Fioretti, Angela N.; Schwartz, Craig P.; Vinson, John; Nordlund, Dennis; Prendergast, David; Tamboli, Adele C.; Caskey, Christopher M.; Tuomisto, Filip; Linez, Florence; Christensen, Steven T.; Toberer, Eric S.; Lany, Stephan; Zakutayev, Andriy

Understanding and control of bipolar self-doping in copper nitride

Published in:
Journal of Applied Physics

DOI:
10.1063/1.4948244

Published: 14/05/2016

Please cite the original version:
Understanding and control of bipolar self-doping in copper nitride

Angela N. Fioretti, Craig P. Schwartz, John Vinson, Dennis Nordlund, David Prendergast, Adele C. Tamboli, Christopher M. Caskey, Filip Tuomisto, Florence Linez, Steven T. Christensen, Eric S. Toberer, Stephan Lany, and Andriy Zakutayev

Citation: Journal of Applied Physics 119, 181508 (2016); doi: 10.1063/1.4948244
View online: http://dx.doi.org/10.1063/1.4948244
View Table of Contents: http://aip.scitation.org/toc/jap/119/18
Published by the American Institute of Physics

Articles you may be interested in
Controlled bipolar doping in Cu$_3$N (100) thin films

Synthesis of a mixed-valent tin nitride and considerations of its possible crystal structures
The Journal of Chemical Physics 144, 144201 (2016); 10.1063/1.4945561

Electrical properties and thermal stability of Pd-doped copper nitride films

Fulfilling the promise of the materials genome initiative with high-throughput experimental methodologies
Applied Physics Reviews 4, 011105 (2017); 10.1063/1.4977487
Understanding and control of bipolar self-doping in copper nitride

Angela N. Fioretti,1,2,a) Craig P. Schwartz,3 John Vinson,4 Dennis Nordlund,3 David Prendergast,5 Adele C. Tamboli,1,2 Christopher M. Caskey,1,2 Filip Tuomisto,6 Florence Linez,6 Steven T. Christensen,5 Eric S. Toberer,1,2 Stephan Lany,1 and Andriy Zakutayev1,1,a)
1National Renewable Energy Laboratory, Golden, Colorado 80401, USA
2Colorado School of Mines, Golden, Colorado 80401, USA
3SLAC National Accelerator Lab, Menlo Park, California 94720, USA
4National Institute of Standards and Technology, Gaithersburg, Maryland 20899, USA
5Lawrence Berkeley National Laboratory, Berkeley, California 94720, USA
6Aalto University, Espoo 02150, Finland

(Received 30 October 2015; accepted 26 February 2016; published online 5 May 2016)

Semiconductor materials that can be doped both n-type and p-type are desirable for diode-based applications and transistor technology. Copper nitride (Cu3N) is a metastable semiconductor with a solar-relevant bandgap that has been reported to exhibit bipolar doping behavior. However, deeper understanding and better control of the mechanism behind this behavior in Cu3N is currently lacking in the literature. In this work, we use combinatorial growth with a temperature gradient to demonstrate both conduction types of phase-pure, sputter-deposited Cu3N thin films. Room temperature Hall effect and Seebeck effect measurements show n-type Cu3N with 10^17 electrons/cm^3 for low growth temperature (~35 °C) and p-type with 10^15 holes/cm^3–10^16 holes/cm^3 for elevated growth temperatures (50 °C–120 °C). Mobility for both types of Cu3N was ~0.1 cm^2/Vs–1 cm^2/Vs. Additionally, temperature-dependent Hall effect measurements indicate that ionized defects are an important scattering mechanism in p-type films. By combining X-ray absorption spectroscopy and first-principles defect theory, we determined that VCu defects form preferentially in p-type Cu3N, while Cu_i defects form preferentially in n-type Cu3N, suggesting that Cu3N is a compensated semiconductor with conductivity type resulting from a balance between donor and acceptor defects. Based on these theoretical and experimental results, we propose a kinetic defect formation mechanism for bipolar doping in Cu3N that is also supported by positron annihilation experiments. Overall, the results of this work highlight the importance of kinetic processes in the defect physics of metastable materials and provide a framework that can be applied when considering the properties of such materials in general.

