Tunable Conductivity and Conduction Mechanism in an Ultraviolet Light Activated Electronic Conductor

Mariana I. Bertoni
Thomas O. Mason
Julia E. Medvedeva
Missouri University of Science and Technology, juliaem@mst.edu
Arthur J. Freeman

et. al. For a complete list of authors, see https://scholarsmine.mst.edu/phys_facwork/414

Follow this and additional works at: https://scholarsmine.mst.edu/phys_facwork

Part of the Physics Commons

Recommended Citation
M. I. Bertoni et al., "Tunable Conductivity and Conduction Mechanism in an Ultraviolet Light Activated Electronic Conductor," Journal of Applied Physics, American Institute of Physics (AIP), Jan 2005. The definitive version is available at https://doi.org/10.1063/1.1899246

This Article - Journal is brought to you for free and open access by Scholars' Mine. It has been accepted for inclusion in Physics Faculty Research & Creative Works by an authorized administrator of Scholars' Mine. This work is protected by U. S. Copyright Law. Unauthorized use including reproduction for redistribution requires the permission of the copyright holder. For more information, please contact scholarsmine@mst.edu.
Tunable conductivity and conduction mechanism in an ultraviolet light activated electronic conductor

Mariana I. Bertoni and Thomas O. Mason
Department of Materials Science and Engineering, Northwestern University, Evanston, Illinois 60208

Julia E. Medvedeva and Arthur J. Freeman
Department of Physics and Astronomy, Northwestern University, Evanston, Illinois 60208

Kenneth R. Poeppelmeier
Department of Chemistry Northwestern University, Evanston, Illinois 60208

Bernard Delley
Paul Scherrer Institut, Villigen CH-5232, Switzerland

(Received 1 November 2004; accepted 8 March 2005; published online 4 May 2005)

A tunable conductivity has been achieved by controllable substitution of an ultraviolet light activated electronic conductor. The transparent conducting oxide system H-doped Ca$_{12-x}$Mg$_x$Al$_{14}$O$_{33}$ ($x=0, 0.1, 0.3, 0.5, 0.8, 1.0$) presents a conductivity that is strongly dependent on the substitution level and temperature. Four-point dc-conductivity decreases with $x$ from 0.26 S/cm ($x=0$) to 0.106 S/cm ($x=1$) at room temperature. At each composition the conductivity increases reversibly with temperature until a decomposition temperature is reached; above this value, the conductivity drops dramatically due to hydrogen recombination and loss. The observed conductivity behavior is consistent with the predictions of our first principles density functional calculations for the Mg-substituted system with $x=0$, 1, and 2. The Seebeck coefficient is essentially composition and temperature independent, the later suggesting the existence of an activated mobility associated with small polaron conduction. The optical gap measured remains constant near 2.6 eV while transparency increases with the substitution level, concomitant with a decrease in carrier content. © 2005 American Institute of Physics.

DOI: 10.1063/1.1899246

INTRODUCTION

Materials with low sheet resistance and optical transparency are sought for many optoelectronic applications. These two properties tend to be mutually exclusive in nature and are usually obtained by creating electron degeneracy in a wide band gap material. Various transparent oxides are rendered electrically conducting by controllably introducing nonstoichiometry and/or appropriate dopants, but this approach is not applicable for the oxides of the main-group metals. Alternative processes are being developed to render these oxide systems conducting, as part of the search for inexpensive and environmentally benign alternatives.

The reported system, 12CaO·7Al$_2$O$_3$ is a well-known insulating oxide widely used in high-alumina cements. The crystal lattice belongs to the cubic system and space group $I43d$ with a lattice parameter of 1.199 nm. It possesses a cage structure with two formula units (12 cages) per unit cell and its empirical formula may be written as [Ca$_{24}$Al$_{28}$O$_64$]$^{4+}$+2O$_2^-$, where the free oxygen ions provide the charge neutrality and are located inside the cages of the framework (Fig. 1). The system was discovered to incorporate hydrogen at elevated temperatures through the following chemical reaction:

$$\text{O}_2^- + \text{H}_2(\text{atmosphere}) \rightarrow \text{OH}^- + \text{H}^-.$$  (1)

