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Charge Density Waves Tuned by Biaxial Tensile Stress

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The electronic orders appearing in condensed matter systems originate from the precise arrangement and nature of crystal atoms. This tenuous relationship can lead to highly different phases, and drive electronic phase transitions. To explore this rich physics, we developed a new device to perform true biaxial mechanical deformation of layered materials at cryogenic temperatures, compatible with x-ray diffraction and transport measurements in the same sample. Here, using this device, we show that a slight deformation of TbTe_3 can have a dramatic influence on the stabilized electronic order. We report on a Charge Density Wave (CDW) orientational transition from \vec{c} to \vec{a} directly driven by the a/c structural parameter, with the occurrence of a tiny coexistence region near $a = c$, without space group change. The CDW transition temperature T_c displays a linear dependence with $|a/c - 1|$, without saturation under deformation while the gap saturates out of the coexistence region. This behaviour is well accounted for within a tight-binding model. Our results question the relationship between the gap and T_c in RTe_3 systems. More generally, our method can be applied to many systems displaying electronic phase transitions and opens a new route towards the study of coexisting or competing electronic orders in condensed matter.

Charge Density Waves (CDW) have raised considerable attention for decades due to their peculiar properties [1, 2], but the interest for this phase significantly in-

creased recently due to its competition with superconductivity in various systems [3]. As they are very sensitive to electron-phonon coupling, application of strain is key to tune their electronic properties. The methods to play on structural parameters are diverse : physical and chemical pressure, epitaxial strain in thin films etc. Recently, application of direct mechanical deformation using piezoelectric actuators [4] was successfully applied in quantum materials to drive electronic transitions. Elastoresistivity measurements performed under uniaxial stress allowed to study nematic susceptibility in iron pnictides [5–8] or heavy fermion materials [9]. Although compatible with cryogenic temperatures, devices applying uniaxial stress have an intrinsic limitation in terms of flexibility, as only one direction of strain is controlled in the sample. It is for instance impossible to get an increase of the lattice parameters in two directions of the crystal. In addition, the true sample deformation is generally not directly measured.

Here, we present results obtained by true biaxial mechanical deformation of quasi-2D materials at cryogenic temperatures, and probe both electronic and structural parameters using resistivity and x-ray diffraction (XRD) measurements in the same sample. To do so, a new device has been developed [10] (see Fig. 1(b)) which allows to measure resistivities along and perpendicular to the deformation axis simultaneously in the temperature range 80-375K (see Fig. 1(c)).

In this work, we focus on TbTe_3 , one of the RTe_3 compounds, where R is a rare-earth element of the Lanthanide family ($R = \text{La, Ce, Pr, Nd, Sm, Gd, Tb, Dy, Ho, Er, Tm}$) [12]. These systems are quasi-tetragonal (see Fig. 1(a)), and display a highly rich phase diagram. A first CDW transition appears along \vec{c} in all RTe_3 systems below T_c with wavevector $\vec{Q}_c = (0, 0, \sim \frac{5}{7}c^*)$, and

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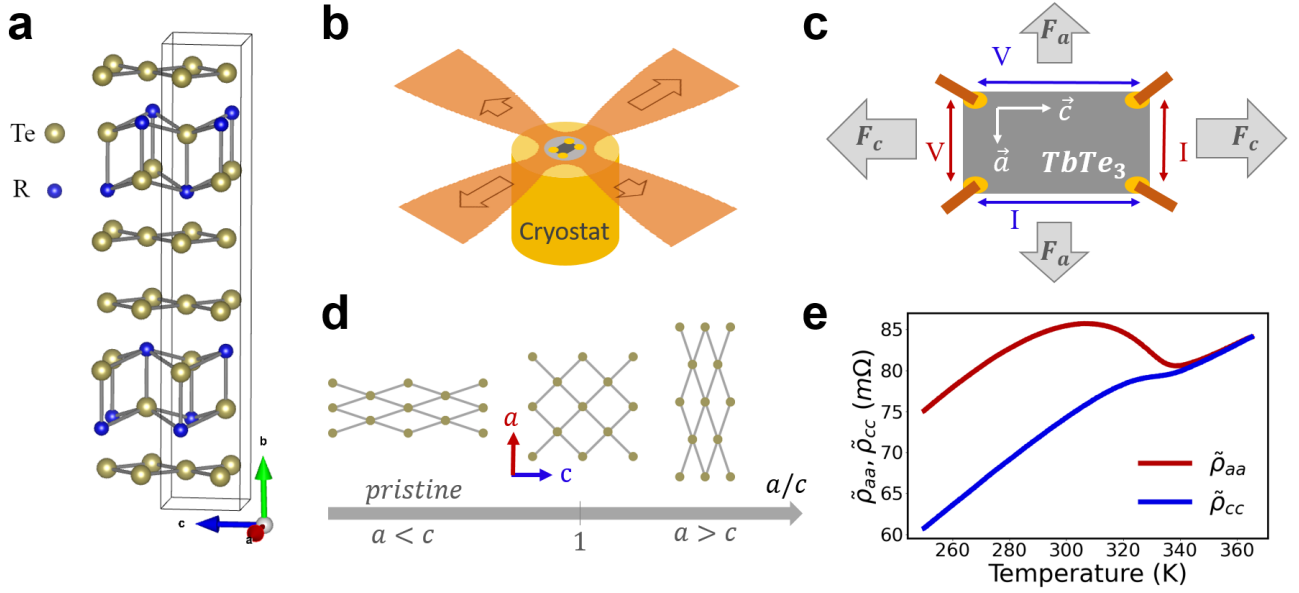