Published by AIP Publishing. [http://dx.doi.org/10.1063/1.4948244]

I. INTRODUCTION

The ability to dope a semiconductor both n-type and p-type is a critical factor in determining the viability of that material for optoelectronics applications (solar cells, light-emitting diodes, photodetectors) and transistor technologies. In cases where bipolar doping is not achievable, selecting a suitable heterojunction partner for diode fabrication can generate a host of additional materials challenges that often hinder device development. This problem has been particularly acute in the n-type nitride-based semiconductors, such as the III-N class of materials. For example, true p-n junctions based on GaN took decades to develop, ultimately leading to a Nobel Prize.1,2 Effective p-type doping is also problematic in solar-absorbing nitride alloys such as (In,Ga)N1,3 and has yet to be shown in other novel nitride semiconductors with solar-matched bandgaps, like ZnSnN2. Given the difficulty in obtaining bipolar doping in many cases, particularly in nitride semiconductors, any material that exhibits this quality warrants pursuit of a deeper understanding of the phenomenon.

In contrast to many other nitride materials, bipolar doping has been experimentally demonstrated4 and externally verified5 in the metastable semiconductor copper nitride (Cu3N). This behavior has been observed as a function of either growth temperature4 or Cu/N flux ratio,5 suggesting some kind of self-doping mechanism. The ability to achieve both conduction types in Cu3N is particularly intriguing due to its direct bandgap of 1.4 eV and indirect bandgap of 1.0 eV, which are well suited for photovoltaic (PV) applications.4,6 Growth temperature would be expected to affect the defect balance (and thus doping type) in Cu3N, since nitrogen chemical potential is a function of both nitrogen pressure and substrate temperature under the ideal gas approximation.6 However, changing Cu3N growth temperature may not only control its doping but can also lead to its decomposition or change the surface kinetics during growth, since Cu3N is a thermodynamically metastable material.6–8 Thus, a sound mechanistic understanding of how bipolar doping occurs in this material is necessary in order to manipulate this property with skill, and such understanding is currently lacking in the literature.

Prior work on Cu3N has focused on characterizing properties such as decomposition temperature,7–9 reflectivity,10–12...
and conductivity as a function of changing deposition parameters. Cu3N was found to exhibit either n-type conduction or quasi-metallic behavior, with carrier concentration or conductivity being directly proportional to substrate temperature during growth. Theoretical calculations later showed that Cu3N has a band structure with anti-bonding states at the valence band maximum (VBM), opposite to that of conventional semiconductors with bonding states at the VBM. Such an unusual band structure causes defects to form as shallow states near the band edges rather than in the mid-gap, leading to a property known as defect tolerance. This finding suggests that intentionally introducing native point defects into Cu3N during growth could be a possible technique for tuning its doping without deteriorating its performance.

In this work, we demonstrate n-type and p-type conduction in Cu3N as a function of substrate temperature during growth and propose a mechanism that explains both the origin and the temperature-dependence of this bipolar doping behavior. The findings presented herein create a new framework for understanding bipolar doping in Cu3N, which is strongly dependent on a balance between surface and bulk kinetic processes. This understanding is achieved using a combinatorial synthesis method paired with high throughput characterization, in addition to advanced characterization techniques and both bulk and surface defect calculations. Finally, the kinetically driven, point-defect-based mechanism developed herein for bipolar doping in Cu3N highlights the important influence of kinetic processes on the defect physics of metastable materials in general.

II. METHODS

A. Thin film deposition

Thin films of Cu3N were deposited via radio frequency (RF) sputtering on 50 mm × 50 mm Eagle XG glass substrates using a combinatorial deposition chamber with 1 × 10^-4 Pa (1 × 10^-6 Torr) residual water base pressure and an RF-plasma atomic nitrogen source. Two sputter guns with 50 mm elemental Cu targets were inclined at 45° with respect to the substrate, and the atomic nitrogen source was positioned at normal incidence angle, to achieve a nominally constant film thickness and to avoid a gradient in nitrogen activity (left side, Fig. 1). Target-substrate distance was kept at 13 cm for all samples. For a more detailed description of the atomic nitrogen source and of the sputter deposition procedures used in this work (see Ref. 6). Target power density for each gun was maintained at 1.5 W cm^-2, and the atomic nitrogen source was powered at 250 W. Nitrogen gas was flowed in through the nitrogen RF source at a rate of 10 cm^3/min and balanced by 10 cm^3/min argon gas for a total chamber pressure of 2.6 Pa (20 mTorr). One side of the glass substrate was kept in contact with a resistive heating unit maintained at constant temperature, so that a continuous temperature gradient was induced across the substrate during growth. For a more detailed description of how the combinatorial temperature gradient was induced in this work see Ref. 16. Hot-side set point temperatures explored in this study were 120°C, 90°C, and no active heating. These set point temperatures represent a continuous growth temperature spread from 120 to 50°C in addition to a growth temperature of 35°C, as calibrated by a thermocouple on the substrate surface.

