The relationship between vacancy ordering and magnetism in TlFe\textsubscript{1.6}Se\textsubscript{2} has been investigated via single crystal neutron diffraction, nuclear forward scattering, and transmission electron microscopy. The examination of chemically and structurally homogenous crystals allows the true ground state to be revealed, which is characterized by Fe moments lying in the ab-plane below 100 K. This is in sharp contrast to crystals containing regions of order and disorder, where a competition between c-axis and ab-plane orientations of the moments is observed. The properties of partially-disordered TlFe\textsubscript{1.6}Se\textsubscript{2} are therefore not associated with solely the ordered or disordered regions. This contrasts the viewpoint that phase separation results in independent physical properties in intercalated iron selenides, suggesting a coupling between ordered and disordered regions may play an important role in the superconducting analogues.

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The microscopic coexistence of superconductivity and magnetism at temperatures up to \(\sim 30\,\text{K}\) in \(\text{A}_{1-y}\text{Fe}_{2-y}\text{Se}_2\) (\(A = \text{K, Cs, Rb}\)) has generated much interest, and a plethora of techniques have been employed to understand the interesting properties of these materials.\[1\]–\[4\] Structural studies on superconducting samples revealed antiferromagnetic order with a local magnetic moment of \(\sim 3\mu_B/\text{Fe}\), and a \(\sqrt{5}\times\sqrt{5}\) superstructure of the ThCr\textsubscript{2}Si\textsubscript{2} structure-type derived from Fe vacancy ordering.\[2\]–\[4\] Other studies have focused on the electronic structure\[5\]–\[7\] and the superconducting pairing mechanism,\[8\]–\[13\] as well as the influence of phase separation.\[11\]–\[19\]

The coexistence of superconductivity and magnetism is now believed to be due to fine-scale phase separation, with each property being associated with a different composition and/or degree of order.\[16\]–\[21\] Phase separation over length scales of \(\sim 10-100\,\text{nm}\) has been observed via transmission electron microscopy (TEM),\[14\]–\[16\] and is inferred from Mössbauer spectroscopy.\[17\] Superconducting samples that are single phase (structurally and chemically homogenous) have yet to be produced and thus the inherent nature of the individual phases remains unclear. Flexibility in the chemical composition near the parent \(A_0\text{Fe}_{1.6}\text{Se}_2\) allows superconductivity to be realized,\[22\]–\[27\] and may do so by promoting phase separation via local inhomogeneity. It has been suggested that the superconducting phase does not have Fe vacancies.\[19\]–\[28\] Similar to other Fe-based superconductors, these superconducting phases may be characterized by a doping level of approximately 0.15 electrons per Fe, such as \(\text{Rb}_{0.3}\text{Fe}_2\text{Se}_2\). The degree of order is also important,\[19\]–\[31\] however, and first principles calculations have shown that the vacancy and magnetic order greatly influence the Fermi surface.\[6\]

Fe vacancies order into the \(\sqrt{5a}\times\sqrt{5a}\) supercell between 460 and 580 K. At or slightly below this temperature, the spins align along the tetragonal \(c\)-axis in a block-checkerboard antiferromagnetic structure (BCAF-\(c\), where \(c\) indicates the alignment of the spins).\[2\]–\[4\] The vacancy order is generally incomplete, however, with partial occupancies observed on at least one Fe site.\[32\]–\[34\]

Superconductivity, which is believed to exist in a phase that does not contain vacancy order, is only observed in samples that also possess regions of vacancy and magnetic order.

