Lithium and Gallium Vacancies in LiGaO$_2$ Crystals

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Lithium and gallium vacancies in LiGaO₂ crystals

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Lithium gallate (LiGaO₂) is a wide-band-gap semiconductor with an optical gap greater than 5.3 eV. When alloyed with ZnO, this material offers broad functionality for optical devices that generate, detect, and process light across much of the ultraviolet spectral region. In the present paper, electron paramagnetic resonance (EPR) is used to identify and characterize neutral lithium vacancies (V₀ Li) and doubly ionized gallium vacancies (V₂ Ga) in LiGaO₂ crystals. These S = 1/2 native defects are examples of acceptor-bound small polarons, where the unpaired spin (i.e., the hole) is localized on one oxygen ion adjacent to the vacancy. Singly ionized lithium vacancies (V⁺ Li) are present in as-grown crystals and are converted to their paramagnetic state by above-band-gap photons (x rays are used in this study). Because there are very few gallium vacancies in as-grown crystals, a post-growth irradiation with high-energy electrons is used to produce the doubly ionized gallium vacancies (V₂ Ga). The EPR spectra allow us to establish detailed models for the two paramagnetic vacancies. Anisotropy in their g matrices is used to identify which of the oxygen ions adjacent to the vacancy has trapped the hole. Both spectra also have resolved structure due to hyperfine interactions with ⁶⁹Ga and ⁷¹Ga nuclei. The V₀ Li acceptor has nearly equal interactions with Ga nuclei at two Ga sites adjacent to the trapped hole, whereas the V₂ Ga acceptor has an interaction with Ga nuclei at only one adjacent Ga site. Published by AIP Publishing. https://doi.org/10.1063/1.5050532

I. INTRODUCTION

Lithium gallate (LiGaO₂) is an ultrawide-band-gap semiconductor¹,² with a wurtzite-like crystal structure. Its optical absorption edge is in the 5.3–5.7 eV range.³–⁷ This material is a ternary analog of ZnO. Replacing half the Zn²⁺ ions with Li⁺ ions and half with Ga³⁺ ions in an ordered arrangement gives LiGaO₂. Alloying LiGaO₂ with ZnO is expected to produce crystals appropriate for ultraviolet optical applications.⁸–¹⁰ As suitable shallow donors and acceptors are identified, the LiGaO₂-ZnO mixed materials system will allow laser diodes and photodetectors to be fabricated that operate across much of the ultraviolet region. LiGaO₂ is also a candidate for phosphor and radiation-detector applications.¹²–¹⁶ This ternary material has both monovalent (Li⁺) and trivalent (Ga³⁺) cation sites and thus provides a variety of doping possibilities. These include transition-metal ions and rare-earth ions, Group I and Group III isovalent impurities, and Group II and Group IV donors and acceptors. Many doping choices offer broad functionality for diverse applications of LiGaO₂ and its alloys with ZnO.

In the present paper, we use electron paramagnetic resonance (EPR) to investigate native acceptors in LiGaO₂ crystals. Similar studies have been reported for LiAlO₂ crystals.¹⁷,¹⁸ Spectra from lithium vacancies and gallium vacancies are observed in our LiGaO₂ crystals. In both cases, the hole is localized on one oxygen ion adjacent to the vacancy, in a small-polaron configuration.¹⁹,²⁰ Anisotropy in the g matrices allows us to determine which oxygen ion traps the hole. The V₀ Li and V₂ Ga spectra in LiGaO₂ have resolved hyperfine structure due to interactions of the trapped hole with the adjacent ⁶⁹Ga and ⁷¹Ga nuclei. Nearly equal interactions with nuclei at two gallium sites are seen in the V₀ Li spectrum, whereas interactions with nuclei at only one gallium site are observed in the V₂ Ga spectrum. These different hyperfine patterns easily allow the V₀ Li and V₂ Ga spectra to be individually recognized. Although their spectra are obtained at low temperature, the paramagnetic charge states of both vacancies are stable at room temperature. Information about the small-polaron characteristics and the thermal stabilities of these native defects will be useful when searching for acceptor dopants for LiGaO₂. Lithium diffusion studies will also benefit from a spectroscopic method that monitors the presence of lithium vacancies.²¹,²²

II. EXPERIMENTAL

Undoped LiGaO₂ crystals, grown by the Czochralski method, were obtained from the MTI Corporation (Richmond, CA). They are highly resistive, thus indicating that the Fermi level is near mid-gap. Lithium vacancies are present in some, but not all, of the as-grown LiGaO₂ crystals supplied by MTI, whereas gallium vacancies are not present in any of the as-grown crystals. Spectra from two LiGaO₂ crystals, labeled Sample A and Sample B, are reported in the present investigation. Sample A was used to investigate neutral lithium vacancies and was only irradiated with x rays (i.e., ionizing radiation). By trapping a hole on an adjacent oxygen ion, the irradiation with x rays (60 kV, 30 mA) at room temperature converted nonparamagnetic lithium vacancies in this sample to a paramagnetic charge state that could...