After hydrogen incorporation the unit cell contains two cages occupied by OH$^-$, another two occupied by H$^-$; and the remaining eight cages of the unit cell are empty. Hydrogen annealing results in no apparent change in the optical and electrical properties of the material. However, upon ultraviolet (UV) irradiation two optical absorption bands are induced, giving rise to a persistent color change from white to green together with a considerable conductivity increase. Our previous density functional investigations revealed that

FIG. 1. One of 12 cages constituting the unit cell of Ca$_{12}$Al$_{14}$O$_{33}$. The charge neutrality is given by the O$^-$ located inside the cage.
the charge transport associated with the photoexcitation of an electron from hydrogen [which is located inside one of the cages, cf. Eq. (1)], occurs by electron variable-range hopping through states of the encaged OH− and H O5 (Ref. 5) and d states of their nearest Ca neighbors (i.e., only 8 out of 24 Ca atoms in the unit cell of mayenite were found to give significant contributions to the density of states near the Fermi level).4 The detailed knowledge of the transport mechanism thus obtained for H-doped UV-activated mayenite predicts the strong dependence of the light-induced conductivity on the atoms participating in the hopping as well as on their spatial arrangement and hence the possibility of varying the conductivity by proper doping. In particular, we expected that Mg substitution may lead to a decrease in the conductivity once Mg substitutes one of the Ca atoms involved in the hopping, since its 3d states will lie much higher in energy than those of Ca.

In this article the conduction mechanism of the Ca12−xMg9Al14O33 (x=0, 0.1, 0.3, 0.5, 0.8, 1.0) system was studied both experimentally and theoretically. We report the change in conductivity, thermopower, optical gap (estimated from the absorption edge measured in a diffuse reflectance experiment), and transparency for the bulk oxide Ca12−xMg9Al14O33. In addition, we present a small polaron conduction mechanism that is consistent with the electronic conduction in this system and discuss the conductivity behavior observed for Cu12−xMg9Al14O33 based on the results and predictions of our first-principles total energy calculations for the Mg-substituted system with x=0, 1, and 2.

EXPERIMENT

Polycrystalline samples of Mg-substituted 12CaO·7Al2O3 were produced by conventional high temperature solid-state reaction. Stoichiometric amounts of CaCO3, Al2O3, and MgO (>99.99% purity metals basis, Alfa Aesar) were mixed in an agate mortar to produce various samples of Ca12−xMg9Al14O33 (x=0, 0.1, 0.3, 0.5, 0.8, 1.0). Once homogeneous mixtures were obtained, 11.6 mm diam × 2–3 mm thick pellets were pressed at 180 MPa. The pressed pellets were heated in air up to 1200 °C in high-purity alumina crucibles and held for 24 h. Once cooled the pellets were reground, repressed, and fired again at 1200 °C for 24 h.

The phase purity of the samples was confirmed by powder x-ray diffraction using Cu Kα radiation (Rigaku, Danvers, MA). A nickel filter was used to remove the Cu Kβ contribution from the diffraction pattern. Powders were scanned between 10° and 80° in 2θ for routine phase analysis.

The subsequent hydrogen treatment was done inside an 18 in. length quartz tube sealed with a water-cooled metal end cap under a constant forming gas (4%-5% H2, N2 balance) flow. The quartz tube was placed inside a high temperature tube furnace where the samples were taken to 1300 °C for 2 h and then rapidly cooled to room temperature by rapid extraction of the quartz tube out of the furnace.

After the hydrogenation, the samples were exposed to UV light under a mercury short arc lamp (Model HAS-200 dc) of total radiation (275–650 nm) 52.3 W, for 40–70 min.

Conductivity measurements were taken using the Van der Pauw technique where a four-point spring-loaded probe touches the sample in four different points close to the edges. A resistance \( R_{AB,CD} \) is defined as the potential difference \( V_{AB} \) between the contacts A and B. Similarly a resistance \( R_{BC,DA} \) can be defined and the following relation holds for a specimen of arbitrary shape:

\[
\exp\left(-\pi \cdot \frac{d}{\rho} R_{AB,CD}\right) + \exp\left(-\pi \cdot \frac{d}{\rho} R_{BC,DA}\right) = 1, \tag{2}
\]

where \( \rho \) is the resistivity of the material and \( d \) is the thickness of the UV-activated slab. Corrections were made for layer thickness and sample diameter in every case. It should be mentioned that corrections for porosity were not performed due to the uncertainty in the properties of the irradiated layer. However, previous conductivity measurements on powder specimens using a powder solution composite technique showed a very good agreement with the conductivity values obtained by the four-point probe method.8