In contrast to the alkali-metal based compounds, TlFe\textsubscript{1.6}Se\textsubscript{2} has a limited compositional window and the Tl site is always fully occupied, which reduces the degree of inherent disorder. While superconductivity is observed for mixed Tl / alkali-metal compounds, and was reported with a small volume fraction in TlFe\textsubscript{1.7}Se\textsubscript{2},\[23\] it has not been reproduced in TlFe\textsubscript{1.6}Se\textsubscript{2}. This is perhaps associated with an improper electron count in TlFe\textsubscript{1.6}Se\textsubscript{2}. Based on the chemical formula, one would expect TlFe\textsubscript{1.6}Se\textsubscript{2} to be a metal. However, TlFe\textsubscript{1.6}Se\textsubscript{2} is observed to be an insulator. TlFe\textsubscript{1.6}Se\textsubscript{2} also displays different magnetic behavior than the alkali-metal analogues.

The BCAF-\(c\) structure is the only magnetic structure observed in the alkali-metal compounds. However, single crystals of ‘partially-disordered TlFe\textsubscript{1.6}Se\textsubscript{2}’ have been shown to possess additional magnetic phase transitions near 100 K and 140 K.\[34\] A detailed structural study revealed these transitions are associated with a canting of the Fe moments toward the \(ab\)-plane, which only occurs between \(\sim 100\) and 150 K.\[35\] These crystals are characterized by crystallographically coherent regions of or-
In the absence of single-phase crystals, it is impossible to know if these magnetic transitions in partially-disordered TiFe$_{1.6}$Se$_2$ are inherent to the ordered or disordered regions, or are the result of some interaction between the two.

In this Letter, we provide a detailed characterization of single crystal TiFe$_{1.6}$Se$_2$ with complete chemical/vacancy order. Such structural and chemical homogeneity has been elusive in A$_{1-x}$Fe$_{2-x}$Se$_2$, and the presence of multiple phases has complicated the structure-property investigations in these complex materials. By obtaining ideal order, a previously unobserved spin reorientation is revealed with spins lying in the $ab$-plane for $T < 100$ K. This suggests a strong interaction between the ordered and disordered regions prevents this ground state from occurring in the partially-disordered crystals at low temperatures.

Crystals with complete vacancy order were grown in melts of nominal composition TiFe$_{1.6}$Se$_2$. In the same furnace, a nominal composition of TiFe$_{1.7}$Se$_2$ produced crystals with magnetization behavior qualitatively similar to the ‘partially disordered’ crystals, which were grown from a melt of nominal composition TiFe$_2$Se$_2$. Therefore, growth in iron-rich environments appears to inhibit the ordering of iron vacancies, perhaps due to minor differences in Fe content that are difficult to detect. In the fully ordered crystals, the vacancy ordering temperature is found to be $\sim 463$ K by differential scanning calorimetry (DSC), and an anomaly in the magnetic susceptibility ($\chi$) was also observed at this temperature. An entropy change of $\sim 7.4$ J/mol-Fe/K was calculated from the DSC data, and this is $\sim 75\%$ of the entropy change expected for the simultaneous ordering of Fe vacancies and Fe magnetic moments. See the Supplemental Information for additional details.

The crystals were observed to be chemically homogenous and fully-ordered via aberration corrected scanning TEM (STEM) using high angular annular dark field (HAADF) imaging. The STEM image shown in Fig. 1a was taken with the electron beam parallel to the [110] direction of the supercell. As a result of this orientation, all of the Fe vacancies are aligned in columns parallel to the electron beam and can be readily viewed as dark spots separated by four iron columns. A crystal structure drawing corresponding to the HAADF image is shown in Fig. 1b to highlight this ideal vacancy ordering, and the atomic layers are labeled for ease of viewing. Specimens viewed along [001] also revealed an ordered and uniform vacancy superstructure, and a selected area diffraction pattern from this orientation is shown in Fig. 1c. The selected area diffraction pattern shows two sets of $\sqrt{5}a \times \sqrt{5}a$ superstructure reflections, indicated by the vectors $q$ and $q'$, due to twinning along the (110) subcell planes. Images of the partially-disordered crystals can be found in Ref. 35. STEM data reveal the disordered regions contain Fe vacancies (modulations in Fe HAADF intensity), and thus these regions are not like the vacancy-free phases being reported in the superconducting analogues.