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be observed at low temperature with EPR. Sample B was only electron irradiated and was used to investigate doubly ionized gallium vacancies. This crystal was irradiated near room temperature with high-energy electrons (1 MeV, 5 μA) from a Van de Graff accelerator at Wright State University. During the irradiation, the crystal was in contact with a water-cooled heat sink to minimize heating by the electron beam. Momentum-conserving displacement events, initiated by the high-energy electrons, produced a large concentration of paramagnetic gallium vacancies in the LiGaO₂ crystal, without requiring a subsequent irradiation with x rays. The EPR spectra reported in this paper represent approximate defect concentrations of 2.3 × 10¹⁸ cm⁻³ for the V⁰_Li acceptors and 1.8 × 10¹⁷ cm⁻³ for the V²⁺/C⁰_Ga acceptors.

LiGaO₂ crystals are orthorhombic (space group Pna2₁), with lattice constants a = 5.402 Å, b = 6.372 Å, and c = 5.007 Å at room temperature. Figure 1 is a ball-and-stick representation of the crystal. The relative x,y,z positions of the ions in Fig. 1 are given in Table I. In this material, each oxygen ion has two lithium neighbors and two gallium neighbors, each lithium ion has four oxygen neighbors, and each gallium ion has four oxygen neighbors. The Li⁺ sites are all equivalent, and the Ga³⁺ sites are all equivalent. Oxygen ions occupy two crystallographically inequivalent sites (these are labeled O_l and O_h in Fig. 1). The two oxygen sites are most easily distinguished by which cation, a Li⁺ or a Ga³⁺, is the nearest neighbor along the c axis (i.e., the [001] direction). The O_l ions have an adjacent lithium ion in the c direction and the O_h ions have an adjacent gallium ion in the c direction.

The EPR spectra were taken with a Bruker EMX spectrometer operating near 9.40 GHz. A Bruker NMR teslameter was used to measure the static magnetic field, and an Oxford helium-gas flow system controlled the sample temperature. The isotropic Cr³⁺ signal (g = 1.9800) in an MgO crystal was used to make corrections for the small differences in magnetic field at the NMR probe tip and the sample position.

### III. NEUTRAL LITHIUM VACANCY (V⁰_Li)

Figure 2(a) shows the EPR spectrum from the neutral lithium vacancy (V⁰_Li) in Sample A. This spectrum, taken at 55 K with the magnetic field along the c axis, was obtained after the as-received crystal was irradiated at room temperature with x rays. Singly ionized lithium vacancies (V⁺/C⁰_Li) in the as-grown crystal are converted to their paramagnetic neutral charge state (V⁰_Li) during the irradiation. Many of the free electrons and holes created by the x rays quickly recombine. A small portion of the holes, however, are trapped on oxygen ions adjacent to lithium vacancies, thus forming the V⁰_Li centers. A corresponding number of electrons are trapped at unidentified defects (possibly oxygen vacancies or impurities). Heating the crystal above 150 °C destroys the V⁰_Li spectrum and returns the crystal to its pre-irradiated state.

![Figure 1. Schematic representation of the LiGaO₂ crystal structure. Lithium ions are green, gallium ions are purple, and oxygen ions are red. The two inequivalent oxygen sites are labeled O_I and O_II. Each lithium and gallium ion has four oxygen neighbors and each oxygen ion has two lithium and two gallium neighbors.](image)

![Figure 2. (a) EPR spectrum of the neutral lithium vacancy (V⁰_Li) in LiGaO₂ (Sample A). The crystal was irradiated at room temperature with x rays, and then the spectrum was taken at 55 K with the magnetic field along the c direction. The microwave frequency was 9.406 GHz. Stick diagrams illustrate the separate hyperfine contributions from the ⁶⁹Ga and ⁷¹Ga nuclei. (b) Simulated spectrum produced with the SimFonia computer program.](image)

**TABLE I.** Relative positions (in Å units) of ions in LiGaO₂ based on the room-temperature lattice parameters reported by Marzio. The ion labeling scheme in Fig. 1 is used.