FIG. 1. **Principle of cryogenic biaxial deformation of TbTe_3 .** (a) The RTe_3 crystal structure is quasi-tetragonal (unit cell shown in black), and made of a succession of RTe slab intercalated by two quasi-square Te planes in the (\vec{a}, \vec{c}) planes. The two in-plane directions are non-equivalent due to the presence of a glide plane along \vec{c} (space group $Cmcm$) which leads to a slight orthorhombicity ($1 - \frac{a}{c} \sim 1.3 \cdot 10^{-3}$ in TbTe_3 at 300K [11]). A view of the nearly-square Te sheet in the (\vec{a}, \vec{c}) plane is shown on the side. (b) Schematic drawing of the biaxial tensile stress device: a thin crystalline sample is glued on a polyimide cross-shaped substrate on which tensile stresses are applied along the arms. The main in-plane directions of the crystal \vec{a} and \vec{c} are aligned with the arms of the cross. The bottom side of the cross is in contact with the cold finger of a nitrogen flow cryostat. (c) Four electrical contacts are deposited at the four corners of the sample, allowing to apply currents and measure voltages along \vec{a} or \vec{c} directions of the crystal. The forces applied along the opposite arms of the cross have same magnitude, both along \vec{a} and \vec{c} (\vec{F}_a and \vec{F}_c respectively). (d) Schematic view of crystal deformation as a function of a/c structural parameter. (e) Temperature dependence of $\tilde{\rho}_{aa} = \rho_{aa}/d$ and $\tilde{\rho}_{cc} = \rho_{cc}/d$ (where d is the sample thickness) in the pristine state of sample 1 extracted using the Montgomery technique (see Methods). They display the typical features observed in TbTe_3 compounds *i.e.* a linear behaviour in the normal state, above T_c , and a resistivity jump below $T_c \sim 337 \pm 1\text{K}$ when entering the CDW phase.

68 a second one at lower temperature T_{c2} with wavevector 92
 69 $\vec{Q}_a = (\sim \frac{5}{7}a^*, 0, 0)$ for the heavier R elements (R = Tb, 93
 70 Dy, Ho, Er and Tm) [11]. Both CDW are incommen- 94
 71 surate with the underlying lattice period [11, 13, 14]. A 95
 72 magnetic phase appears at low temperature for the heav- 96
 73 iest R as well as SC under pressure at $\sim 1\text{K}$ for some
 74 compounds in the series [15, 16].

75 Recently, elastoresistivity and elastocaloric measure- 97
 76 ments performed under uniaxial stress in ErTe_3 and 98
 77 TmTe_3 suggested a possible CDW orientational switch-
 78 ing from \vec{c} to \vec{a} [17]. A significant change of T_{c2} was also 99
 79 reported when the sample is deformed along \vec{a} , while only 100
 80 a very slight change of T_c was observed. However, in the 101
 81 latter study, the change of a and c lattice parameters, 102
 82 could not be measured or changed simultaneously and, 103
 83 resistivity measurements are only performed along the 104
 84 applied stress direction, which prevents to get the full in- 105
 85 formation on both CDW in all experimental conditions. 106

86 Here, we use biaxial in-plane tensile stress to indepen- 107
 87 dently apply controlled deformation and measure resis- 108
 88 tivities along both crystallographic axis \vec{a} and \vec{c} of TbTe_3 110
 89 (see Fig. 1(c)). The resistivities obtained in the pristine 111
 90 state of sample 1, are presented in Fig. 1(e), where $\tilde{\rho}$ in- 112
 91 dicates resistivity ρ divided by sample thickness. $\tilde{\rho}_{aa}$ is 113

particularly sensitive to the transition towards the CDW
 along \vec{c} due to the gap opening at Fermi level in this
 direction that annihilates electronic states with veloci-
 ties along \vec{a} . This consequently induces a much larger
 increase of $\tilde{\rho}_{aa}$ than $\tilde{\rho}_{cc}$ [18].

I. STRUCTURAL EVOLUTION OF LATTICE AND CDW UNDER BIAXIAL STRESS

XRD measurements were performed to follow the
 Bragg peaks associated to the main crystal structure
 during deformation (see Methods), as well as satellite
 peaks associated to the CDWs along \vec{a} and \vec{c} (referred
 to as CDW_a and CDW_c in the following). Three non-
 colinear Bragg reflections (0 16 0, 0 16 1 and 1 15 0)
 were measured to retrieve the three lattice parameters at
 all forces F_a and F_c applied along \vec{a} and \vec{c} respectively
 (see Fig 2(a)). Their evolution is plotted as a function
 of F_a and $-F_c$ in Fig. 2(b). a and c follow a quasi-linear
 behaviour when applying uniaxial forces with an in-plane
 Poisson ratio $\nu_{ac} \sim 1$ while b decreases both when apply-
 ing forces F_a and F_c with an out-of-plane Poisson ratio
 $\nu_{ab} \sim \nu_{bc} \sim 0.1$. This strong difference can be explained