B. Characterization

Each deposition performed as described above yielded a library of samples grown at a range of temperatures dictated by the hot-side temperature. Long-range phase purity was determined using X-ray diffraction (XRD). Conductivity of each library was measured using four-point probe sheet resistance measurements combined with the thickness measured by profilometry. XRD, four point probe, and profilometry measurements were taken using a combinatorial 44-point grid (right side, Fig. 1). Film microstructure was imaged at representative regions via scanning electron microscopy (SEM). Carrier concentration and mobility were measured on selected regions of each library using room temperature Hall effect in the van der Pauw configuration and room temperature Seebeck effect. Regions selected for Hall and Seebeck measurements are indicated in Fig. 1 with white asterisks and were removed from the original libraries in 7 mm × 7 mm pieces. Temperature-dependent Hall effect measurements were performed on a region of another p-type library grown isothermally at ≈160°C. Error bars shown for the conductivity data give the standard deviations among the four repeated rows of each combinatorial sample. Error bars shown for the Hall effect data give the standard deviations among 5–10 repeated measurements of each of the selected pieces measured from each library. Further information on the experimental methods used herein can be found in prior publications.

Experimental X-ray absorption spectra (XAS) were taken on representative p-type and n-type Cu3N samples at beamline 8–2 at the Stanford Synchrotron Radiation Lightsource (SSRL). A nominal resolution of ∆E/E of 10^-4 is provided by this beamline. Data were collected both by using largely bulk-sensitive total fluorescence yield (TFY), which has a penetration of ≈100 nm, and by the more surface sensitive total electron yield (TEY), which has a penetration of ≈10 nm. Additionally, both n- and p-type Cu3N samples were analyzed using positron annihilation spectroscopy (PAS). The positron irradiation energy was tuned from 0.5 keV to 25 keV to probe the samples from the surface up to a mean implantation depth.

FIG. 1. Diagram depicting the combinatorial chamber geometry used to obtain a gradient only in temperature (left) and the layout of the 44-point combinatorial grid used for all mapping characterization described in this work (right). The asterisks denote regions of the film that were removed and subjected to non-mapping-style measurements.
of 1.3 μm (i.e., into the substrate). After implantation and thermalization in the layer, each positron diffuses and annihilates with an electron. Each such event results in two photons with an energy of 511 ±ΔEkeV, which are measured by a high-purity Ge detector with an energy resolution of 1.3 keV full width at half maximum. The Doppler broadening, ΔE, depends on the momentum distribution of the annihilating electrons. Typically, the Doppler broadened energy spectrum narrows when positrons are trapped by vacancy defects. The shape of the spectrum is characterized using S and W shape parameters referring to the fraction of positron annihilation with low-momentum electrons (pL < 4 a.u.) and high-momentum electrons (1.6 a.u. < pL < 4 a.u.), respectively.21

C. Computation

The defect formation energies ΔHD for the copper vacancy (VCu), copper interstitial (CuI), the N vacancy (VN), and substitutional oxygen (ON) were determined using the results of previously published supercell calculations22 using the GGA+U functional with U = 5 eV for Cu-d. In order to determine the ΔHD for the actual growth conditions, we assumed here a quasi-equilibrium of Cu3N with Cu-metal (∆μCu = 0) and an activated source of N (∆μN = +0.76 eV). The value for ∆μN was obtained from the stability condition (3 × ∆μCu) + ∆μN = ΔHd(Cu)N = +0.76 eV. For the chemical potential of oxygen, we assumed ∆μO = −0.84 eV, which corresponds to the ideal gas law chemical potential for T = 200°C (upper T limit for Cu3N growth) and pO2 = 10−3 Pa (base pressure of the growth chamber). Thermodynamically, the formation of CuO could further reduce the O chemical potential to ∆μO = −1.71 eV. However, this requires that the kinetics allows for nucleation and precipitation of a CuO phase. While we observe a CuO-like surface due to oxidation in ambient conditions, we do not have evidence for CuO formation during deposition. Thus, we consider ∆μO = −0.84 eV as an upper bound. In order to calculate the dependence of the VCu and CuI defect formation energy as a function of the distance from the surface, we used a slab supercell with 248 atoms modeling the partially oxidized (001) surface of Cu3N. This surface termination, formed by substituting every other N atom of the (001) surface by an O atom, was previously found to yield an exceedingly low surface energy.4 The charge compensation was maintained by changing the O/N ratio at the surface upon introducing the defects. These supercell calculations were performed using the VASP code,23 with otherwise identical computational settings as in Ref. 24.