Room temperature thermopower data were collected on bar-shaped samples cut from the irradiated pellets. The bars of approximately 10 mm × 3 mm × 3 mm had UV-activated conductive layers on one lateral face and both ends. These bar-shaped samples were sandwiched between two gold foil contacts. The bars were painted on the contact faces with a silver colloidal suspension to improve the electrical and thermal contact between the small UV-activated layer and the two gold electrodes. One gold contact was in thermal equilibrium with a 23 W heating element and the other was in thermal equilibrium with a cylindrical steel slug that rested on an insulating ceramic brick. A type S (Pt-Pt/10%Rh) thermocouple bead was welded to both gold contacts. A thermal gradient was created by switching on the heating element and allowing it to reach 100 °C, at which point the heating element was switched off, letting the system thermally relax. The temperature difference (\( \Delta T \)) and the voltage difference (\( \Delta V \)) were measured at regular intervals (3 s) using a programmable scanner (Keithley 705, Cleveland, OH) and a digital multimeter (Keithley 195A, Cleveland, OH) connected through an IEEE port to a personal computer. Thermopower was calculated by fitting the temperature and voltage gradient data with a least-squares fit as the sample approached equilibrium using the concept presented by Hong et al.9

\[
Q = -\lim_{\Delta T \to 0} \frac{\Delta V}{\Delta T}. \tag{3}
\]

A correction for the contribution of the Pt thermocouple to the overall thermopower has to be made using the polynomial fit of Hwang.10

Since thin films of these materials were unavailable, optical data were obtained from diffuse reflectance measurements. The spectra for the specimens were collected on a Cary 500 UV visible-near infrared spectrophotometer (Varian Instruments Inc., Palo Alto, CA) using a diffuse reflectance accessory between 400 and 700 nm with a lead sulfide detector. This accessory has the ability to collect most
reflected radiation, remove any directional preferences, and present an integrated signal to the detector. The data were corrected with a spectrum obtained from a polytetrafluoroethylene reference specimen. The optical gap was estimated from the absorption edge that was determined by the intersection of a line drawn through the sloped portion of the transition region between high and low transmission and the baseline of the low-transmission portion of each spectrum.

THEORY

First-principles all-electron density functional electronic structure calculations for pure and H-doped \( \text{Ca}_{12-x}\text{Mg}_{x}\text{Al}_{14}\text{O}_{33} \) \( (x = 0, 1, \text{and 2}) \) were performed using the DMOl method, with the Perdew–Wang generalized gradient exchange-correlation functional and a doubled numeric basis set with local basis cutoffs of 11.0 a.u. Summations over the Brillouin zone were carried out using 24 special \( k \) points in the irreducible wedge. The structures were modeled within the cell of mayenite with 1 f.u. (i.e., 59 atoms per cell which combine into six cages) with periodic boundary condition. The equilibrium relaxed geometry of the structures was determined via total energy and atomic forces minimization; during the optimization, all atoms were allowed to move in \( x, y, \) and \( z \) directions, while the volume of the unit cell was fixed to the experimental value of the mayenite.\(^5\)

RESULTS AND DISCUSSION

The temperature dependence of the conductivity for the system \( \text{Ca}_{12-x}\text{Mg}_{x}\text{Al}_{14}\text{O}_{33} \) \( (x = 0, 0.1, 0.3, 0.5, 0.8, 1.0) \) is shown in Fig. 2. Although the systematic error is on the order of 5% due to uncertainty in geometric factors, the random uncertainty is on the order of the symbol size or less. As shown in Fig. 2(a), the conductivity is strongly dependent on the substitution level and rises reversibly from room temperature to approximately 130 °C. Once the temperature exceeds 135 °C \( (x < 0) \), the conductivity drops irreversibly as can be seen in Fig. 2(b). This drop is related to the amount of hydrogen released when the system crosses the decomposition temperature and will be discussed in detail later. Figure 2(a) also shows a slight shift of the decomposition temperatures to higher values as the substitution level increases.