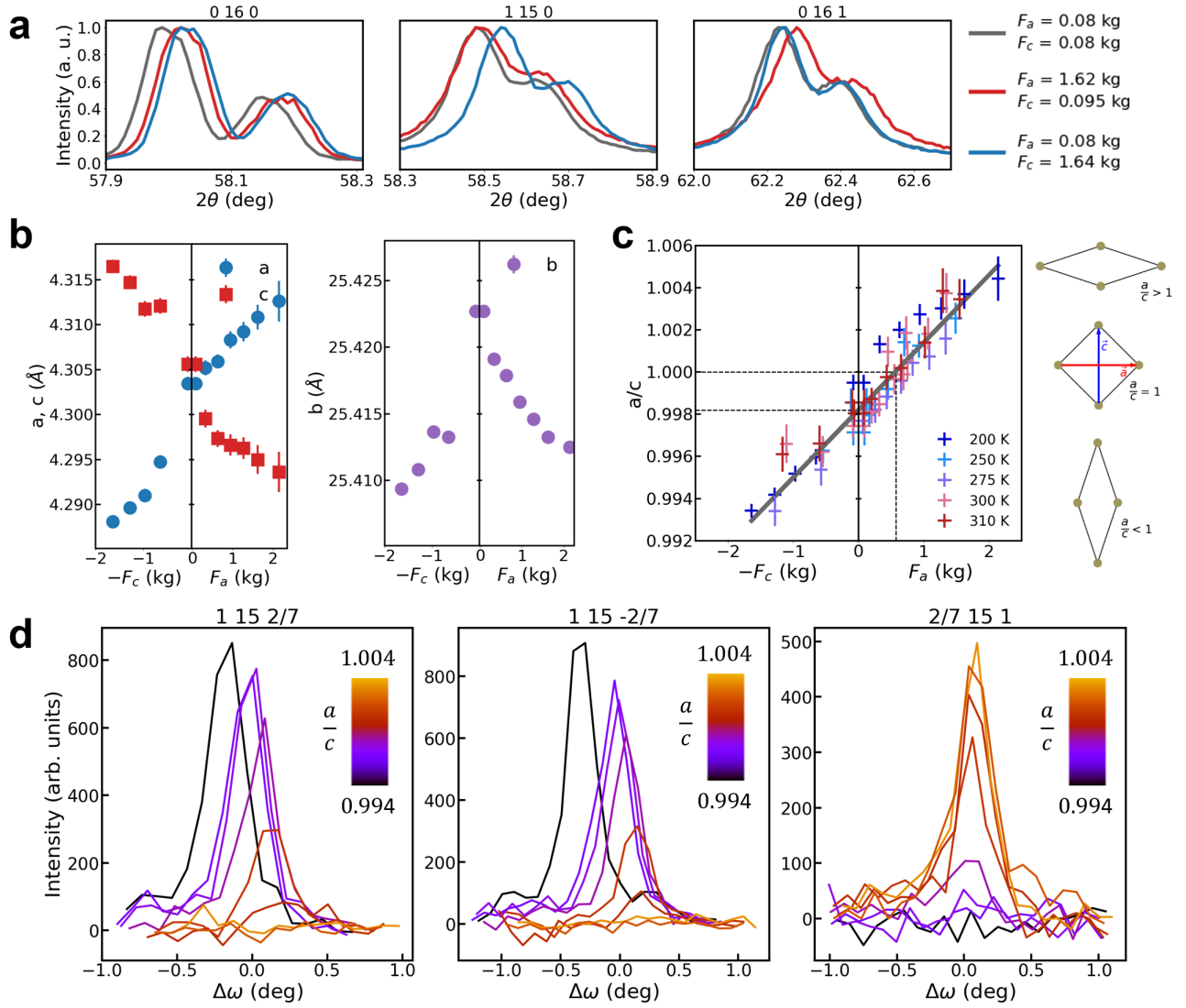


FIG. 2. **Lattice and CDW probed by XRD under uniaxial stresses.** (a) 0 16 0, 1 15 0 and 0 16 1 Bragg peak intensities normalized to 1, plotted along the longitudinal 2θ direction, for 3 sets of forces: without applied force (grey curves), with $F_a = 1.62$ kg (red curves) and $F_c = 1.64$ kg (blue curves). Each Bragg reflection display the K_{α_1} and K_{α_2} components of the x-ray source; (b) Evolution of the in-plane and out-of-plane lattice constants of TbTe₃, obtained from the fit of the 3 non-colinear Bragg peaks shown in (a) as a function of F_a and $-F_c$. $\Delta a/a \sim -\Delta c/c \sim 0.3\%$ and $\Delta b/b \sim -0.03\%$ at maximum deformation which corresponds to an in-plane (resp. out-of-plane) Poisson ratio $\nu_{ac} \sim 1$ (resp. $\nu_{ab} = \nu_{cb} \sim 0.1$). (c) Evolution of the a/c ratio as a function of the same forces, at several temperatures between 200K and 310K. The grey line is obtained by a linear fit of all data points: $a/c = \alpha F + \beta$, with $F = F_a$ for $F > 0$ and $F = -F_c$ for $F < 0$, $\alpha = 0.0032\text{kg}^{-1}$ and $\beta = (a/c)_{F=0} \sim 0.9982 \pm 0.001$ which is consistent with the expected a/c ratio in the pristine state. $a/c = 1$ is obtained for $F_a \sim 0.6 \pm 0.1$ kg. (d) Rocking scans on the 1 15 2/7, 1 15 -2/7 peaks associated to the CDW along \vec{c} and 2/7 15 1 peak associated to the CDW along \vec{a} , as a function of a/c ratio, at $T=250$ K. The a/c ratio was computed from the uniaxial forces F_a and F_c using the linear fit shown in (c). The rocking angle ω is taken relative to the peak position in the pristine state.

114 by the weak van der Waals coupling between layers along 122
 115 \vec{b} . The evolution of the a/c ratio is plotted as a function 123
 116 of uniaxial forces in Fig 2(c) for several temperatures. It 124
 117 evolves linearly as a function of applied uniaxial force, 125
 118 and this direct correlation is used to express all relevant 126
 119 quantities in terms of a/c in the following. 127

120 The CDW reflections can also be tracked as a function 128
 121 of applied force by XRD. Indeed, as CDWs add a new 129

periodicity in the system, additional reflections appear around lattice Bragg reflections and provide important information about the static or dynamical CDW structure [19–22]. The intensity of those satellites is related to the amplitude of the periodic lattice distortion (PLD), their position relative to the Bragg peak position gives the CDW wavevector, and their width is linked to its correlation length. Here, we measured the 1 15 $\pm 2/7$

satellite reflections associated to CDW_c as a function of applied stress, as well as the $2/7$ 15 1 associated to CDW_a as it was shown to appear transiently after laser excitation [23] and suggested to appear under mechanical stress [17]. The rocking curves obtained on these satellites at $T=250K$ are presented in Fig. 2(d) as a function of a/c . In the pristine state, the 1 15 $\pm 2/7$ display a single peak, while the $2/7$ 15 1 is completely absent, expected from a sample without twin domains. When a/c decreases down to the lowest value reached here, the 1 15 $\pm 2/7$ intensities increase with no clear change of width while the $2/7$ 15 1 is still absent. On the contrary, when a/c increases, the 1 15 $\pm 2/7$ peak intensities decrease until complete disappearance at the highest $a/c = 1.004$. Concomitantly, the $2/7$ 15 1 peak increases up to $\sim 2/3$ of the maximum intensity of the 1 15 $\pm 2/7$ with a similar width as the 1 15 $2/7$ (~ 0.5 deg). Thus, when a/c increases, CDW_c progressively disappears while CDW_a appears with a similar correlation length as the previous CDW_c , with an intermediate coexistence region.