The modeled XAS structures were generated as detailed previously, with the exception of the CuO surface structure.4 The previously calculated bulk structures consist of approximately 100 atoms with specific defects inserted into them. The surface structures consisted of at least 50 atoms. The CuO structure was generated by placing CuO on top of a slab of Cu3N and relaxing the structure within VASP.24 The X-ray calculations were performed in two ways, both largely detailed previously.25,26 First, X-ray core hole (XCH) spectra were calculated only at the nitrogen K-edge. Within an ultra-soft pseudopotential formalism, the excitation is simulated by replacing the excited atom with a 1s core hole, such that the excited nitrogen now is represented as 1s2s2p6. K-point sampling is used to converge the spectra. The unoccupied p-type orbitals are then projected onto the excited atom and the resultant stick intensity as a function of energy is broadened by 0.05 eV and the process is repeated for every atom. Second, calculations based on solving the Bethe-Salpeter equation (BSE) were performed within oceAn software package for Cu L-edge as previously detailed, using Density Functional Theory (DFT) as a basis.27 The electronic structure was calculated within the local-density approximation using the Quantum ESPRESSO code,28 and sampling of the Brillouin zone was improved through the use of the orthogonal basis function interpolation scheme. The number of bands included hundreds (>1000) of the first conduction bands, and the screening of the core-hole potential included thousands (>4000) bands. The K-space and mesh sampling were checked for convergence. The broadening was 0.2 eV. The spectra were aligned based on the Fermi level and potential on the core.
III. RESULTS

A. Structure

Crystalline Cu3N films with long-range phase-purity were achieved at all growth temperatures explored in this work. Long-range phase-purity can be seen in Fig. 2(a), which displays XRD patterns for Cu3N grown at various temperatures. Based on comparison to the peak intensities of the Cu3N powder diffraction pattern from the Inorganic Crystal Structure Database (ICSD), these films exhibit no preferential orientation. The absence of peak intensity at 50.5° and at 43.3° confirms the absence of any long-range metallic copper second phase, indicating that the films in this work represent a comparison between phase-pure Cu3N films grown at different conditions and not between Cu3N and a phase-separated mixture of Cu3N and Cu0.

As shown in Figs. 2(b) and 2(c), two distinct film morphologies were observed for the thin films prepared in this study. Films grown at a slightly elevated substrate temperature (i.e., 50°C–120°C) exhibited spheroid grain termination with the grain size on the order of 10 nm–50 nm (Fig. 2(b)). For films grown without actively heating the substrate (i.e., the only source of substrate heating was from the impinging sputtered atom flux), pyramidal grain termination with 50 nm–100 nm grains was observed (Fig. 2(c)). Observing larger grains for unheated growth and smaller grains for slightly elevated growth temperature is related to the metastable nature of Cu3N. At elevated growth temperature, slightly below the decomposition temperature of Cu3N, it is possible that partial evaporation leads to spherical grain termination such that the surface area is minimized. A more detailed description of this counterintuitive grain size phenomenon and its relation to metastability can be found in previous works.6,15

B. Transport

Fig. 3 shows the electrical transport properties of Cu3N films as a function of growth temperature. Designations of n-type and p-type in all panels of Fig. 3 are based on the sign of the Hall coefficients and corresponding Seebeck coefficients, summarized in Table I. Films from the 120°C region were unable to be reliably measured by Hall effect, due to contact degradation with repeated measurement. The same effect was not observed while collecting Seebeck effect data, which allowed the high-temperature samples to be designated p-type. To fill the gap in the Hall effect data, a sample grown isothermally at 160°C in a previous work4 was used for obtaining temperature-dependent Hall effect data. The data from that sample are indicated with an asterisk in Table I.