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![Image 1. Schematic representation of the LiGaO₂ crystal structure. Lithium ions are green, gallium ions are purple, and oxygen ions are red. The two inequivalent oxygen sites are labeled O_I and O_II. Each lithium and gallium ion has four oxygen neighbors and each oxygen ion has two lithium and two gallium neighbors.](image)

![Image 2. (a) EPR spectrum of the neutral lithium vacancy (V⁰_Li) in LiGaO₂ (Sample A). The crystal was irradiated at room temperature with x rays, and then the spectrum was taken at 55 K with the magnetic field along the c direction. The microwave frequency was 9.406 GHz. Stick diagrams illustrate the separate hyperfine contributions from the ⁶⁹Ga and ⁷¹Ga nuclei. (b) Simulated spectrum produced with the SimFonia computer program.](image)
A related thermoluminescence (TL) peak is observed near 110 °C in our x-ray irradiated LiGaO₂ crystals. The emitted light associated with this TL peak has a maximum near 350 nm.

The S = 1/2 spectrum from the neutral lithium vacancies (V₀ Li) consists of a symmetrical pattern of resolved hyperfine lines with differing intensities caused by interactions with ⁶⁹Ga and ⁷¹Ga nuclei. Both isotopes have I = 3/2 nuclear spins and they are 60.1% and 39.9% abundant, respectively. Their nuclear magnetic moments are ⁶⁹μ = +2.0166μ₀ and ⁷¹μ = +2.5623μ₀.⁴⁴ The complicated V₀ Li spectrum in Fig. 2(a) is explained by having nearly equal hyperfine interactions with Ga nuclei at two gallium sites adjacent to the trapped hole. This is consistent with our detect model since each with Ga nuclei at two gallium sites adjacent to the trapped hole on the oxygen ion. Hyperfine lines from the ⁷Li nucleus are not resolved in the V₀ Li spectrum, whereas lines from the ⁶⁹,⁷¹Ga nuclei are well resolved. This agrees with our model, as atomic calculations²⁵,²⁶ predict that the isotropic ⁷Li hyperfine parameters will be approximately a factor of 30 smaller than the isotropic ⁶⁹,⁷¹Ga parameters when similar amounts of unpaired s-like spin density are on the Li and Ga ions.

With two adjacent sites for Ga, there are three combinations of the two Ga isotopes that contribute to the observed hyperfine pattern in Fig. 2(a). These are (i) two ⁶⁹Ga nuclei, (ii) one ⁶⁹Ga nucleus and one ⁷¹Ga nucleus, and (iii) two ⁷¹Ga nuclei. The relative distributions of these three combinations are 36.1%, 48.0%, and 15.9%, respectively. In Fig. 2(a), each combination is represented by a stick diagram above the experimental spectrum. Although the hyperfine interactions at the two Ga sites are not exactly equal, the stick diagrams are drawn for equal interactions. The relative lengths of the vertical lines in these diagrams reflect the natural abundances of the two Ga isotopes. The uppermost stick diagram illustrates the seven lines (with relative intensities of 1:2:3:4:3:2:1) that are produced when the unpaired spin interacts equally with two ⁶⁹Ga nuclei. The lowest stick diagram in Fig. 2(a) is the sum of the three upper stick diagrams and should be directly compared to the experimental spectrum. An EPR spectrum with a hyperfine pattern very similar to our V₀ Li spectrum has been recently reported for the doubly ionized gallium vacancy (V₂ Ga) in a β-Ga₂O₃ single crystal.²⁷

Evidence that the hyperfine interactions with the nuclei at the two neighboring Ga sites, although similar, are not equal is found in the relative intensities of the lines in the experimental spectrum in Fig. 2(a). If the two Ga sites have equal interactions, the intensity of the middle line in the spectrum should be a factor of 14.3 greater than the intensity of the lowest-field line. The experimental ratio in Fig. 2(a), however, is only about 8.5. This observed lower ratio suggests that the center line is slightly split due to inequivalent interactions at the neighboring Ga sites. This splitting is not resolved in the spectrum, but it does significantly reduce the intensity of the center line. The lowest-field line does not split when the two Ga sites have inequivalent interactions. Thus, unlike the center line, its intensity is unaffected by the inequivalency.