II. RESISTIVITIES UNDER BIAxIAL STRESS

$\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ were obtained at several uniaxial forces between 250K and 375K, well below and above T_c and are plotted in Fig 3(a-c) as a function of a/c for $0.993 < a/c < 1.004$.

When $a/c = 0.993$, $\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ are similar to the ones found in the pristine state, but T_c is shifted to higher temperatures. When a/c increases, the resistivity jump of $\tilde{\rho}_{aa}$ (resp. $\tilde{\rho}_{cc}$) at T_c decreases (resp. increases) until $\tilde{\rho}_{aa}$ is similar to the initial shape of $\tilde{\rho}_{cc}$, and inversely. In addition, all curves shift first to lower temperatures then to higher ones. These features are well seen on the anisotropy curves $\frac{\rho_{aa}}{\rho_{cc}}$ that tends to one for $T > T_c$, and displays a jump below T_c . This jump is positive for low a/c values, and negative for high ones. The evolution of T_c is spotted by dots in the inset of Fig. 3(c).

The results obtained in the case of equibiaxial deformation are highly interesting. Indeed, no clear change is observed on $\tilde{\rho}_{aa}$, $\tilde{\rho}_{cc}$ and anisotropy in this case (see Fig. 3(d-e)) meaning that changing the absolute values of a and c keeping the a/c ratio constant does not affect the pristine CDW_c state.

III. A TRANSITION AT $a = c$

The evolution of the main parameters measured by XRD and resistivity data are plotted as a function of the a/c parameter in Fig 4. More specifically, in Fig 4(a), we computed the integrated intensities of the 1 15 $2/7$ and $2/7$ 15 1 satellite reflections shown in Fig 2. As previously described, the intensity of the 1 15 $2/7$ decreases while $2/7$ 15 1 increases when a/c increases from 0.994 to 1.004, following a sigmoid shape for the $2/7$ 15 1 and an inverse-sigmoid shape for 1 15 $2/7$, having the same width, and

both centered at the same position $a/c = 1$ (within the error bars). The saturation value of the $2/7$ 15 1 is $\sim 2/3$ that of the saturation value of the 1 15 $2/7$, and both go down to zero when they reach their minimum value. When $a/c < 0.999$ (resp. $a/c > 1.002$), we only measure a satellite along \vec{c} (resp. \vec{a}). We thus observe a continuous transformation of CDW_c into CDW_a when increasing the a/c ratio, an intermediate coexistence phase for $0.9985 < a/c < 1.002$, and a crossing point at $a/c = 1$.

The same kind of plot has been performed for the resistivity jump $\Delta\tilde{\rho}$ obtained from $\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ by taking the maximum value of $\tilde{\rho} - \tilde{\rho}_n$, where $\tilde{\rho}_n$ is the linear resistivity in the normal state ($T > T_c$). The values $\Delta\tilde{\rho}_{aa}$ and $\Delta\tilde{\rho}_{cc}$ obtained with this method are shown in Fig 4(b). The shape of both curves is identical to the ones obtained for the satellite intensities in Fig. 4(a): $\Delta\tilde{\rho}_{aa}$ has an inverse-sigmoid shape and $\Delta\tilde{\rho}_{cc}$ a sigmoid shape, and the same widths. They also cross around $a/c = 1$ (within error bars), with an intermediate region in the range $0.9985 < a/c < 1.0005$. Also, similarly to the satellite intensity curves, the saturation value obtained for $\Delta\tilde{\rho}_{cc}$ is $\sim 2/3$ the saturation value of $\Delta\tilde{\rho}_{aa}$. In contrast, the resistivity jumps are constant for equibiaxial deformations (see Fig. 3(f)).

Finally, a value of T_c has been extracted from the resistivity curves by taking the second derivative of the resistivity curves and spotting the local peak corresponding to the inflection point of the resistivities (see Supp. Mat.). By doing so, the T_c obtained from $\tilde{\rho}_{aa}$, $\tilde{\rho}_{cc}$ and anisotropies are consistent. The evolution of T_c is plotted as a function of a/c in Fig. 4(c). T_c decreases linearly for $a/c < 1$, and then increases linearly again for $a/c > 1$, with a minimum value of 330K found at $a/c = 1$. T_c was increased by 30K compared to the pristine state, but could potentially go on increasing to much larger values if the a/c parameter could be explored in a wider range of values. The linear dependence is consistent with the behaviour reported in [17], but with a much greater amplitude. Interestingly, in the case of equibiaxial deformations, T_c is found to be constant within error bars (see Fig. 3(f)).