Hall and Seebeck effect measurements of the three Cu3N libraries reveal a switch from n-type to p-type conduction as the growth temperature increased from 35°C to 160°C. This switch in majority carrier type coincides with a U-shaped conductivity trend measured as a function of increasing growth temperature in the 35°C–120°C range (Fig. 3(a)). This correlation suggests competing formation of donor and acceptor defects, with donor defects dominating during unheated growth and acceptor defects dominating at elevated temperatures. Such a mechanism would best be described as defect compensation, in which the observed majority carrier type is determined by whether donor or acceptor defects are present in higher concentration at a given growth temperature. Support for this defect compensation hypothesis was obtained by investigating the transport properties of films in this work using carrier density and mobility data from Hall effect.
Compensated semiconductors would be expected to display transport properties consistent with ionized defect scattering. Evidence for ionized defect scattering was observed via temperature-dependent Hall effect for p-type samples. This can be seen in Fig. 3(b), in which the carrier density and mobility were inversely proportional to each other as a function of decreasing temperature for p-type Cu$_3$N grown at 160 °C. Fig. 3(b) also shows that the carrier density has an activated character as a function of inverse temperature, which indicates carrier freeze out at low temperatures. For p-type samples grown between 85 °C and 50 °C (Fig. 3(c)), it was found that mobility exhibited an exponential decrease as a function of increasing carrier density; again, this is characteristic of ionized defect scattering and is consistent with a competing defect formation hypothesis. Collectively, these findings support the conclusion that competing defect formation as a function of growth temperature leads to compensation, which results in the observed bipolar self-doping.

An Arrhenius analysis of the carrier density–temperature data $n(T)$, shown in Fig. 3(b), yields a pre-factor $n_0 = 7.5 \times 10^{19}$ cm$^{-3}$ and an activation energy $E_{\text{act}} = 0.188$ eV (dashed black line). However, these numbers should not directly be associated with the acceptor density and ionization energy. Rather, the $n(T)$ function can be modeled by a detailed balance approach, such as that used for calculation of defect equilibria, taking into account ionization probabilities, the effective masses at the band edges, as well as the mutual compensation of acceptors and donors. In particular, there is an inherent contribution to the activation energy due to the temperature dependence of the so-called effective density of states. In fact, for the case of an uncompensated acceptor, we need to assume an acceptor ionization energy as large as 0.33 eV in order to reconcile the Hall data. However, this ionization energy is much larger than $E_a = 0.12$ eV obtained by the first principles calculations (see Fig. 4), and according to the discussion above, a significant compensation is very likely.

We show in Fig. 3(b) two sets of detailed balance simulations, which can be viewed as limits for the range of physically meaningful solutions that are consistent with the Hall data. For these simulations, we use a band gap of 1.0 eV and an effective hole mass of $m_h/\hbar^2 = 2.8$, as obtained from GW calculations [http://materials.nrel.gov]. The first set (red curve) is obtained for a moderate total acceptor concentration of $N_A = 4 \times 10^{17}$ cm$^{-3}$, a compensation ratio of $N_D/N_A = 0.12$, and an acceptor ionization $E_a = 0.20$ eV. The second set (solid black line) is obtained for a very high total acceptor concentration of $N_A = 10^{22}$ cm$^{-3}$, a high compensation ratio $N_D/N_A = 0.86$, and $E_a = 0.15$ eV. The small but significant differences in the functional form of these two datasets (Fig. 3(b)) suggest that such simulations could be utilized in the future in conjunction with more detailed Hall measurements for characterization of electrical properties in terms of a defect model. The results of the simulations also suggest that the calculated acceptor level $E_a = 0.12$ eV is somewhat underestimated, which is very common in DFT calculations, unless specific self-interaction corrections are employed.