The gradual shift of the decomposition temperature for the Mg-substituted mayenite results from two factors. First, the magnitude and resulting slope of conductivity versus temperature in the reversible conductivity regime (up to the decomposition temperature) decrease monotonically with increasing Mg content. The origin of this behavior is described below. Second, the postpeak behavior is virtually the same for all the Mg-substituted specimens, which is noticeably different from the nonsubstituted mayenite. This indicates that the kinetics of decomposition are different for the Mg-substituted materials than for the host material. This is not attributable to changes in microstructure; the grain sizes (5–10 \( \mu \)m) were very similar for all the specimens and there was no systematic change in specimen density with the Mg level. A possible explanation can be obtained from density functional calculations for the H-doped \( \text{Ca}_{11}\text{Mg}_{1}\text{Al}_{14}\text{O}_{33} \). From a comparison of the total energies of the 12 fully relaxed structures with different Mg site locations, we found that the H\(^-\) relaxes toward the Mg atom and demonstrates a strong Mg–H bonding, (cf. Fig. 3). This fact combined with the tendency for Mg atoms to cluster (see below), could account for the sluggish decomposition kinetics in the Mg-substituted samples.

![Fig. 2.](image)

FIG. 2. (a) Conductivity vs temperature of \( \text{Ca}_{12-x}\text{Mg}_{x}\text{Al}_{14}\text{O}_{33} \) \( (x = 0, 0.1, 0.3, 0.5, 0.8, 1) \) and (b) conductivity vs temperature of pure mayenite \( (\text{Ca}_{12}\text{Al}_{14}\text{O}_{33}) \) showing irreversible loss of conductivity above the decomposition temperature (135 °C).

![Fig. 3.](image)

FIG. 3. Calculated total energy, in eV, as a function of the H\(^-\) location between Mg and Ca atoms in H-doped \( \text{Ca}_{12}\text{Mg}_{1}\text{Al}_{14}\text{O}_{33} \). To obtain this curve, the positions of all atoms in the cell except H\(^-\) were fixed. In the insert: The calculated distances, in Angstroms, between the H\(^-\) ion and its nearest Ca (Mg) neighbors in the fully relaxed (i.e., all atoms in the cell were allowed to move) structures of: (a) \( \text{Ca}_{12}\text{Al}_{14}\text{O}_{33} \) and (b) \( \text{Ca}_{11}\text{Mg}_{1}\text{Al}_{14}\text{O}_{33} \) with the Mg atom located in the most energetically favorable position.
The thermoelectric coefficient was measured at room temperature for the different substitution levels ($x = 0.1, 0.3, 0.5, 0.8, 1.0$), and in the range between room temperature and 120 °C for the pure Ca$_{12}$Al$_{14}$O$_{33}$ specimen. Results are shown in Fig. 4. The coefficient obtained was negative ($n$ type) and, within experimental error, temperature independent with a value of approximately $-206 \pm 12 \, \mu V/K$. The fact that the conductivity is thermally activated while the Seebeck coefficient is temperature independent confirms that an activated mobility exists in the Ca$_{12-x}$Mg$_x$Al$_{14}$O$_{33}$ system; and this is indicative of a small polaron conduction mechanism. A small polaron is a localized charge carrier and its concomitant lattice polarization field, which migrates via a thermally activated hopping process.  

The thermoelectric coefficient for small polaron conductors is given by 

$$Q = \pm \frac{k}{e} \left( \ln \frac{2(1-c)}{c} \right),$$  

where $k/e$ is 86.14 $\mu V/K$, $c$ is the fraction of conducting ions of higher valence, the factor 2 accounts for the spin degeneracy, the entropy of the transport term is neglected, and the sign is determined by the nature of the polaron. It is negative if it forms around the trapped electron and positive for the trapped hole.

Based on the experimental result of the thermopower $-206 \, \mu V/K \pm 12 \, \mu V/K$ (see Fig. 2), the fraction of conducting species is constant and can be estimated as $0.155 \pm 0.02$, which is consistent with two carriers per unit cell moving along a 12-site hopping path ($c = 2/12 = 0.166$) as was suggested in our previous articles.

The expression for the small polaron conductivity is 

$$\sigma = \frac{\sigma_0}{T} \exp \left( \frac{-E_H}{kT} \right),$$

where $E_H$ is the activation or hopping energy and the pre-exponential factor $\sigma_0$ is given by

$$\sigma_0 = \frac{gNc(1-c)e^2a^2\nu_0}{k},$$

where $g$ is a geometrical factor on the order of unity (related to the coordination number of equivalent sites), $a$ is the jump distance between equivalent sites, $\nu_0$ is a lattice vibrational frequency, $N$ is the total density of conducting sites, and $c$ is the fraction of conducting species.