IV. DISCUSSION

All the results presented above show that the key parameter for the CDW orientational switching from \vec{c} to \vec{a} is the structural parameter a/c . This result is of prime importance as it shows the direct relationship between the crystal structure and the appearance of CDWs in $TbTe_3$, and presumably in all RTe_3 systems. Indeed a hint of this transition was also reported in $TmTe_3$ and $ErTe_3$ in [17]. Our results also show that the glide plane does not play any role in the CDW stabilisation as it is still present during crystal deformation - reflections forbidden by symmetry never appeared in our experiments. When $a = c$, both CDW_c and CDW_a coexist, as revealed by XRD and transport measurements. Those results are

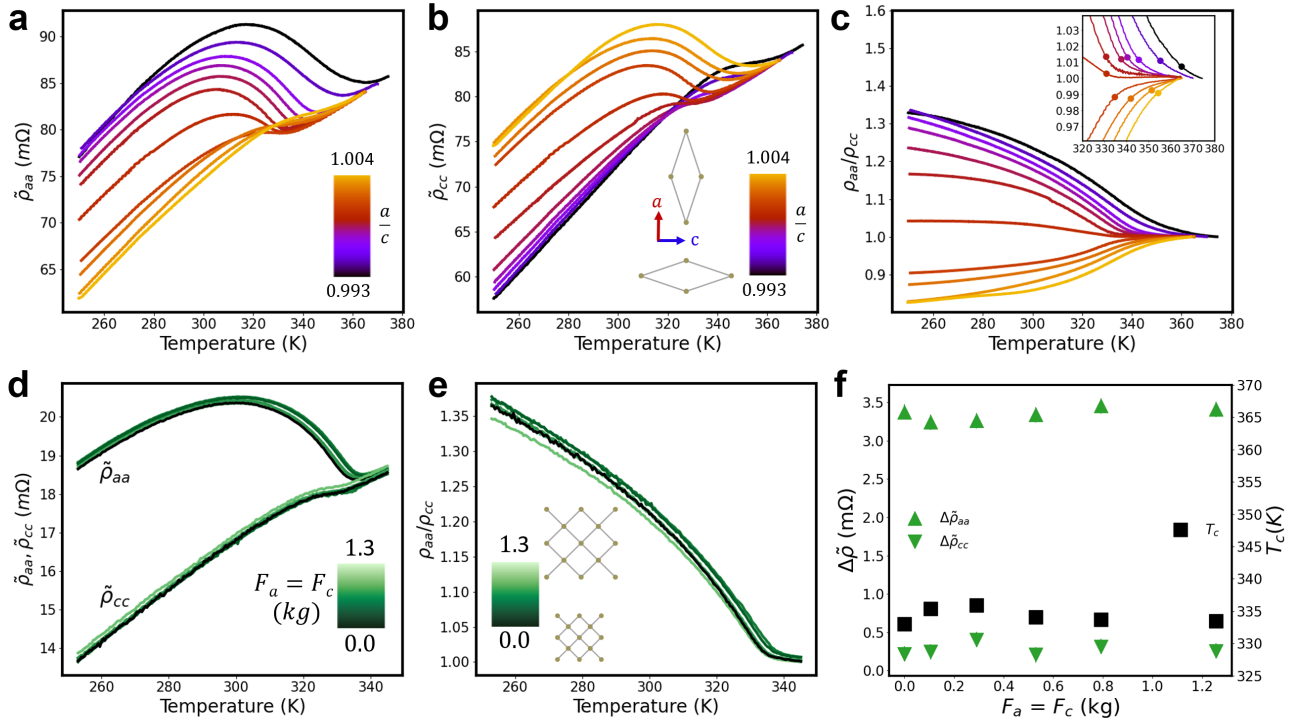


FIG. 3. **Evolution of CDW transition under uniaxial and biaxial stress through resistivity measurements.** (a) $\tilde{\rho}_{aa}$ and (b) $\tilde{\rho}_{cc}$ resistivity curves obtained along \tilde{a} and \tilde{c} respectively for uniaxial measurements, between 250K and 375K, as a function of a/c , varied between 0.993 and 1.004 in Sample 1. (c) Anisotropy ρ_{aa}/ρ_{cc} in the same temperature range, obtained from the curves shown in (a) and (b). Inset: zoom on the anisotropy curves. The dots indicate the position of T_c extracted from each anisotropy curve. (d) $\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ resistivity curves obtained by equibiaxial deformation in sample 2, as a function of temperature. The equibiaxial deformation is depicted by the sketch on the left. (e) Anisotropy curves extracted from the resistivity curves shown in (d) as a function of temperature. (f) Evolution of resistivity jumps (green triangles) and T_c (black squares) for equibiaxial deformations, plotted with the same scale as uniaxial data presented in Fig 4. No change of these parameters take place under equibiaxial deformation. Note that sample 2 behaves exactly as sample 1 under uniaxial stress, with an inversion of $\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ for $F_a > 0.9$ kg (see Supp. Mat.).

extremely reproducible, both when changing the a/c ratio from one phase to another, and also from one sample to another (see Suppl. Mat.).

In XRD experiments, the intensity of the satellite reflections associated to the CDW is generally related to the amplitude of the PLD, itself proportional to the CDW gap squared. Here, the intensity of the satellite associated to CDW_c decreases when a/c increases, while the one associated to CDW_a increases. The fact that we do not observe any change of width, within the resolution of the present measurement, does not support the appearance of domains during the transition from CDW_c to CDW_a. The observed behaviour suggests that the amplitude of the PLD along \tilde{c} vanishes while the one along \tilde{a} grows, with the same correlation length for CDW_c and CDW_a. Consequently the gap must close along \tilde{c} and appear along \tilde{a} with a mixed state when $a = c$.

This is consistent with the resistivity measurements. Indeed, the amplitude of the jumps $\Delta\tilde{\rho}_{aa}$ and $\Delta\tilde{\rho}_{cc}$ are related to the gap along \tilde{c} and \tilde{a} respectively. This is again consistent with a gap closure along \tilde{c} when a/c increases while it opens along \tilde{a} , with a mixed state in

the region $a = c$. The inversion of anisotropy, that is also associated to the gap, is also a proof of this gap switching. In the pure CDW_a phase (for $a/c > 1.002$), XRD and transport data consistently suggest that the gap of the CDW_a phase is lower than the one in the pure CDW_c phase. The gap that opens along \tilde{a} should thus be smaller than the one along \tilde{c} in the CDW_c phase. All this should appear in ARPES experiments upon tensile deformation of the sample along \tilde{a} .

The coexistence region $0.999 < a/c < 1.002$ demonstrates that a mixed state can be stabilized, with gaps in both directions. It should be investigated whether they appear in separate domains in the sample, in different Te planes, or if they are superimposed in the same Te planes.