Demonstration in this work of bipolar self-doping in Cu$_3$N confirms the reproducibility of this phenomenon as reported elsewhere but does not offer sufficient insight into the specific origins of this property nor into the identity of the defects giving rise to each majority carrier type. Arguably, a deeper understanding of bipolar doping in copper nitride will be necessary if this desirable property is to be manipulated with precision. Recognizing this necessity, we used different computational methods of defect identification coupled with X-ray absorption spectroscopy and positron annihilation spectroscopy to investigate the atomic-scale origins of bipolar self-doping in Cu$_3$N.
C. Bulk defect formation enthalpy

Fig. 4 shows the defect formation enthalpies as a function of the Fermi level, assuming a quasi-equilibrium between Cu$_3$N, Cu-metal, and activated N. Only defects that are stable and could contribute to n-type or p-type doping are shown. It is apparent from Fig. 4 that only VCu defects could contribute to p-type doping in this material, as this is the only defect with a positive slope as the Fermi level moves down in energy from the conduction band minimum (CBM) to the valence band maximum (VBM). This is consistent with other Cu$^{1+}$-based semiconductors, such as Cu(In,Ge)Se$_2$ and Cu$_2$O, that exhibit p-type conductivity. For n-type doping, three stable donor defects were found: VN, O$_N$, and Cu$_i$. Oxygen substitutional defects were included in these calculations, even though O$_N$ is not a native defect because oxygen incorporation from background pressure is common in nitride films. The formation enthalpy of VN was found to be 1.5 eV higher than O$_N$ and Cu$_i$ for the entire range of Fermi energies plotted, making it the least favorable donor defect to form and therefore an unlikely contributor to n-type doping in copper nitride. Considering defect formation enthalpies alone, the results presented in Fig. 4 suggest that VCu defects give rise to p-type doping and O$_N$ plus Cu$_i$ defects give rise to n-type doping.

The formation enthalpies shown in Fig. 4 suggest that even when the incorporation of O is suppressed by reducing or eliminating O from the growth process (effectively increasing the $\Delta H_D$ of O$_N$), a true quasi-equilibrium situation would lead to n-type doping due to an excess of Cu$_i$ over VCu. However, such a situation might be difficult to achieve, because it requires a sufficiently high temperature to enable bulk diffusion, which at the same time also facilitates the decomposition of the metastable Cu$_3$N compound. Therefore, we employed additional computational and experimental methods to gather more information about the atomic-scale origins of bipolar doping.

D. X-ray absorption spectroscopy

Using near edge X-ray absorption fine structure (NEXAFS) measurements and modeling performed using OCEAN software package,$^{25,33}$ it was found that O$_N$ defects are unlikely to be dominant contributors to bipolar doping in Cu$_3$N. This finding is in addition to the finding in Section III C that VN defects are the most energetically unfavorable donor defects to form in this material. Taken together, these results suggest that Cu$_i$ and VCu defects make the most significant contributions to Cu$_3$N bipolar doping behavior. NEXAFS spectra for the Cu L-edge and the N K-edge for representative n-type and p-type samples are shown in Fig. 5, along with modeled spectra corresponding to Cu$_i$, VCu, and O$_N$ defect structures. Both Cu$_i$ and VCu defects were found to be present in both p-type and n-type Cu$_3$N, which supports the hypothesis that competing donor and acceptor defect formation is the mechanism behind bipolar self-doping in this material.

To arrive at the above conclusions, it was necessary to use a formalism for modeling NEXAFS spectra that was not
Three defect structures for Cu$_3$N were simulated using the BSE-based method described above: Cui, V$_{Cu}$, and ON. These defect structures were chosen based on the point defect formation enthalpies presented in Section III C. The resulting modeled spectra for each edge are displayed in Figs. 5(c) and 5(d). A vertical gray line is overlaid on each set of panels to indicate the experimental peak position of the first leading edge peak that is not due to the Cu$^{II+}$ surface layer. Looking specifically at the Cu L-edge (Figs. 5(a) and 5(c)), the only defect structure that is unlikely to be a major contributor to the NEXAFS signal is O$_N$ substitution (red curve), due to the modeled spectrum being significantly red-shifted compared to experiment. This is not to say that O$_N$ defects are not present in the films in this work only that O$_N$ substitutions in the bulk must exist at a far lower concentration than either Cui or V$_{Cu}$. Turning to the N K-edge (Figs. 5(b) and 5(d)), it is clear that none of the other defect structures are shifted enough to be reasonably ruled out as contributing to the overall spectra. It should be noted that the resemblance of the simulated defect spectra in panel (d) to the CuO surface layer spectrum in panel (b) is due to the s-p transition of the nitrogen K-edge being highly sensitive to broken local symmetry. This explains why the simulated defect spectra shown in panel (c) more closely resemble the simulated perfect crystal spectra shown in panel (a), rather than resembling the simulated CuO surface layer: because the p-d transitions probed at the Cu L-edge are not as sensitive to broken local symmetry, and therefore the defect spectra are not dominated by these effects as much as the N k-edge spectra. The conclusion that O$_N$ defects are not major contributors to the NEXAFS spectra is consistent with the films in this work being grown under activated nitrogen atmosphere, in which substitution of oxygen impurities on nitrogen sites is mitigated by the dramatic improvement in nitrogen incorporation. Overall, the combination of calculated defect formation enthalpies and modeled NEXAFS spectra points to Cui and V$_{Cu}$ defects as being the most prominent point defects in both p-type and n-type Cu$_3$N.