Figure 5(a) shows Arrhenius plots for the as-prepared and partially decomposed nonsubstituted sample ($x=0$). It can be seen that there is no difference between the slopes of the graphs, confirming that the hopping energy $E_H$ is the same for both cases and equal to 0.12 eV. This value compares favorably with values typical of small polaron behavior. The other hand, the values of the pre-exponential factor are significantly different, showing a 35% decrease between the values before and after decomposition. Considering the values of $a$, $g$, and $\nu_0$ in the pre-exponential expression [Eq. (6)] to be approximately constant, and also recalling the constant value obtained for the fraction of occupied sites ($c$) from thermopower data, it can be concluded that the variation of the total density of conducting sites ($N$) is responsible for the 35% drop in the pre-exponential factor. To confirm this, secondary ion mass spectrometry measurements were performed on similarly treated deuterated samples. The amount of deuterium was detected before and after the decomposition temperature, giving a 38% loss that matches nicely with the 35% drop in the total number of available sites. This result confirms not only that hydrogen loss occurs above a certain decomposition temperature, but also that the hydrogenous species are involved in the hopping path of the small polaron conduction model.
sites in the four conducting cages. Neesium ions have a tendency to occupy any of the calcium ions in the cage structure of mayenite.4

If magnesium ion eliminates all of the 12 sites on the conducting path, the conductive path will be 12 per unit cell. Using these values and Eqs. (5) and (6), we would expect an initial slope of −3 on Fig. 6, given by the following equation:

\[
\frac{\sigma(x)}{\sigma(0)} = \frac{\sigma_0}{\sigma(0,x)} = 1 - 3\lambda \chi (1 - \delta \chi).
\]

(7)

As can be seen in Fig. 6, this slope is greater than the initial slope of the experimental values, suggesting that not all magnesium atoms are occupying conducting sites. Indeed, under the rapid cooling of the sample annealed at 1300 °C, some of the Mg atoms can become “frozen” into the positions located far away from the hopping path although the corresponding total energy is found to be higher by at least 57 meV as compared to the most energetically favorable structure in which the Mg atom substitutes one of the hopping centers of the UV-activated system.

Now, to further understand the conductivity behavior with an increase of the Mg concentration, we compared the total energy of 11 structures of Ca_{10}Mg_{2}Al_{14}O_{33} with different site locations of one of the two Mg atoms (the other Mg atom was located at the most energetically favorable position as obtained from the calculations for Ca_{11}MgAl_{14}O_{33}). We found that the second Mg atom prefers to be located in the same cage with the first one,20 which thus demonstrates a tendency for Mg atoms to cluster.

Based on these results, we modify Eq. (7) by introducing the factor \(\chi\) which represents the fraction of magnesium in conducting cages. Moreover, if we consider that a second magnesium atom occupying a calcium site in a conducting cage results in no additional reduction of the conducting sites, the propensity of magnesium ions to cluster in the conducting cages can be accounted for by the term \(\delta\) in the following equation:

\[
\frac{N(x)}{N(0,x)} = \frac{\sigma_0}{\sigma(0,x)} = 1 - 3\lambda \chi (1 - \delta \chi).
\]

(8)

The best fit to the experimental data in Fig. 6 is \(\lambda=0.52\) (approximately 48% of the Mg ions occupying nonconducting cages) and \(\delta=0.5\) (a significant tendency for Mg to cluster).

This small polaron model also allows us to estimate the maximum conductivity achievable in the nonsubstituted mayenite sample, based on Eqs. (5) and (6). We assume that the fraction of conducting sites occupied by carriers remains 2/12, a jump frequency of \(\sim 10^{13}\) s\(^{-1}\), and an average jump distance of 0.3 nm.4 If we furthermore assume that all the \(\text{O}^{2-}\) reacts with \(\text{H}_2\) in the hydrogenation process, and that every \(\text{H}^+\) releases an electron after UV irradiation, giving four conducting cages per unit cell, the total density of conducting sites will be 12 per unit cell. Using these values and setting the hopping energy to zero, Eq. (5) gives a maximum conductivity of \(\sim 100\) S/cm, which agrees well with the results obtained by Matsushi et al. on Ca-treated, fully reduced (no \(\text{O}^{2-}\) species) mayenite.21

