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Coherent Phonon Coupling to Individual Bloch States in Photoexcited Bismuth

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We investigate the temporal evolution of the electronic states at the bismuth (111) surface by means of time- and angle-resolved photoelectron spectroscopy. The binding energy of bulklike bands oscillates with the frequency of the A1g phonon mode, whereas surface states are insensitive to the coherent displacement of the lattice. A strong dependence of the oscillation amplitude on the electronic wave vector is correctly reproduced by ab initio calculations of electron-phonon coupling. Besides these oscillations, all the electronic states also display a photoinduced shift towards higher binding energy whose dynamics follows the evolution of the electronic temperature.

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It is well known that most elements crystallize in structures with hexagonal or cubic symmetry. One notable exception is bismuth, which instead crystallizes in the A7 rhombohedral structure. It has been originally proposed by Jones and Peierls that cubic bismuth would be unstable towards a lattice distortion leading to the doubling of the unit cell [1]. Due to the distorted nature of the ground state, the atomic positions are very sensitive to the electronic distribution in the conduction band. This conjecture has been experimentally verified by suddenly changing the occupation number of the electrons via the absorption of a femtosecond laser pulse [2–4]. Due to photoexcitation, a large coherent phonon modulates the distance between the two atoms in the unit cell along the (111) direction. The displacement vector indicates that the lattice coordinates move towards a hypothetical phase with a single atom per unit cell. In reality the material displays a non-thermal melting before the symmetric phase can be attained [5].

The atomic motion following the photoexcitation of bismuth is nowadays well understood. Time resolved X-ray diffraction and density functional theory (DFT) calculations could accurately describe the amplitude of the oscillations and the frequency softening at high-excitation densities [6–8]. On the other hand, the temporal evolution of electronic states has never been directly observed. Some aspects of the electronic dynamics can be inferred from general arguments. The generation mechanism of coherent phonons with large amplitude implies that low-energy electronic excitations are coupled to the A1g mode. It is also expected that the electron-phonon coupling of such distorted material varies with the electronic wave vector. Nonetheless, the determination of the coupling matrix elements for individual Bloch states is still an experimental challenge. We approach this task by time resolved photoelectron spectroscopy measurements. The electronic states of the bismuth (111) surface are characterized by angle-resolved photoemission and first principles calculations. We confirm that the electronic structure displays a rich combination of bulklike bands, surface states, and surface resonances [9,10]. Upon photoexcitation, these electronic states display a dynamics that depends on wave vector and band index. The connection of our data with DFT calculations provides a full characterization of the coupling between electrons and the A1g mode. In addition, we observe that the electronic states do not follow rigidly the motion of the lattice displacement. Instead, the bands also display a purely electronic shift towards higher binding energy.

The experiments were performed with the FemtoARPES setup, using a Ti:sapphire laser that generates 35 fs pulses centered at 810 nm with repetition rate of 250 kHz. Part of the beam is employed to generate the fourth harmonic by frequency doubling in BBO crystals (β-BaB2O4). The 205 nm probe and the 810 nm pump are focused on the sample with a spot diameter of 100 μm and 200 μm, respectively. Their cross-correlation in a BBO crystal has a full width at half maximum (FWHM) of 80 fs. An electrostatic spectrometer with energy resolution better than 10 meV and angular resolution better than 0.5° analyzes the emitted photoelectrons. The overall energy resolution is nonetheless limited to 50 meV by the bandwidth of the 205-nm beam. The (111) surface of bismuth has been obtained by sputtering and annealing cycles of a single crystal. All measurements have been performed at the base temperature of 130 K and at the base pressure of 7 × 10⁻¹¹ mbar. The out-of-equilibrium spectra have been collected with incident pump fluence of 0.6 mJ/cm².

Figure 1(a) shows an intensity map of the photoelectrons emitted along the Γ-M direction of the Brillouin zone.
As previously reported, the breakdown of translational symmetry in the (111) direction generates surface states that intersect the Fermi level and give rise to a Fermi surface [9]. These evanescent wave functions are localized at the topmost layers, thus conferring to the surface of bismuth good metallic properties. Although the band structure supports surface states only near the Fermi level, some bulklike states and surface resonances are clearly visible at higher binding energy [11]. Two bands of different nature are indicated in Fig. 1(a) by letter a and b. Due to the matrix elements of the photoemission process, band b is clearly visible only when the parallel component of the electronic wave vector is \( k_\parallel < 0.25 \, \text{Å}^{-1} \). Band a has larger binding energy, disperses more strongly, and intersects band b at the \( \Gamma \) point. The overall agreement between our measurements and one step photoemission calculations is remarkable [10]. A. Kimura et al. [10] also show that band a has a large spin polarization whereas band b does not. This finding suggests that a is a surface resonance, whereas b is a bulk band. In order to confirm it, we performed DFT calculations of bulk bismuth within the framework of the local density approximation and the ABINIT [12] package. Since the photoemission process does not conserve the perpendicular component of the electronic wave vector, the connection between a peak in the photocurrent and a calculated band is formally not allowed [13]. Nonetheless, Fig. 1(a) shows that the band calculated along the \( \Gamma^\prime-M \) direction for perpendicular wave vector \( k_\perp = 0.45 \, \text{Å}^{-1} \) matches reasonably well the experimental intensity distribution. More insights on the dispersion of band b can be obtained by mapping in the reciprocal space the photoelectrons at fixed kinetic energy. Figure 1(b) shows an intensity map of photoelectrons acquired at different wave vectors for binding energy \( \epsilon = -260 \, \text{meV} \). The structure b is very intense along \( \Gamma^\prime-M^\prime \), whereas it vanishes in the direction \( \Gamma^\prime-M^\prime \). The experimental evidence that band b is invariant for a rotation of 120° but not for a rotation of 60° corroborates the bulk character of band b [9] (see also the Supplemental Material [14]).