The experimental EPR spectrum in Fig. 2(a) has a set of underlying weak lines that slightly distort the V₀ Li signals in the magnetic field region above 331.5 mT. Consequently, Ga hyperfine parameters that describe the c axis V₀ Li spectrum were obtained from the undistorted low-field side. The lowest line and the middle line in Fig. 2(a) are separated by 9.675 mT. According to the ⁷¹Ga-⁷¹Ga stick diagram, the averaged value of Aₐ (⁷¹Ga) for these nuclei at the two Ga sites is equal to one-third of this separation. (Here, Aₐ represents the hyperfine interaction for each nucleus when the magnetic field is along the c direction.) This gives an averaged value of 3.22 mT for Aₐ (⁷¹Ga). A corresponding averaged value of 2.53 mT for Aₐ (⁶⁹Ga) is obtained using the ratio of magnetic moments. A simulated c axis spectrum, produced with the SimFonia program from Bruker, is shown in Fig. 2(b). In this simulation, the hyperfine interactions at the two neighboring gallium sites are 4% different. The ⁶⁹Ga and ⁷¹Ga parameters used in the simulation were 2.59 and 3.29 mT for one gallium site and 2.49 and 3.16 mT for the other gallium site.

Averaged hyperfine values for the ⁶⁹,⁷¹Ga nuclei at the two Ga sites adjacent to the trapped hole were also obtained for the V₀ Li acceptor when the magnetic field was aligned along the a and b directions in the crystal. Results for the three directions (a, b, and c) are listed in Table II. For these high-symmetry directions, all crystallographically equivalent orientations of the defect are also magnetically equivalent, and the EPR spectra have their simplest form. Nearly identical hyperfine patterns were observed for each of the three directions of magnetic field, thus establishing the isotropic nature of the Ga interactions. The results in Table II provide direct evidence for the small-polaron model of the V₀ Li acceptor. Specifically, small values for the strength of the Ga hyperfine interactions indicate that only 1.1% of the unpaired spin is in 4s orbitals on the two Ga neighbors.²⁵ This leaves nearly all the remaining unpaired spin density in a p orbital on the oxygen ion.

The g values in Table II were obtained from measurements of the position of the center line in the V₀ Li spectrum when the static magnetic field was along the a, b, and c directions. For these directions, the crystallographically equivalent orientations of the defect are all magnetically equivalent and the centers of the hyperfine patterns are easily identified. The EPR spectrum reached a minimum field

<table>
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<th>Direction of magnetic field</th>
<th>g value</th>
<th>⁶⁹Ga (mT)</th>
<th>⁷¹Ga (mT)</th>
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<tbody>
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<td>3.09</td>
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<tr>
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<td>c crystal axis</td>
<td>2.0366</td>
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<td>3.22</td>
</tr>
</tbody>
</table>

Table II. Parameters describing the EPR spectra of the neutral lithium vacancy (V₀ Li) in a LiGaO₂ crystal. The oxygen ion trapping the hole has two slightly inequivalent Ga neighbors. A g value and the average of the hyperfine parameters for the ⁶⁹,⁷¹Ga nuclei at the two sites were obtained when the magnetic field was along each of the three crystallographic axes. The estimated error is ±0.0005 for the g values and ±0.05 mT for the hyperfine values.
position (i.e., a turning point) when the magnetic field was along the c direction and reached a maximum field position when the magnetic field was near the a direction. These observations indicate that the principal axes of the g matrix are near the a, b, and c crystal directions and that the values listed in Table II are very near the principal values of the g matrix. The strong overlap of lines caused by the presence of two magnetically inequivalent orientations of the defect prevented useful measurements of the angular dependence of the g matrix within the a-b, b-c, and c-a planes.

