

may also explain the larger extent of quiet Sun spicules<sup>1,2</sup> where p-mode power and granular flows are stronger by up to 50% and magnetic fields are more inclined owing to the presence of opposite polarity<sup>24,25</sup>.

A natural consequence of our model is that the quasi-periodic shocks driving the spicules propagate upwards into the low corona, where they may lead to intensity oscillations with properties that are similar to those of longitudinal oscillations observed by TRACE<sup>26</sup> in 1 MK coronal loops<sup>27</sup>. □

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## A universal scaling relation in high-temperature superconductors

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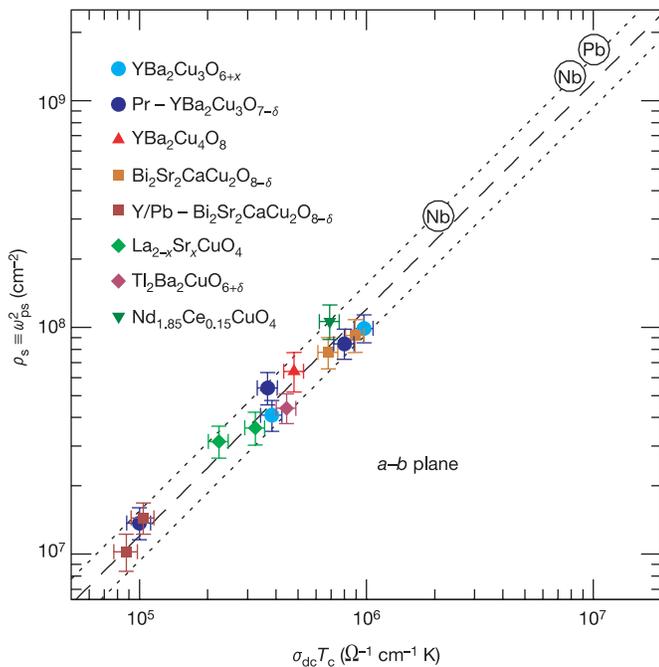
Since the discovery of superconductivity at elevated temperatures in the copper oxide materials<sup>1</sup> there has been a considerable effort to find universal trends and correlations amongst physical quantities, as a clue to the origin of the superconductivity. One of the earliest patterns that emerged was the linear scaling of the superfluid density ( $\rho_s$ ) with the superconducting transition temperature ( $T_c$ ), which marks the onset of phase coherence. This is referred to as the Uemura relation<sup>2</sup>, and it works reasonably well for the underdoped materials. It does not, however, describe optimally doped (where  $T_c$  is a maximum) or overdoped materials<sup>3</sup>. Similarly, an attempt to scale the superfluid density with the d.c. conductivity ( $\sigma_{dc}$ ) was only partially successful<sup>4</sup>. Here we report a simple scaling relation ( $\rho_s \propto \sigma_{dc} T_c$ , with  $\sigma_{dc}$  measured at approximately  $T_c$ ) that holds for all tested high- $T_c$  materials. It holds regardless of doping level, nature of dopant (electrons versus holes), crystal structure and type of disorder<sup>5</sup>, and direction (parallel or perpendicular to the copper-oxygen planes).

We first demonstrate scaling for the  $a$ - $b$  plane (that is, parallel to the copper-oxygen planes) properties<sup>6–12</sup> of single- and double-layer copper oxide materials (Supplementary Table 1), as well as for the conventional metals<sup>13,14</sup> Nb and Pb (elemental superconductors with relatively high values of  $T_c$ ). The values for  $\rho_s$  and  $\sigma_{dc}$  are obtained simultaneously from studies of the reflectance of these materials. The results for the scaling relation are shown on a log-log plot in Fig. 1. The dashed line is a linear fit to the data, while the dotted lines form what are effectively an upper and lower bound for the data; this is described by  $\rho_s = (120 \pm 25)\sigma_{dc} T_c$  (where  $\rho_s$  is in  $\text{cm}^{-2}$ ,  $\sigma_{dc}$  is in  $\Omega^{-1} \text{cm}^{-1}$ , and  $T_c$  is in K). The remarkable result contained in this plot is that within error all of these points fall onto a single line with a slope of unity. This is significant, as the optimally and overdoped materials, which fall well off of the Uemura plot, now scale with the underdoped materials onto a single line.