Photoexcitation by an intense and ultrafast laser pulse generates sizable effects on the photoelectron current (see movie in the Supplemental Material [14]). Figure 2(a) shows the spectrum acquired at \( k_\parallel = 0.12 \, \text{Å}^{-1} \) along the \( \Gamma-M \) direction for two different pump-probe delays. Note the large reduction of spectral weight taking place 300 fs after the arrival of the pump beam. The partial depletion of states a and b can be approximately described by a Fermi-Dirac distribution with an effective temperature of 2080 K. As shown by Fig. 2(b), the electronic temperature decays

![Intensity map of photoelectrons](image1)

**FIG. 1** (color online). (A) Intensity map of photoelectrons emitted for parallel wave vector along the \( \Gamma-M \) direction. Band a and band b are the surface resonance and the bulklike state, respectively. The blue line is a bulk band calculated for perpendicular wave vector \( k_\perp = 0.45 \, \text{Å}^{-1} \). (B) Intensity map of photoelectrons emitted at binding energy \( \epsilon = -0.26 \, \text{eV} \). An intense structure originating from band b is visible along the \( \Gamma-M \) direction but not along \( \Gamma^\prime-M^\prime \).

![Photoelectron spectra](image2)

**FIG. 2** (color online). (A) Photoelectron spectra acquired 100 fs before and 300 fs after the arrival of the pump beam along the \( \Gamma-M \) direction at \( k_\parallel = 0.12 \, \text{Å}^{-1} \). (B) The effective temperature of photoexcited electrons as a function of pump-probe delay extracted from the data (marks) and simulated with the model of Ref. [8] (red line). (C) Intensity map of photoelectrons emitted at wave vector \( k_\parallel = 0.12 \, \text{Å}^{-1} \) along the \( \Gamma-M \) direction as a function of pump-probe delay.
as a function of pump-probe delay with a time constant of 6 ps. The relaxation of the electronic energy density at the surface is due to the combined effect of electronic diffusion and heat dissipation into the lattice bath. We could accurately reproduce the temporal evolution of the electronic temperature by solving the coupled differential equations in Ref. [8] with incident pump fluence of 0.6 mJ/cm², electron thermal conductivity $\kappa_0 = 0.04$ Wm⁻¹K⁻¹ (fitting parameter), and a temperature-dependent electron-phonon coupling (calculated ab initio).

Besides the large changes of the electronic distribution, we also observe strong variations in the structure of the electronic states. As shown by Fig. 2(a), both peak $a$ and $b$ shift towards higher binding energies after the arrival of the pump pulse. In addition, the position of peak $b$ displays periodic modulations at the frequency of the $A_{1g}$ mode. Such a dynamics arises from the coupling of the coherent phonon to electronic states and can be directly observed on the photoelectron intensity map of Fig. 2(c). The oscillations are large for the bulk band $b$ whereas they fall below the detection limit for the surface resonance $a$ and for all other surface states (see also Supplemental Material [14]). This observation leads to two possible conclusions: either the amplitude of the $A_{1g}$ mode is smaller on the topmost bilayer or the surface states have a weak coupling to this mode.