The present results can be compared to the behaviour observed in hydrostatic pressure experiments or with chemical pressure obtained by changing the rare-earth element, and which were shown to induce similar effects on the CDWs [24, 25]: when the lattice parameters are compressed, the CDW wavevector, the gap and T_c evolve similarly in both cases. However, here, when a and c

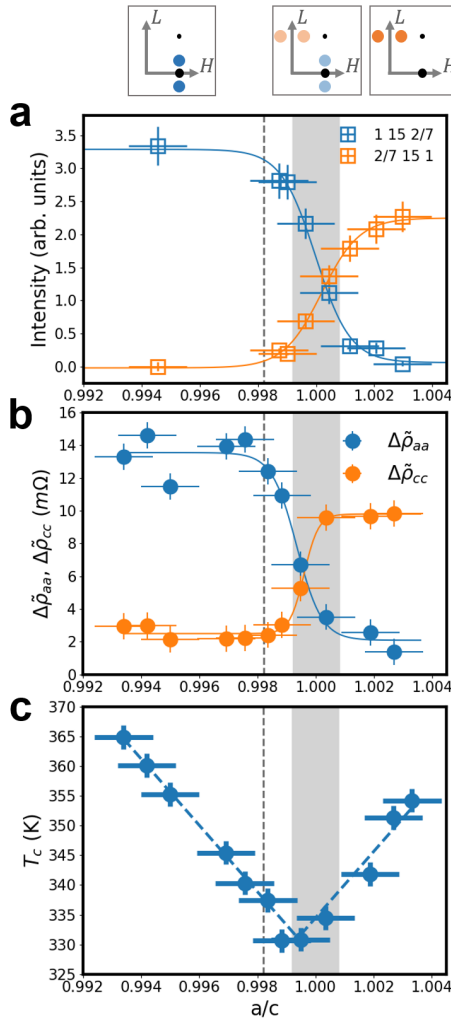


FIG. 4. **Distinct evolution of CDW gap and T_c under mechanical deformation.** (a) Evolution of the 1 15 2/7 (blue squares) and 2/7 15 1 (orange squares) satellite intensities measured as a function of a/c . The blue (resp. orange) solid line is an inverse-sigmoid (resp. sigmoid) fit to the experimental points obtained on the 1 15 2/7 (resp. 2/7 15 1) satellite reflection. The three sketches in the top represent a (H 15 L) plane in reciprocal space. The big (resp. small) black dot is the 1 15 0 (resp. 1 15 1) Bragg reflection, and the blue (resp. orange) dots are the 1 15 $\pm 2/7$ (resp. $\pm 2/7$ 15 1) reflections, with a color strength depending on the measured intensity in each a/c region. (b) Resistivity jumps $\Delta\tilde{\rho}_{aa}$ (blue dots) and $\Delta\tilde{\rho}_{cc}$ (orange dots) extracted from the experimental curves shown in Fig. 3, plotted as a function of a/c (see method in Supp. Mat.). The blue (orange) solid line is an inverse-sigmoid (resp. sigmoid) fit of $\Delta\tilde{\rho}_{aa}$ (resp. $\Delta\tilde{\rho}_{cc}$). (c) T_c computed from the resistivity curve taking the average of the local peaks found in the second derivative of $\tilde{\rho}_{aa}$, $\tilde{\rho}_{cc}$ and ρ_{aa}/ρ_{cc} as a function of a/c (see the T_c determination method Suppl. Mat.). For all three panels, the dashed gray line is the a/c value in the pristine state, and the gray filled region marks the position $a/c = 1$ with a width equal to the error bar ($\pm 0.8 \cdot 10^{-3}$).

tensile stress, no change in resistivity is observed. We conclude that the gap and T_c do not change in the negative pressure region. To observe variations of gap and T_c , the parameter a/c has to change. In the experiments reported in [24], the parameter a/c changes and tends to 1 when pressure is applied (above 3GPa in CeTe₃). In these conditions, T_c decreases, which is in qualitative agreement with our observations: T_c is minimum when $a/c = 1$. Quantitatively, the variation of T_c reported in [24] is much larger than reported here. Concerning the evolution of the gap, they also report vanishing CDW satellite intensities, suggesting a disappearance of the gap at high pressures, while $a/c = 1$. Here, no change is observed in equibiaxial deformation, but the experiments should be repeated in equibiaxial conditions while $a/c = 1$. The role of the relative variation of the b parameter should also be investigated further to compare our experiment to hydrostatic pressure measurements.

The behaviour observed here is very similar to the one observed after fs laser excitation reported in [23], with the CDW appearing along \vec{a} while the one along \vec{c} disappears. Contrary to the equilibrium CDWs at low temperatures in some RTe₃ compounds, the CDWs appearing along a and c in this reference and in our study have only a tiny coexistence region. This makes the new CDW _{a} unique and different from the equilibrium coexistence of CDW _{c} and CDW _{a} at low temperatures. Following the results presented in [23], the lattice parameters of LaTe₃ after laser pulse excitation does not vary by more than 0.02% when the transient CDW _{a} appears, which indicates that the driving mechanism is different from the one we report here. In our case, both CDWs are stable as long as the a/c parameter is kept constant.

Finally, the most striking feature is the evolution of T_c compared to the evolution of the gap, as extracted from XRD and resistivity measurements. Both satellite intensities and $\Delta\tilde{\rho}$ saturate when $a/c < 0.9985$ and $a/c > 1.002$, and therefore the gap values saturate in this region. However, T_c keeps diverging linearly, which questions its relation with the gap. The linear dependence of T_c with respect to a/c can be understood within the tight-binding model, with good quantitative agreement. The model predicts an increase of T_c $\delta T_c \approx 26$ K for $\delta a/a \approx 0.2\%$, which is comparable to the experimental values reported here. The same applies to deformations along \vec{c} , and explains the linear dependence of T_c as a function of deformation. The system actually chooses the CDW along the axis with the highest T_c . For $a > c$, T_c is higher for CDW _{a} , while for $a < c$, T_c is higher for CDW _{c} . At $a = c$, the transition temperatures are equal, and the system is degenerate.