Conceptually, if both Cui and V$_{Cu}$ defects are present in both conduction types of Cu$_3$N, then it follows that the donor defect (Cui) must dominate in n-type copper nitride and the acceptor defect must dominate in p-type copper nitride. For insulating Cu$_3$N, both defect types must be present in equal numbers and lead to full defect compensation. What is not clear from this analysis is why the donor defect would form preferentially at low growth temperature (~35°C) and the acceptor defect (V$_{Cu}$) would do so at elevated growth temperature (120°C). To answer this question and thus propose a complete mechanism for bipolar doping in Cu$_3$N, an additional set of first principles calculations were performed. Specifically, depth-dependent defect formation enthalpies were calculated to determine whether each defect type (Cui and V$_{Cu}$) could form preferentially at the surface. Based on these surface calculation results, a kinetic mechanism was proposed to explain the observed temperature dependence of the doping type in Cu$_3$N.
The surface. This finding implies that the prevalence of Cu
slightly lowered and even increased for a Cu adatom on top of
observed p-type character of Cu$_3$N at the elevated deposition
defect formation enthalpies for Cu$_3$N were recalculated as a
function of depth from the film surface. Only formation enthal-
pies for Cui and VCu defects were recalculated because both
VN defects (Fig. 4) and ON defects (Fig. 5) were already identi-
fied as not being major contributors to the bipolar self-doping
phenomenon. The results of these calculations, shown in Fig.
6, indicate that VCu formation energy is sharply lowered
towards the surface, whereas the Cu$_i$ formation energy is only
slightly lowered and even increased for a Cu adatom on top of
the surface. This finding implies that the prevalence of Cu$_i$
over VCu in the bulk (Fig. 4) could be inverted close to the
growing surface, creating a net p-type doping. It would then
lead to enhanced formation of Cui donors over VCu acceptors
at any given time is given by
$$c_e = \frac{1}{c_0 e^{-\frac{E_a}{k_B T}}}$$
where $K = \frac{K_0 e^{-E_a/k_B T}}{c_0}$ is the T-dependent desorption rate determined by
activation energy $E_a$ (proportional to residence time of the
atoms at the surface). This inherent Cu vs. N asymmetry
leads to enhanced formation of Cu$_i$ donors over VCu acceptors
and thus produces n-type conductivity at these lower sub-
strate temperatures.

Next, we address n-type conductivity at lower substrate
temperature (≈35 °C, Fig. 3(a)). It is likely that in this tem-
perature regime, equilibration of the surface defects as dis-
cussed above does not fully occur. In this case, a non-
equilibrium defect distribution becomes frozen in the bulk
when the next monolayer of Cu and N atoms arrives. Note
that Cu atoms are more likely to be frozen in the bulk than N
atoms because the Cu-Cu diatomic species probably has a
longer residence time at the surface than the gaseous N-N
species (i.e., N$_2$ molecule). From a kinetic point of view, the
atom concentration $c$ at the surface of the growing Cu$_3$N film
at any given time is given by $c = c_0 e^{-\frac{E_a}{k_B T}}$, where $K = $
Based identification of the vacancy-type defects in the films by sluggish kinetics at this lower deposition temperature. Supporting the hypothesis that the n-type doping is enabled at the growth temperature than the rest of the n-type film, interfacial p-type layer can be attributed to longer time spent at the interface with the substrate. The presence of this nominally n-type sample resembles that of the p-type film/substrate interface. Hence, the defect structure of the bulk diffusion are kinetically limited (no active heating), n-type Cu₃N is observed. See manuscript text for further explanation.