The small and positive g shifts observed for the V\textsubscript{Li}\textsuperscript{0} acceptor are consistent with a model that has the unpaired spin (i.e., the hole) located in a p orbital on one oxygen ion adjacent to the lithium vacancy. Acceptor-bound small polarons of this type have been extensively investigated in oxide crystals.\textsuperscript{17,19,20,27} The oxygen trapping the hole is an O\textsuperscript{−} ion with a 2p\textsuperscript{5} configuration (2p\textsubscript{x},2p\textsubscript{y},2p\textsubscript{z}). The local crystalline electric field removes the threefold orbital degeneracy of this 2P state (L = 1, S = 1/2) and forms three energy levels (E\textsubscript{1}, E\textsubscript{2}, and E\textsubscript{3}). In this simplified picture, E\textsubscript{1} is the ground state with the hole in the p\textsubscript{x} orbital and E\textsubscript{2} and E\textsubscript{3} are excited states with the hole in the p\textsubscript{x} and p\textsubscript{y} orbitals of the ion, respectively. The spin-orbit interaction then mixes these excited states with the ground state and gives the following first-order expressions for the principal g values:\textsuperscript{28}

\[ g_1 = g_e - \frac{2\lambda}{E_2 - E_1}, \quad g_2 = g_e - \frac{2\lambda}{E_3 - E_1}, \quad g_3 = g_e - \frac{2\lambda}{E_3 - E_1}. \quad (1) \]

The principal direction corresponding to the g\textsubscript{1} principal value is parallel to the unique axis of the p\textsubscript{x} orbital. In these equations, the spin-orbit coupling constant \( \lambda \) for an O\textsuperscript{−} ion\textsuperscript{29} is −135 cm\textsuperscript{−1} and g\textsubscript{e} is 2.0023. The positive g shifts (i.e., values greater than 2.0023) reported in Table II for the V\textsubscript{Li}\textsuperscript{0} acceptor are a result of the negative sign for \( \lambda \).

The value of 2.0088 for g\textsubscript{e} in Table II is close to 2.0023, which suggests that the p\textsubscript{x} orbital containing the unpaired spin is oriented near the a direction in the crystal. Figure 3 illustrates our proposed model for the V\textsubscript{Li}\textsuperscript{0} acceptor. The lithium vacancy is at the Li(1) position and the hole is localized on the adjacent O\textsubscript{iii}(2) ion. Nuclei at the Ga(6) and Ga(7) sites are responsible for the resolved hyperfine structure seen in the EPR spectra. Before lattice relaxation, the Li(1) and O\textsubscript{ii}(2) sites are 1.995 Å apart and the line joining them makes an angle of 18.2° with the a axis of the crystal. The p\textsubscript{x} orbital representing the hole, shown in blue in Fig. 3, is pointing toward the center of the Li vacancy. This orientation of the p\textsubscript{x} orbital corresponds to the minimum for the energy of the ground state of the neutral acceptor and establishes the importance of the electrostatic attraction between the positive hole and the “effective” negative charge of the lithium vacancy. The difference between the value of 2.0088 for g\textsubscript{e} and an anticipated g\textsubscript{1} value nearer 2.0023 is explained by the relatively small 18.2° angle that the unique axis of the p\textsubscript{x} orbital (and thus the principal direction for g\textsubscript{1}) makes with the a direction in the crystal. In other words, the measured value of g\textsubscript{e} is slightly greater than 2.0023 because the a direction, although close, is not a principal direction of the g matrix for the V\textsubscript{Li}\textsuperscript{0} acceptor.

IV. DOUBLY IONIZED GALLIUM VACANCY (V\textsubscript{Ga}\textsuperscript{2−})

Figure 4(a) shows the EPR spectrum from the doubly ionized gallium vacancy (V\textsubscript{Ga}\textsuperscript{2−}) in Sample B. This spectrum was taken at 93 K with the magnetic field along the c axis. There are six resolved hyperfine lines and g\textsubscript{e} is 2.0032. These paramagnetic V\textsubscript{Ga}\textsuperscript{2−} acceptors were produced by a “knock-on” process in the crystal during an irradiation near room temperature with 1 MeV electrons. A subsequent exposure to ionizing radiation was not needed, as the doubly ionized charge state of the gallium vacancy was thermally stable at room temperature in our crystal.

Similar to the V\textsubscript{Li}\textsuperscript{0} acceptor, the V\textsubscript{Ga}\textsuperscript{2−} acceptor has a hole localized on one oxygen ion adjacent to the vacancy. With fewer hyperfine lines than the V\textsubscript{Li}\textsuperscript{0} spectrum, the experimental V\textsubscript{Ga}\textsuperscript{2−} spectrum in Fig. 4(a) is explained by interactions with 69Ga and 71Ga nuclei at only one neighboring Ga site. This is consistent with our defect model since each oxygen ion in the LiGaO\textsubscript{2} crystal has two Li\textsuperscript{+} ions and two Ga\textsuperscript{3+} ions as nearest neighbors. With one of the gallium ions missing, the oxygen ion with the trapped hole has two neighboring Ga\textsuperscript{3+} ions and one neighboring Ga\textsuperscript{3+} ion. Hyperfine interactions with the adjacent 7Li nuclei are not resolved in the V\textsubscript{Ga}\textsuperscript{2−} spectrum, whereas interactions with the 69,71Ga nuclei at the one neighboring Ga site are easily observed.