Refinements of the neutron diffraction data indicate complete occupancy of the Fe1 site (Wyckhoff position 16d), and zero occupation of the Fe2 site (Wyckhoff position 4d), which yields the ideal $\sqrt{5}a \times \sqrt{5}a$ superstructure in TiFe$_{1.6}$Se$_2$. In Fig. 2a, the intensity of the vacancy order (020) peak is shown to be independent of temperature below $\sim 460$ K, indicating a saturation of vacancy order at all temperatures probed by neutron diffraction.

The magnetic susceptibility of fully ordered TiFe$_{1.6}$Se$_2$ single crystals is shown in Fig. 2a, where a phase transition is clearly observed near 100 K. Above this transition, the susceptibility is largest when the magnetic field ($H$) is perpendicular to $c$, and for temperatures below the transition the susceptibility is largest for $H \parallel c$. This indicates a switch in the easy magnetic direction, with moments parallel to $c$ above $\sim 100$ K and perpendicular to $c$ below $\sim 100$ K.

The reorientation of the magnetic moment is confirmed via single crystal neutron diffraction (Fig. 2b,c) and synchrotron radiation based nuclear forward scattering (Fig. 3). Figure 2a shows the integrated intensity for three Bragg peaks observed via neutron diffraction on a $\sim 50$ mg single crystal. The intensities of the two peaks with magnetic contributions change rapidly near 100 K. The (121) peak intensity is associated with mo-
ments aligned with the $c$-axis and is beginning to saturate before it quickly diminishes near 100 K. At this temperature, the intensity of the (105) peak increases sharply due to the onset of the new magnetic order. The (105) peak gains intensity when the $ab$-plane component of the moment increases, as well as when the Fe layer develops a corrugation. The moment reorientation is highlighted in Fig. 2, where the refined moments are presented, and reach $\sim 3\mu_B$/Fe in both magnetic structures. A similar moment is obtained via first principles calculations for the BCAF-$c$ (Supplemental Information), and the current calculations are consistent with previous reports.

The canting of the Fe moment in the partially-disordered TFe$_{1.6}$Se$_2$ between 100 and 150 K is shown in Figure 2. A competition between the BCAF-$c$ and an in-plane structure exists in the partially-disordered crystals, where the BCAF-$c$ structure is present below 100 K and above 150 K, and thus the corresponding moment has no $ab$-plane component. When this canting is lost below $\sim 100$ K, an abrupt increase in the $c$-lattice parameter is observed (see Fig. 2). The magnetic transition in the fully ordered crystals is not associated with an anomaly in the $c$ lattice parameter, though, and this is consistent with the smooth behavior in $c$ near $\sim 150$ K where the canting of the moment begins in the partially-disordered crystals.

The moment reorientation is also observed via nuclear forward scattering (NFS), the time-analogue of Mössbauer spectroscopy. NFS measurements were completed at beam line ID18 of the European Synchrotron Radiation Facility, with the crystallographic $c$-axis parallel to the x-ray beam. The spectra were fit by a conventional routine.

The NFS data at temperatures slightly above and below magnetic transition at 100 K are shown in Fig. 3. These results clearly indicate the change of the hyperfine structure across the transition, and confirm the moment orientation changes from parallel to $c$ above 100 K to perpendicular to $c$ below 100 K. At 110 K, the data are well described by a model with AFM Fe moments aligned with the $c$-axis, yielding a magnetic field $H=26.1$ T, a quadrupole splitting of $\Delta E_Q=1.18$ mm/s, and the angle between main axes of those interactions is $46^\circ$. These values are in excellent agreement with those observed for Rb$_{0.8}$Fe$_{1.6}$Se$_2$.