In this article, we report *in-situ* x-ray diffraction and transport to follow the structural and electronic properties of the CDWs in TbTe₃ during biaxial mechanical deformation between 250K and 375K. By directly measuring both CDW satellite intensities and resistivities along \vec{a} and \vec{c} , we show that the CDW orientation continuously flips from \vec{c} to \vec{a} when the a/c ratio is increased, with a

are increased by the same amount applying equibiaxial

coexistence of both orientations when $a = c$. As those two parameters are linked to the gap, we can infer that the gap position changes from one band to another, and that both bands are gapped when $a = c$. Moreover, the transition temperature displays a linear behaviour with respect to a/c , with a minimum at $a = c$, which is well accounted for by a 2D tight-binding model in the Te planes, with good quantitative match between theory and experiment. For the pure CDW_a and CDW_c phases, the gap saturates while T_c does not, which questions the relationship between those quantities in RTe₃ systems. Our study opens new perspectives for the exploration of many electronic phase transitions in condensed matter systems with application of biaxial tensile stress at cryogenic temperatures.

V. METHODS

Samples: The TbTe₃ samples were grown by the self-flux method, as described in [13]. All samples were cut to have an in-plane rectangular shape, and were mechanically exfoliated down to few μm thickness to get homogeneous deformation in the volume. Results obtained on two different TbTe₃ single-crystals are presented in the main text. Sample 1 was probed both by XRD and transport to get the full structural and electronic properties, with uniaxial deformation, and sample 2 was probed by transport under uniaxial and biaxial deformation. The clearly different behaviour of ρ_{aa} and ρ_{cc} allowed us to directly determine the crystal orientations and T_c (see Supp. Mat.) [18]. Other samples were measured by XRD or transport and give the same results, as presented in Supp. Mat. Sample 1 (resp. sample 2) has a 1:1.17 (resp. 1:2) in-plane aspect ratio, and $2.5\mu\text{m}$ (resp. $10\mu\text{m}$) thickness.

Biaxial Tensile stress device: The principle of the new device developed here is similar to the one described in [26], with the sample glued on a deformable substrate that is mechanically stretched, and adapted to host a nitrogen-flow cryostat to reach temperatures in the range 80-375K. The samples are glued at the center of a $125\mu\text{m}$ -thick polyimide cross-shaped substrate, with \vec{a} and \vec{c} in-plane directions aligned with the arms of the polyimide cross. The four arms of this substrate are attached to four independent motors that can pull on each branch separately. The forces applied along the four arms are measured using calibrated force gauges, and are given in kg. The center of the cross is covered with a thin gold layer on its bottom surface and lies on the cold finger of a Konti-Micro cryostat from CryoVac GmbH, allowing to reach temperatures in the range 80-375K. Apiezon grease is used to get a good thermal transfer between the cold finger and the polyimide cross. In practice, the same motor displacement is used for opposite arms to keep the sample at the same position in the device. This whole setup is enclosed in a vacuum chamber for cryogenic operation, and allows access for incoming and

outgoing x-rays to perform x-ray diffraction in reflection geometry through a $300\mu\text{m}$ -thick Polyether-ether-ketone dome. Four wires are also available in the cryostat for 4-point transport measurements.

XRD measurements: The experiment shown in the main text were performed at LPS with a 8keV x-ray beam generated by a Cu rotating anode source (Rigaku RU-300B), equipped with a multilayer monochromator suppressing the K_β emission line but allowing the K_{α_1} and K_{α_2} ones. The sample mounted in the biaxial tensile stress device was positioned at the center of rotation of a Huber Eulerian 4-circle diffractometer to perform wide-angle x-ray diffraction, and detection was performed with a 2D detector (Timepix from ASI) located 82 cm after the sample. The 3 Bragg reflections (0 16 0, 1 15 0 and 0 16 1) recorded on the 2D detector were projected along the 2θ direction of reciprocal space to compute the three lattice parameters a , b and c for each set of applied forces (see Suppl. Mat.). The absence of twin domains was demonstrated by checking that the forbidden 0 15 1 reflection was indeed not measurable.

Transport measurements: Four contacts are deposited at the four corners of the crystal, and the Montgomery method [27–30] is used to get both $\tilde{\rho}_{aa}$ and $\tilde{\rho}_{cc}$ resistivities along the a and c respectively by applying a fixed current with a Keithley 2611 Sourcemeter, and measuring the voltage with a Keithley 2182a Nanovoltmeter. For each measurement, the forces F_a and F_c were changed at 375K, and the resistances were measured during cooling down to 250K at a fixed rate of 1.5K/min. An electronic device allowed us to switch I^+ and V^- every second to get R_{aa} and R_{cc} during the same temperature ramp. The ratio of transverse dimensions of the sample was obtained by fixing the anisotropy ρ_{aa}/ρ_{cc} to c^2/a^2 in the normal state and used to compute the resistivities normalized by sample thickness d : $\tilde{\rho}_{aa} = \rho_{aa}/d$ and $\tilde{\rho}_{cc} = \rho_{cc}/d$. See Supp. Mat. for detailed formalism. Resistivity measurements presented in Fig. 3(a)-(c) were performed in Sample 1 after measuring the XRD data, and the ones in Fig. 3(d)-(f) in Sample 2. See Supp. Mat. for complete resistivity measurements in Sample 2.