It should be noted that detailed positron annihilation-based identification of the vacancy-type defects in the films in this work was not possible due to the lack of a reference sample with low defect density. For an additional representation of the PAS data, showing the variation of W with S for n-type and p-type samples, please see Fig. S2 of the supplementary material.

Our current understanding of bipolar self-doping in Cu₃N is summarized in Fig. 8. At lower substrate temperatures, Cu₃N shows n-type conductivity due to Cuᵩ defects frozen in the bulk that dominate over VCu, defects as a result of surface diffusion barriers. As the growth temperature increases, such that the thermal energy becomes comparable to the surface diffusion barriers, the conductivity switches from n-type to p-type due to the decreased formation energy of VCu, and increased formation energy of Cuᵩ, at the growing Cu₃N surface (Fig. 6). In both of these temperature regimes, both VCu and Cuᵩ are present in Cu₃N, highlighting the compensated character of this semiconductor. Finally, at the highest substrate temperature, Cu₃N decomposes into metallic Cu and N₂, which corresponds to bulk equilibration of this metastable semiconducting material. Note that this mechanistic understanding is consistent with a number of experimental observations, including temperature-dependent conductivity type (Fig. 3), synchrotron-based NEXAFS measurements (Fig. 5), depth-dependent positron annihilation (Fig. 7), and Cu₃N growth/decomposition experiments (Ref. 7).

**IV. CONCLUSIONS**

We have presented an explanation for bipolar self-doping in the metastable material Cu₃N. Using a combination of high-throughput synthesis, advanced characterization, and different computational approaches, we were able to address the outstanding question of why bipolar self-doping behavior is observed in this material. The proposed mechanism implicates two native point defects, specifically the donor defect Cuᵩ and the acceptor defect VCu, as giving rise to n-type and p-type Cu₃N, respectively. Doping type was found to be temperature dependent, with p-type doping observed for samples grown at the elevated temperature (50°C–120°C) and n-type doping observed for samples grown with no active heating (≈35°C). On average, p-type films in this work exhibited carrier density on the order of 10¹⁵ holes/cm³–10¹⁶ holes/cm³ with μ = 0.1 cm² V⁻¹ s⁻¹–1.5 cm² V⁻¹ s⁻¹ and n-type films exhibited ≈10¹⁷ electrons/cm³ with μ = 1.5 cm² V⁻¹ s⁻¹. Additionally, both the donor and the acceptor defects (Cuᵩ and VCu) were found to be present in both conduction types of Cu₃N based on NEXAFS analysis, indicating that competing defect formation as a function of temperature determines the ultimate majority carrier type. Indeed, defect compensation was found to be necessary in order to model the temperature-dependent carrier density data using a detailed balance approach. This hypothesis was supported by comparing bulk and surface defect formation energies, in which it was shown that the acceptor defect formation is preferentially enhanced and the donor defect formation is preferentially suppressed at elevated growth temperatures just below the decomposition temperature of Cu₃N. The kinetics-driven doping mechanism presented herein explains this competition as a balance...
between surface and bulk kinetic processes. When surface diffusion is dominant and bulk diffusion is frozen (50°C–120°C), p-type Cu₃N is observed. When both surface and bulk diffusion are kinetically limited (no active heating), n-type Cu₃N is observed. This mechanistic understanding of copper nitride bipolar doping is consistent with several experimental observations, both from this work and from previous works. Overall, the results of this work highlight the importance of kinetic processes in the defect physics of metastable materials and provide a framework that can be applied when considering the properties of such materials in general.

ACKNOWLEDGMENTS

This work was supported by the U.S. Department of Energy under Contract No. DE-AC36-08GO28308 with the National Renewable Energy Laboratory (NREL). A.N.F. was supported by the Renewable Energy Materials Research Science and Engineering Center under Contract No. DMR-0820518 at the Colorado School of Mines. Computational Science and Engineering Center under Contract No. DMR-0820518 at the Colorado School of Mines. Certain commercial equipment, instruments, or materials are identified in this paper in order to specify the experimental procedure adequately. Such identification is not intended to imply recommendation or endorsement by the National Institute of Standards and Technology, nor is it intended to imply that the materials or equipment identified are necessarily the best available for the purpose.


29See supplementary material at http://dx.doi.org/10.1063/1.4948244 for Figs. S1 and S2.