The NFS data at 90 K were fitted with 2 Fe sites with moments aligned in $ab$-plane, and the angle between them is utilized as a fit parameter. This model can generally describe any complex, non-collinear magnetic structure with moments in the plane. Independent of the model utilized, a change of the carrier frequency from that related to the beats of $1^{\text{st}}$ and $6^{\text{th}}$ lines of Mössbauer spectrum (at 110 K) to that related to the beats of $2^{\text{nd}}$ and $5^{\text{th}}$ lines (at 90 K) is observed, indicating a reorientation of the Fe moments. An appearance of the strong $2^{\text{nd}}$ and $5^{\text{th}}$ lines indicates the $ab$-plane alignment of the Fe magnetic moments below 100 K. See the Supplemental Information for further details on NFS data collection and analysis.

Below 100 K, the neutron diffraction data were modeled using several different magnetic structures. First, the eight possible magnetic structures within the expected tetragonal symmetry were examined. Of these, the best fit ($R=0.052$ at 5 K) corresponds to the non-collinear magnetic structure ($I4/m$) shown in Fig. 4, with moments pointing 10.8° away from the vacancies. Fixing the moments towards the vacancies results in a similar quality fit.

Alternative magnetic models require lowering the symmetry of the magnetic structure and assuming equal twin fractions of collinear magnetic domains. The monoclinic model ($I2'/m$) shown in Fig. 4 (and a twin generated by rotating 90 degrees about the $c$-axis) can describe the data well ($R=0.045$ with 2.99(4)$\mu_B$/Fe, at 5 K). Other possible twinned collinear magnetic struc-
data provided an equal mixture of domains rotated 90° from the ferromagnetic structure that describes the neutron diffraction of the crystal structure. (b) An in-plane, block-checker antiferromagnetic structure describes the neutron diffraction data and retains the symmetry of the magnetic structures that describe the neutron diffraction data.

FIG. 4: (color online) The Fe (blue) lattice with different magnetic moments aligned with the c-axis at 110 K, and lying in the ab-plane at 90 K. The insets are the model simulations of Mössbauer spectra, with the lines numbered in the inset of (b).

The transitions at 100 K in the fully-ordered and 140 K in the partially-disordered crystals are similar. In both cases, the moments develop an ab-plane component upon cooling through the transition. The structures in Fig. 4 were not observed via neutron diffraction for the partially-disordered crystals, though the NFS data suggest the Fe moments lie within the ab-plane at 110 K (see Supplemental Information). In either case, the BCAF-c order with spins along c is the ground state in the partially-disordered crystals below 100 K. Therefore, the disordered regions, which are likely to be non-magnetic, have a strong influence on the observed magnetic behavior. Consistent with these results, it has been recently suggested that superconductivity is induced in KFe$_2$Se$_2$ when this KFe$_2$Se$_2$ stoichiometry interacts with the vacancy-ordered K$_{0.8}$Fe$_{1.6}$Se$_2$ phase.

Given the magnetoelastic nature of these materials, it is not surprising that disordered and ordered (non-magnetic and magnetic) regions would be strongly coupled. These regions are crystallographically coherent and approximately 10-20 nm in ab-plane and 2-4 unit cells along c, which allows them to interact via lattice strain. A detailed analysis of the STEM data for partially-disordered TiFe$_{1.6}$Se$_2$ reveals the c lattice parameter is approximately 2% larger in the disordered regions than in the ordered regions, and the change in a is less than ~0.3%. Therefore, modulations in c are the most likely source for the strain interaction.

As demonstrated in this Letter, the absence of disordered regions allows TiFe$_{1.6}$Se$_2$ to complete a spin reorientation at ~100 K, thereby unveiling the true magnetic ground state associated with the $\sqrt{5}a \times \sqrt{5}a$ supercell. This demonstrates a strong influence of the secondary phase on the behavior of the primary phase, and thus an interaction between the ordered and disordered regions determines the properties of the partially-disordered crystals. These results have implications for the superconducting alkali-metal iron selenides, which also possess regions of order and disorder separated on a fine scale. Understanding the interaction between these regions may prove critical in identifying the true nature of superconductivity in those materials.

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