We also searched for scaling relations along the poorly conducting  $c$  axis, where the charge transport is thought to be incoherent<sup>15</sup>. Previous work focused on scaling between  $\rho_s$  and  $\sigma_{dc}$  only<sup>16,17</sup>. Whereas this approach yields good results for the underdoped materials, in a fashion reminiscent of the Uemura plot, significant deviations from linearity are encountered for optimally and overdoped materials; this was thought to signal the onset of more conventional three-dimensional behaviour. Figure 2 demonstrates

that the *c*-axis data<sup>11,17–20</sup> for all of the single and double-layer materials (Supplementary Table 2) are again well described by a line with slope of unity. What is perhaps most remarkable is that the *a*–*b*-plane and *c*-axis results may all be described by the same universal line shown in Fig. 2, even though the two results correspond to very different ranges of  $\rho_s$ . The combined data span nearly five orders of magnitude, from the insulating behaviour along the *c* axis in the underdoped systems, to the metallic behaviour in the *a*–*b* planes of the overdoped copper oxides.

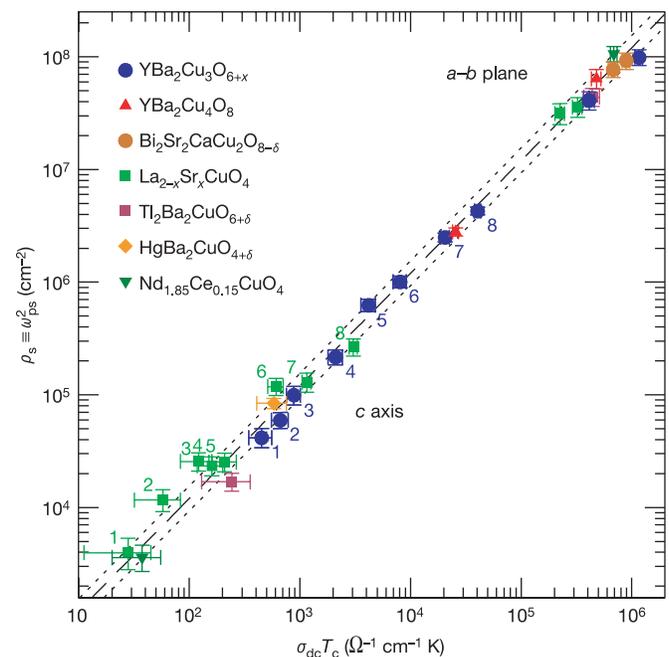
The scaling relation for the *a*–*b* planes can be interpreted in a number of different ways. One of the most direct is the assumption that all of the spectral weight (the area obtained from the integral of the optical conductivity) associated with the free-carriers of the normal state ( $n_n$ ) collapses into the superconducting condensate<sup>21</sup> ( $n_s \equiv n_n$ ) below  $T_c$ . Allowing that the low-frequency conductivity at  $T \approx T_c$  can be described by the simple Drude theory for a metal,