Each Ga isotope has an I = 3/2 nuclear spin and produces four equally spaced hyperfine lines, as illustrated by the stick diagrams above the experimental spectrum in Fig. 4(a). Only six lines are resolved in this spectrum because of the strong overlap of the middle lines within each set of four. The difference in the separation of hyperfine lines in the two sets is directly related to the ratio of the nuclear magnetic moments for 69Ga and 71Ga. From the line positions in Fig. 4(a), we find A\textsubscript{e}(69Ga) = 3.59 mT and
AcGa = 4.60 mT for the V2/C0 Ga acceptor. Using these results, the simulated c-axis spectrum shown in Fig. 4(b) was produced with the SimFonia program. Table III contains the experimental g values and hyperfine parameters for the V2/C0 Ga acceptor when the magnetic field is along the a, b, and c directions. These results show that the 69Ga and 71Ga hyperfine matrices for the V2/C0 Ga acceptor are nearly isotropic. The center of the EPR spectrum reached a minimum field position when the magnetic field was along the b direction and reached a maximum field position when the magnetic was near the c direction, thus indicating that the g values listed in Table III are very near the principal values of the g matrix.

The anisotropy of the g matrix is used to construct a model for the V2/C0 Ga acceptor. When a hole is trapped on one oxygen ion next to a gallium vacancy, Eq. (1) predicts that one principal value of the g matrix will be very near 2.0023 and the other two principal values will have small, but positive, g shifts. The value of 2.0032 that we measure for gc is very close to 2.0023, whereas the values for ga and gb are 2.0155 and 2.0551, respectively. This strongly suggests that the pz orbital containing the unpaired electron spin is aligned along the c direction in the crystal. Our model for the V2/C0 acceptor in LiGaO2 is shown in Fig. 5. The gallium vacancy is at the Ga(4) site and the hole is localized on the adjacent OII(4) ion. Nuclei at the Ga(3) site are responsible for the resolved hyperfine structure seen in the EPR spectra.

V. SUMMARY

Electron paramagnetic resonance (EPR) has been used to identify and characterize native acceptors in wurtzite-like LiGaO2 crystals. Neutral lithium vacancies (V0Li) and doubly ionized gallium vacancies (V2/C0 Ga) are observed. These defects provide clear examples of acceptor-bound small polarons, where the unpaired spin (i.e., the hole) is localized on one oxygen ion adjacent to the vacancy. Resolved hyperfine structure from neighboring 69,71Ga nuclei and anisotropy in the g matrices are used to construct specific models for these acceptors. In both defects, the hole is located at an OII ion, as it forms the shortest bond with neighboring lithium and gallium ions.

The thermal stability of the paramagnetic charge states of these cation vacancies at room temperature is days for V0Li and more than one year for V2/C0 Ga. This suggests that they are deep levels, as expected for acceptor-bound polarons in oxide crystals. If acceptors such as Mg can be placed on a Ga site in Li-rich, Ga-poor material, their neutral state may also be deep, and thus not useful for devices, because of the formation of similar small polarons. In the search for suitable p-type dopants for LiGaO2, nitrogen ions replacing oxygen ions may be a more likely solution.

### Table III

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</tbody>
</table>

FIG. 4. (a) EPR spectrum of the doubly ionized gallium vacancy (V2Ga) in LiGaO2 (Sample B). The crystal was irradiated near room temperature with 1 MeV electrons. The spectrum was then taken at 93 K with the magnetic field along the c direction. The microwave frequency was 9.404 GHz. (b) A simulation spectrum produced with the SimFonia computer program.

FIG. 5. Model of the doubly ionized gallium vacancy (V2Ga) in LiGaO2. The trapped hole (shown in blue) is localized in a p orbital on the OII(4) oxygen ion with the gallium vacancy at the Ga(4) position. The 69Ga and 71Ga nuclei at the Ga(3) site are responsible for the observed hyperfine.
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