Furthermore, the carrier content \(N_c\) can be estimated by combining the fraction of occupied sites (from thermopower) with the total density of conducting sites (from \(\sigma_0\) coefficient). By combining the carrier content values with the conductivity, the mobility for the different substitution levels...
can be obtained as shown in Table I. These results are consistent with mobility values typical for small polaron conductivity (<1 cm²/V s).22

Finally, diffuse reflectance spectra shown in Fig. 8 characterize the optical properties of the Ca₁₂₋ₓMgₓAl₁₄O₃₃ system. Transmission increases with the doping level; however, the energy of the optical gap remains constant at 2.6 eV and is in good agreement with previous results.2 Increased transmission is consistent with the drop in the number of carriers from the electrical measurements (see Table I).

CONCLUSIONS

Electrical property measurements of Ca₁₂₋ₓMgₓAl₁₄O₃₃ (x=0, 0.1, 0.3, 0.5, 0.8, 1.0) offer evidence for small polaron conduction. The electrical conductivity is activated, whereas the thermoelectric coefficient remains temperature independent. The activation energy obtained for all the compositions is the same and equal to 0.12 eV. Carrier contents on the order of 10²⁰–10²¹ cm⁻³ were calculated from the conductivity pre-exponential factors and the thermopower values (≈206 μV/K). The mobilities obtained for Mg-substituted mayenite at 125 °C range from 4×10⁻³ to 7×10⁻³ cm²/V s and are consistent with the small polaron transport mechanism. This study shows that hydrogen is intimately involved in the conduction process, and loss of hydrogen above a decomposition temperature leads to a permanent reduction in conductivity. The optical absorption peak induced by the irradiation is 2.6 eV and does not shift with Mg concentration. For wavelengths longer than 500 nm there is a monotonic decrease in adsorption as the substitution level increases, which is consistent with the drop in electron population.

Finally, we found, both theoretically and experimentally, that Mg in the Ca₁₂Al₁₄O₃₃ system acts as a blocking agent on the conduction path. Although this produces a compositionally tunable conductivity, the resulting conductivities are limited by the low mobilities associated with small polaron conduction.

TABLE I. Transport data for Ca₁₂₋ₓMgₓAl₁₄O₃₃ at room temperature and 125 °C

| Substitution level (x) | Carrier content (cm⁻³) | Conductivity 25 °C (S/cm) | Mobility 25 °C (cm²/V s) | Conductivity 125 °C (S/cm) | Mobility 125 °C (cm²/V s) |
|------------------------|------------------------|---------------------------|-------------------------|---------------------------|-------------------------|
| 0.0                    | 1.08×10²¹              | 0.270                     | 1.57×10⁻³               | 0.711                     | 4.12×10⁻³               |
| 0.1                    | 6.43×10²⁰              | 0.242                     | 2.35×10⁻³               | 0.534                     | 5.18×10⁻³               |
| 0.3                    | 5.94×10²⁰              | 0.209                     | 2.19×10⁻³               | 0.515                     | 5.41×10⁻³               |
| 0.5                    | 4.72×10²⁰              | 0.165                     | 2.18×10⁻³               | 0.384                     | 5.07×10⁻³               |
| 0.8                    | 3.19×10²⁰              | 0.131                     | 2.56×10⁻³               | 0.318                     | 6.22×10⁻³               |
| 1.0                    | 2.08×10²⁰              | 0.106                     | 3.19×10⁻³               | 0.234                     | 7.04×10⁻³               |

FIG. 8. Optical properties for Ca₁₂₋ₓMgₓAl₁₄O₃₃.

ACKNOWLEDGMENTS

This work was supported by the MRSEC program of the National Science Foundation (Grant No. DMR-0076097) at the Materials Research Center at Northwestern University and by the DOE (Grant No. DE-FG02-88ER45372). M.I.B is supported by the U. S. Department of State through a Fulbright Scholarship. Computational resources have been provided by the DOE supported NERSC.