**Figure 1** Plot of the superfluid density ( $\rho_s$ ) versus the product of the d.c. conductivity ( $\sigma_{dc}$ ) and the superconducting transition temperature ( $T_c$ ) for a variety of copper oxides and some simple metals. ( $\sigma_{dc}$  is measured just above the transition, and parallel to the copper–oxygen (*a*–*b*) planes; data are shown on a log–log plot; see Supplementary Table 1 for details, including errors.) The values for  $\sigma_{dc}$  and  $\rho_s$  are obtained from optical measurements of the reflectance. The reflectance is a complex quantity consisting of an amplitude and a phase; in an experiment only the amplitude is usually measured. However, if the reflectance is measured over a wide frequency range, the Kramers–Kronig relation may be used to obtain the phase. Once the complex reflectance is known, then other complex optical functions may be calculated (for example, the dielectric function or the conductivity). The  $\sigma_{dc}$  used in this scaling relation has been extrapolated from the real part of the optical conductivity  $\sigma_{dc} = \sigma_1(\omega \rightarrow 0)$  at  $T \approx T_c$ . For  $T \ll T_c$ , the response of the dielectric function to the formation of a condensate is expressed purely by the real part,  $\epsilon_1(\omega) = \epsilon_\infty - \omega_{ps}^2/\omega^2$ , which allows the superconducting plasma frequency  $\omega_{ps}$  to be calculated from  $\omega_{ps}^2 = -\omega^2 \epsilon_1(\omega)$  in the  $\omega \rightarrow 0$  limit, where  $\omega_{ps}^2 = 4\pi n_s e^2 / m^*$  is proportional to the number of carriers in the condensate. The strength of the condensate ( $\rho_s$ ) is simply  $\rho_s \equiv \omega_{ps}^2$ . The dashed and dotted lines are described by  $\rho_s = (120 \pm 25)\sigma_{dc}T_c$ . Within error, all the data for the copper oxides are described by the dashed line. The data for the conventional superconductors Nb and Pb, indicated by the atomic symbols within the circles, lie slightly above the dashed line.

$\sigma_1(\omega) = \sigma_{dc}/(1 + \omega^2\tau^2)$  (where  $\omega$  is frequency), which has the shape of a Lorentzian centred at zero frequency with a width at half-maximum given by the scattering rate  $1/\tau$ , the area under this curve may be approximated simply as  $\sigma_{dc}/\tau$ . Transport measurements for the copper oxides<sup>22</sup> suggest that  $1/\tau$  near the transition scales linearly with  $T_c$ , so the strength of the condensate is just  $\rho_s \propto \sigma_{dc}T_c$ , in agreement with the observed scaling relation. This result requires that these materials approach the clean limit ( $1/\tau \ll 2\Delta$ , where  $2\Delta$  is the superconducting energy gap).

However, this approach cannot be applied to the properties along the *c* axis, because it is generally conceded that transport in this direction is incoherent, and therefore hopping rather than scattering governs the physics<sup>15</sup>. The quasi-two-dimensional nature of the copper oxides, which often includes a semiconducting or activated response of the resistivity along the *c* axis, has motivated the description of the superconductivity in this direction in terms of a Josephson-coupling picture<sup>16,17,23–26</sup>. The *c*-axis penetration depth  $\lambda$  is then determined by the Josephson current density  $J_c$  and is  $\lambda^2 = \hbar c^2 / 8\pi d e J_c$ , where  $J_c = (\pi \Delta / 2e R_n) \tanh(\Delta / 2k_B T)$ ,  $d$  is the separation between the planes, and  $R_n = d/\sigma_{dc}$  is the normal-state tunnelling resistance<sup>24</sup>. There is convincing evidence that the energy gap in the copper oxides is *d*-wave in nature, containing nodes at the Fermi surface<sup>27,28</sup>, making the determination of  $J_c$  difficult. However, if the coupling between the planes originates at the  $(0, \pi)$ ,  $(\pi, 0)$  points<sup>29</sup> where the gap is a maximum,  $\Delta_0$ , then we can approximate  $\Delta \approx \Delta_0$ . Furthermore, if  $\Delta_0 \propto T_c$ , then  $J_c \propto T_c/R_n$  and  $\rho_s \propto \sigma_{dc}T_c$ , which yields the observed scaling behaviour in the *c*-axis direction. Despite the different nature of the transport properties parallel and perpendicular to the *a*–*b* planes, the universal scaling pertaining to both directions is an unusual and surprising result that should provide new insights into the origins of the superconductivity in these materials. □



**Figure 2** As Fig. 1 but for copper oxides only, and including data for the poorly conducting *c* axis. The values for  $\rho_s$  and  $\sigma_{dc}$  are obtained from optical measurements, as described in Fig. 1 legend. In addition to the published results, new data are also included for  $\text{HgBa}_2\text{CuO}_{4+\delta}$  and  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$ . Within error, all of the data fall on the same universal (dashed) line with slope of unity, defined by  $\rho_s = 120\sigma_{dc}T_c$ ; the dotted lines are from  $\rho_s = (120 \pm 25)\sigma_{dc}T_c$ . See Supplementary Table 2 for details, including errors.