Tight-binding Model: In this description, the electron dispersion of the two bands formed by the p_x and p_z Te orbitals ε_\pm keeps its form (where x and z are along \vec{a} and \vec{c} respectively):

$$\varepsilon_\pm = -2t_\parallel \cos(k_x^* \pm k_z^*) - 2t_\perp \cos(k_x^* \mp k_z^*) \quad (1)$$

where $k_x^* = k_x a/2$ and $k_z^* = k_z c/2$, and t_\parallel and t_\perp are the transfer integrals parallel and perpendicular to the orbital direction. Both lattice parameters and transfer integrals depend on the applied stress. At $a = c$, Eq. 1 coincides with the accepted tight-binding dispersion given by Eq.(1) in [31]. The CDW transition temperature is given by the condition :

$$U(\vec{Q}_0) \chi(T_c, \vec{Q}_0) = 1 \quad (2)$$

448 where the static electron-electron interaction $U(\vec{Q})$ in-472
 449 cludes both Coulomb and phonon-mediated interaction, 473
 450 and the Lindhard susceptibility reads : 474

$$\chi(T_c, \vec{Q}) = \sum_{\alpha, \alpha'} \sum_{k_x, k_z} 16 \frac{n_F(E_{\vec{k}, \alpha}) - n_F(E_{\vec{k}+\vec{Q}, \alpha'})}{E_{\vec{k}+\vec{Q}, \alpha'} - E_{\vec{k}, \alpha}} \quad (3)$$

451 where $n_F(\varepsilon) = 1/(1 + \exp[(\varepsilon - E_F)/T])$ is the Fermi-
 452 Dirac distribution function, $\alpha, \alpha' = \pm$ label the subbands
 453 and $E_{\vec{k}, \alpha}$ differs from Eq 1 only near the intersection
 454 points of two bands in momentum space, as given by 477
 455 Eq.(2) in [32]. As ε_{\pm} and $E_{\vec{k}, \pm}$ depend only on k_x^* and 478
 456 k_z^* , and as $\chi(T_c, \vec{Q})$ has a $x \leftrightarrow z$ symmetry, an equi-479
 457 axial stress does not affect the Lindhard susceptibility. 480
 458 The main difference in the transition temperatures of 481
 459 the CDW along \vec{a} and \vec{c} thus comes from the difference 482
 460 of $U(\vec{Q})$ when $a \neq c$. The electron-electron coupling 483
 461 mainly comes from the Coulomb interaction, screened by 485
 462 conducting electrons: $U(r) = e^2 \exp(-\zeta r)/r$, where the 486
 463 inverse Debye screening radius $\zeta = \sqrt{4\pi e^2 \rho_F}$ and ρ_F 487
 464 is the density of states at the Fermi level. In TbTe₃, 488
 465 $\zeta \approx a^{-1} \approx (4.3)^{-1} \text{ \AA}^{-1}$. The Fourier transform of the
 466 screened Coulomb potential is :

$$U(\vec{Q}) \approx \frac{4\pi e^2}{\vec{Q}^2 + \zeta^2} \quad (4)$$

467 The CDW wavevector \vec{Q}_0 is given by the maximum
 468 of the Lindhard susceptibility, Eq. 3, and gives a fixed
 469 product $Q_{0x}a = Q_{0z}c \approx 10\pi/7$, when $a = c$. An increase
 470 of the lattice constant a decreases the CDW wavevector
 471 Q_{0x} and increases the CDW coupling $U(\vec{Q}_0)$ according

to Eq. 4. An increase of a by $\delta a/a = 0.1\%$ results in a
 decrease of Q_{0x}^2 by 0.2%. Since $\zeta \ll Q_{0x}$, according to
 Eq. 4, this gives an increase of $U(\vec{Q}_0)$ by 0.2%. Accord-
 475 ing to Eq. 2, an increase δU of $U(\vec{Q}_0)$ raises the CDW
 transition temperature T_c by δT_c given by Eq. 2:

$$\frac{\delta U}{U} = \frac{-\delta \chi}{\chi} = \frac{d\chi(T, \vec{Q}_0)}{dT} \frac{-\delta T_c}{\chi(T, \vec{Q}_0)} \quad (5)$$

because the electron susceptibility $\chi(T, \vec{Q}_0)$, approxi-
 mately given by Eq. 3 decreases with the increase of T .
 The temperature-dependence of the Lindhard suscepti-
 bility was calculated in [32] for the second CDW in ErTe₃
 and in [33] for the Q-dependence of the first CDW for var-
 ious parameters of electron dispersion. To find $d\chi/dT$ for
 the first CDW in relevant temperature range and for the
 transfer integrals $t_{\parallel} \approx 2eV$, $t_{\perp} \approx 0.37eV$ and the Fermi
 energy $E_F \approx 1.48eV$ in TbTe₃ [31], we performed new
 calculations of $\chi(T, \vec{Q}_0)$ using Eq. 3 (see Supp. Mat.),
 from which the slope of the temperature-dependent sus-
 ceptibility can be extracted :

$$\eta = \frac{d[\ln \chi(T, \vec{Q}_0)]}{dT} = \chi^{-1} \frac{d\chi}{dT} \approx -1.5 \cdot 10^{-4} K^{-1} \quad (6)$$

We thus can compute δT_c for an increase $\delta a/a \approx 0.2\%$,
 comparable to the experiment described here:

$$\delta T_c = \frac{\delta U(Q_0)}{\eta U(Q_0)} \approx -\frac{2\delta a}{a\eta} \approx \frac{4 \cdot 10^{-3}}{1.5 \cdot 10^{-4}} \approx 26K \quad (7)$$

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VI. CONTRIBUTIONS

A.H.-A. and P.M. grew and provided the TbTe₃ samples. A.A.S. and A.G.-F. prepared the samples for transport and x-ray experiments. A.G.-F. and V.L.R.J. performed and analyzed the laboratory XRD experiments. A.G.-F., A.A.S. and V.L.R.J. performed and analyzed the transport measurements. A.G.-F., A.A.S., D.G., L.O., V.L.R.J., D.L.B., P.Go., P.-O.R., D.T. and E.B. participated in the experiments at synchrotron SOLEIL. P.Gr. developed the theoretical part. V.L.R.J. wrote the initial draft and all authors participated in the discussion and correction of the manuscript. V.L.R.J. and D.L.B. led the project.