1B. Lewis and D. Paine, MRS Bull. 25, 22 (2002).
2K. Hayashi, S. Matsuuchi, T. Kamiya, M. Hirano, and H. Hosono, Nature (London) 419, 462 (2002).
3V. Bartl and T. Scheller, Neues Jahrb. Mineral., Monatsh. 35, 547 (1970); A. N. Christensen, Acta Chem. Scand., Ser. A 42, 110 (1987).
4J. E. Medvedeva, A. J. Freeman, M. I. Bertoni, and T. O. Mason, Phys. Rev. Lett. 93, 016408 (2004); J. E. Medvedeva and A. J. Freeman, Appl. Phys. Lett. 85, 955 (2004).
5There is some uncertainty regarding the precise UV-activated mechanism. An alternative involves the reaction: H−→H⁺+2e− and is the subject of ongoing work.
6H. Hosono and Y. Abe, Inorg. Chem. 36, 1192 (1997).
7T. P. Perdue and T. M. Mason, Key Eng. Mater. 125–126, 163 (1997).
8J. Ell, B. J. Wood, S. E. Dorris, and T. O. Mason, J. Solid State Chem. 82, 247 (1989).
9A. F. Joffe, Physics of Semiconductors (Inforsearch Ltd., London, 1960).
10H. L. Tuller and A. S. Nowick, J. Phys. Chem. Solids 38, 859 (1977).
11R. Dieckmann, C. A. Witt, and T. O. Mason, Ber. Bunsenges. Phys. Chem. 87, 495 (1983).
12E. Gartstein, T. O. Mason, and J. B. Cohen, Bull. Am. Ceram. Soc. 60, 375 (1981).
13E. Gartstein, T. O. Mason, and J. B. Cohen, Bull. Am. Ceram. Soc. 60, 375 (1981).
14J. P. Medvedeva, A. J. Freeman, M. I. Bertoni, and T. O. Mason, Appl. Phys. Lett. 85, 955 (2004).
15J. E. Medvedeva and Y. Wang, Phys. Rev. B 45, 13244 (1992); W. A. Shapley and D. P. Chong, Int. J. Quantum Chem. 81, 34 (2001).
16J. E. Medvedeva, A. J. Freeman, M. I. Bertoni, and T. O. Mason, Phys. Rev. Lett. 93, 016408 (2004); J. E. Medvedeva and A. J. Freeman, Appl. Phys. Lett. 85, 955 (2004).
17There is some uncertainty regarding the precise UV-activated mechanism. An alternative involves the reaction: H−→H⁺+2e− and is the subject of ongoing work.
18R. Dieckmann, C. A. Witt, and T. O. Mason, Ber. Bunsenges. Phys. Chem. 87, 495 (1983).
19H. L. Tuller and A. S. Nowick, J. Phys. Chem. Solids 38, 859 (1977).
20J. Z. Wang, S. Ford, and T. M. Mason, Key Eng. Mater. 125–126, 163 (1997).
21B. Delley, J. Chem. Phys. 113, 7756 (2000); 92, 508 (1990).
22B. Delley, J. Chem. Phys. 113, 7756 (2000); 92, 508 (1990).
23J. P. Perdue and W. A. Shapley and D. P. Chong, Int. J. Quantum Chem. 81, 34 (2001).
24J. Ell, B. J. Wood, S. E. Dorris, and T. O. Mason, J. Solid State Chem. 82, 247 (1989).
25A. F. Joffe, Physics of Semiconductors (Inforsearch Ltd., London, 1960).
26H. L. Tuller and A. S. Nowick, J. Phys. Chem. Solids 38, 859 (1977).
27R. Dieckmann, C. A. Witt, and T. O. Mason, Ber. Bunsenges. Phys. Chem. 87, 495 (1983).
28D. P. Karim and A. T. Aldred, Phys. Rev. B 20, 2255 (1979).
29T. O. Mason and H. K. Bowen, J. Am. Ceram. Soc. 64, 237 (1981).
30E. Gartstein, T. O. Mason, and J. B. Cohen, Bull. Am. Ceram. Soc. 60, 375 (1981).
31E. Gartstein, T. O. Mason, and J. B. Cohen, Bull. Am. Ceram. Soc. 60, 375 (1981).
32J. E. Medvedeva, A. J. Freeman, M. I. Bertoni, and T. O. Mason, and K. R. Poppelsmeier (unpublished).
33S. Matsuishi, Y. Toda, M. Miyakawa, K. Hayashi, T. Kamiya, M. Hirano, I. Tanaka, and H. Hosono, Science 301, 626 (2003).
34H. C. Chen, E. Gartstein, and T. O. Mason, J. Phys. Chem. Solids 43, 991 (1982).