The excitation of coherent phonons in semimetals has been the subject of many experimental and theoretical works [2–8]. Since bismuth absorbs photons at 810 nm, the generation mechanism is mainly due to a dispersive excitation of the charge density [2,3]. The DFT calculations can help us to illustrate this point: Figure 3(a) displays a color scale plot of the electronic density $\rho(r,T_i)$ at $T_i = 130$ K in the (110) plane. The sudden increase of electronic temperature from 130 K to $T_f = 2000$ K changes the spatial distribution of electronic density. We show in Fig. 3(b) that $[\rho(r,T_f)-\rho(r,T_i)]/\rho(r,T_i)$ is negative in the region between nearest neighbours (atoms 1 and 2). The transfer of electronic charge out of the bonding region increases the repulsion between the ion cores. As a consequence, the atoms oscillate around a new equilibrium position until the energy dissipation into other phonon modes brings the system back to the initial state. The lattice motion changes the binding energy of the Bloch state $|b,k_{||}|$ by $D_{b,k_{||}}u$, where $u$ is the $A_{1g}$ phonon coordinate and $D_{b,k_{||}}$ is the deformation potential. Figure 3(c) shows the binding energy of peak $b$ at the base temperature of 130 K as a function of pump-probe delay (see also Supplemental Material [14]). The oscillations induced by the $A_{1g}$ mode have frequency $\nu = 2.97 \pm 0.05$ THz and damping time $\gamma = 2.6 \pm 0.2$ ps. These values compare well with the frequency and damping time observed by transient reflectivity in comparable experimental conditions [4]. We reproduce the oscillations of peak $b$ by solving the model proposed in Ref. [8] and using the value of the computed deformation potential $D_{b,k_{||}} = 0.91$ eV/Å for $k_{||} = 0.12$ Å⁻¹. The simulated lattice dynamics attains a maximum displacement value $u = 1.6$ pm, corresponding to 0.14% of the body diagonal length $c = 11.86$ Å.

As shown in Fig. 3(d), the measured amplitude of oscillation is 20 meV for $k_{||} < 0.05$ Å⁻¹ and decreases down to 10 meV for $k_{||} < 0.2$ Å⁻¹. At larger wave vectors the photoelectron intensity of band $b$ is too low for a reliable extraction of the $D_{b,k_{||}}u$ value. The large $k_{||}$ dependence of $D_{b,k_{||}}$ has been previously reported for the amplitude mode.
energy has also been observed in the surface states of
mental conditions, the photoinduced increase of binding
follows the electronic temperature. However, the depopu-
states. As shown in Fig. 4, the dynamics of peak
shifts peak
Besides coherent phonon oscillations, an additional effect
states does not depend simply on the lattice coordinates. This aspect of the coupling can be
understood by a simple tight-binding model on a linear
chain: if the lattice displacement modifies the overlap
between atomic orbitals, the binding energy of the Bloch
states will change differently for different wave vectors.
Such $k$ dependence of the electron-phonon coupling is
essential for several problems of solid state physics: it
accounts for the presence of charge-density waves that
do not respect the nesting condition [16] and it allows for
the correct estimate of the high transition temperature in
superconducting MgB$_2$ [17].

In the case of bismuth, our data offer a unique oppor-
tunity to test the deformation potential extracted by DFT.
We calculate the band structure of bismuth for several
values of the displacement $u$. Figure 3(e) shows the dis-
persion of band $b$ for $u$ between 0.1% and 0.3% of the body
diagonal length $c = 11.86$ Å. In agreement with the mea-
surements, the binding energy of band $b$ strongly varies
near the $\Gamma$ point whereas it is less sensitive to the phonon
coordinate for $k|| = 0.2$ Å$^{-1}$. On the other hand, Fig. 3(d)
shows that the wave-vector dependence of $D_{bk_b}$ is stronger
in the calculations than in the experiment. A more accurate
simulation of the measured signal would require the com-
putation of one-step photoemission spectra [10].

It is clear in Fig. 3(c) that the dynamics of electronic
states does not depend simply on the lattice coordinates.
Besides coherent phonon oscillations, an additional effect
shifts peak $b$ towards negative values. This photoinduced
shift occurs also in the surface resonance and surface
states. As shown in Fig. 4, the dynamics of peak $a$ nearly
follows the electronic temperature. However, the depopu-
luation of the electronic states does not account for it (see
also Supplemental Material [14]). Under similar experi-
mental conditions, the photoinduced increase of binding
energy has also been observed in the surface states of
gadolinium [18]. An opposite behavior has instead been
reported in the electronic states of materials with photo-
induced phase transitions [15,19].

The detailed analysis of the electronic shift as a function
of $k$ will be part of forthcoming work. Nonetheless, we
notice that the magnitude of this component depends on the
electronic wave vector and band index. A wave-vector
dependent stiffening of the electronic states in the bonding
band can be expected because of the reduced ion core
screening at elevated electronic temperature. However,
the finite temperature DFT calculations indicate that the
charge transfer of Fig. 3(b) accounts at maximum for
2 meV to the shift observed in band $b$. This discrepancy
may be due to the fact that our bulk model of the electronic
states does not take into account charge redistribution and
carrier transport at the surface of the sample. Alternatively,
the dynamical aspect of the electronic screening may
require simulations that go behind the DFT method [20].

In conclusion, the electronic states of photoexcited
bismuth oscillate with the frequency of the $A_{1g}$ mode. By
analyzing the photoelectron spectra collected at different
angles, we reconstruct the deformation potential of specific
Bloch states. A quantitative agreement with DFT calcula-
tions confirms that the photoinduced charge transfer ac-
counts for the amplitude of the observed oscillations.
Finally, we mention that a purely electronic effect adds
to the temporal evolution of the electronic states with the
dynamics of the electronic temperature.

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