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**Supplementary Information** accompanies this paper on [www.nature.com/nature](http://www.nature.com/nature).

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## Magnetic phase control by an electric field

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The quest for higher data density in information storage is motivating investigations into approaches for manipulating magnetization by means other than magnetic fields. This is evidenced by the recent boom in magnetoelectronics and 'spintronics'<sup>1</sup>, where phenomena such as carrier effects in magnetic semiconductors<sup>2</sup> and high-correlation effects in colossal magnetoresistive compounds<sup>3</sup> are studied for their device potential. The linear magnetoelectric effect—the induction of polarization by a magnetic field and of magnetization by an electric field—provides another route for linking magnetic and electric properties. It was recently discovered that composite materials and magnetic ferroelectrics exhibit magnetoelectric effects that exceed previously known effects<sup>4,5</sup> by orders of magnitude<sup>6–10</sup>, with the potential to trigger magnetic or electric phase transitions. Here we report a system whose magnetic phase can be controlled by an external electric field: ferromagnetic ordering in hexagonal  $HoMnO_3$  is reversibly switched on and off by the applied field via magnetoelectric interactions. We monitor this process using magneto-optical techniques and reveal its microscopic origin by neutron and X-ray diffraction. From our results, we identify basic requirements for other candidate materials to exhibit magnetoelectric phase control.

Hexagonal  $HoMnO_3$  displays ferroelectric ordering at Curie temperature  $T_C = 875$  K (ref. 11), antiferromagnetic  $Mn^{3+}$  ordering at Néel temperature  $T_N = 75$  K (ref. 12), and magnetic  $Ho^{3+}$  ordering at  $T_{Ho} = 4.6$  K (ref. 13). The ferroelectric phase possesses  $P6_3cm$  symmetry and a polarization  $P_z = 5.6 \mu C cm^{-2}$  (ref. 11) along the hexagonal *z* axis. Figure 1 shows that it is made up by three magnetic sublattices with  $Mn^{3+}$  ( $3d^3$ ) ions at 6c positions and  $Ho^{3+}$  ( $4f^{10}$ ) ions at 2a and 4b positions<sup>14</sup>. Anisotropy confines the  $Mn^{3+}$  spins to the basal *x-y* plane where frustration leads to four possible triangular antiferromagnetic structures<sup>12,15</sup>. In contrast, the  $Ho^{3+}$  sublattices are assumed to order Ising-like along *z* showing antiferromagnetism or ferri-/ferromagnetism according to Table 1.

Magnetic  $Mn^{3+}$  ordering was monitored by optical second harmonic generation (SHG) as detailed elsewhere<sup>12,15</sup>: Light at frequency  $\omega$  is incident on a crystal, inducing an electromagnetic polarization at frequency  $2\omega$ , which acts as source for a SHG light wave emitted from the crystal. The magnetic symmetry determines the polarization  $\mathbf{P}(2\omega)$  of the SHG wave relative to that of the fundamental wave at  $\omega$ , so that in turn  $\mathbf{P}(2\omega)$  reveals the underlying arrangement of  $Mn^{3+}$  spins. The relation between SHG polarization and  $Mn^{3+}$  ordering is tabulated elsewhere<sup>12,15</sup>. Magnetic  $Ho^{3+}$  ordering was monitored by neutron powder diffraction and optical Faraday rotation, that is, the rotation  $\Phi \propto \mathbf{B}$  of the plane of polarization of linearly polarized light by the magnetic field  $\mathbf{B}$  in a transmission measurement. The microscopic mechanisms driving magnetoelectric phase control were revealed by neutron and X-ray powder diffraction.

Optical measurements were performed at the MBI on flux-grown, polished, *z*-oriented  $HoMnO_3$  platelets ( $\sim 50 \mu m$  thick) using previously described transmission set-ups<sup>15,16</sup>. The static