamonte, A. Rusydi, S. Smadici, *et al.*, Nature Phys., **1**, 155 (2005).
- <sup>43</sup> Y.-J. Kim, G. D. Gu, T. Gog, *et al.*, Phys. Rev. B, **77**, 064520 (2008).
- <sup>44</sup> M. Hücker, M. von Zimmermann, G. Gu, *et al.*, Phys. Rev. B, **83**, 104506 (2011).
- <sup>45</sup> J. Tranquada, B. Sternlieb, J. D. Axe, *et al.*, Nature, **375**, 561 (1995).
- <sup>46</sup> J. Fink, E. Schierle, E. Weschke, *et al.*, Phys. Rev. B, **79**, 100502 (2009).
- <sup>47</sup> J. Fink, V. Soltwisch, J. Geck, *et al.*, Phys. Rev. B, **83**, 092503 (2011).
- <sup>48</sup> We note that in Refs. 46 and 47 the correlation length was defined as  $\xi = \frac{2\pi}{\text{HWHM}}$  resulting in reported correlation lengths being larger by a factor of  $2\pi$  than presented in the present work. The definition in the present work is consistent with previous x-ray and neutron work.
- <sup>49</sup> C. T. Chen, F. Sette, Y. Ma, *et al.*, Phys Rev Lett, **66**, 104 (1991).
- <sup>50</sup> H. Romberg, M. Alexander, N. Nücker, *et al.*, Phys. Rev. B, **42**, 8768 (1990).
- <sup>51</sup> S. Partzsch and S. B. Wilkins, Unpublished.
- <sup>52</sup> J. M. Tranquada, J. D. Axe, N. Ichikawa, *et al.*, Phys. Rev. B, **54**, 7489 (1996).
- <sup>53</sup> M. von Zimmermann, A. Vigliante, T. Niemoller, *et al.*,

- Europhys. Lett., **41**, 629 (1998).
- <sup>54</sup> A. Rusydi, P. Abbamonte, H. Eisaki, *et al.*, Phys Rev Lett, **97**, 016403 (2006).
- <sup>55</sup> L. M. Rodriguez-Martinez and J. P. Attfield, Phys. Rev. B, **54**, R15622 (1996).
- <sup>56</sup> B. Nachumi, Y. Fudamoto, A. Keren, *et al.*, Phys. Rev. B, **58**, 8760 (1998).
- <sup>57</sup> J. M. Tranquada, N. Ichikawa, and S. Uchida, Phys. Rev. B, **59**, 14712 (1999).
- <sup>58</sup> J. D. Axe and M. K. Crawford, Journal of Low Temperature Physics, **95**, 271 (1994).
- <sup>59</sup> S. Katano, J. Fernandez-Baca, S. Funahashi, *et al.*, Physica C: Superconductivity, **214**, 64 (1993).
- <sup>60</sup> D. Reznik, L. Pintschovius, M. Ito, *et al.*, Nature, **440**, 1170 (2006).
- <sup>61</sup> R. D. Shannon, Acta Crystallographica Section A, **32**, 751 (1976).
- <sup>62</sup> H. Kimura, M. Kofu, Y. Matsumoto, *et al.*, Phys Rev Lett, **91**, 067002 (2003).
- <sup>63</sup> B. Buchner, M. Breuer, A. Freimuth, *et al.*, Phys Rev Lett, **73**, 1841 (1994).
- <sup>64</sup> P. G. Radaelli, M. Marezio, H. Y. Hwang, *et al.*, Phys. Rev. B, **54**, 8992 (1996).