Microwave Directional Dichroism Resonant with Spin Excitations in the Polar Ferromagnet GaV$_4$S$_8$

Y. Okamura,[1] S. Seki,[2] S. Bordács,[3,4] Á. Butykai,[3] V. Tsurkan,[5] I. Kézsmárki,[3] and Y. Tokura[1,2]

[1]*Department of Applied Physics and Quantum Phase Electronics Center, University of Tokyo, Tokyo 113-8656, Japan*
[2]*RIKEN Center for Emergent Matter Science (CEMS), Wako 351-0198, Japan*
[3]*Department of Physics, Budapest University of Technology and Economics*
and MTA-BME Lendulet Magneto-optical Spectroscopy Research Group, 1111 Budapest, Hungary
[4]*Hungarian Academy of Sciences, Premium Postdoctor Program, 1051 Budapest, Hungary*
[5]*Experimental Physics V, Center for Electronic Correlations and Magnetism, University of Augsburg,*
86159 Augsburg, Germany

We have investigated the directional dichroism of magnetic resonance spectra in the polar ferromagnet GaV$_4$S$_8$. While four types of structural domains are energetically degenerated under a zero field, the magnetic resonance for each domain is well separated by applying magnetic fields due to uniaxial magnetic anisotropy. Consequently, a directional dichroism as large as 20% is clearly observed without domain cancellation. The present observation therefore demonstrates that not only magnetoelectric monodomain crystals but also magnetoelectric multidomain specimens can be used to realize microwave (optical) diodes owing to the lack of inversion domains.

Light-matter interaction plays a fundamental role in various fields of physics, chemistry, and engineering. The optical response is highly sensitive to the material properties even beyond the static ones and often helps to realize versatile functionalities. Multiferroics, materials with simultaneously broken spatial inversion and time reversal symmetry, provide a unique arena to study novel optical phenomena that cannot show up in conventional ferromagnets or ferroelectrics [1–4]. Such an unconventional optical response, called the optical magnetoelectric (ME) effect, occurs when coexisting magnetic and ferroelectric orders simultaneously interact with the oscillating electric field and magnetic field of light. The discovery of this phenomenon was first reported for a noncentrosymmetric antiferromagnet Cr$_2$O$_3$ [5], where the polarization rotation of light is reversed depending on the propagation direction of the light, although the effect is rather small. Another type of the optical ME effect, termed nonreciprocal directional dichroism (DD), was demonstrated subsequently, where the light beams traveling in opposite directions are absorbed differently [6–21]. It has been revealed that the DD can become large and even show a cloaking function, thus being particularly important from the viewpoint of future applications [13,14].

The DD is observed for various types of excitations from x-ray to microwave-frequency regions in multiferroic materials [6–21]. Among them, the DD can often be large for the collective spin excitations that are both electric- and magnetic-dipole active owing to the strong ME coupling. These novel elementary excitations, referred to as the magnetoelectric resonances, are accompanied by the resonant motion of the magnetization (M) as well as that of the electric polarization (P), which enhances the ME coupling and leads to large DD. Recently, as a new guiding principle towards large DD, magnetoelectric resonances in the type-I multiferroics, in which magnetic orders develop within a preexisting ferroelectric or pyroelectric phase [22], have been attracting growing interest because this class of materials often exhibits large spin-induced changes of the electric polarization [15–17]. The lacunar spinel GaV$_4$S$_8$ is a new member of the type-I multiferroics, which is an excellent candidate material to show large DD since it is a rare polar ferromagnet [23–27].

In this Letter, we have studied the microwave DD at the ferromagnetic resonance in the polar magnetic semiconductor GaV$_4$S$_8$. While the sample consists of multiple polar rhombohedral structural domains in which the magnetic states are degenerate in zero magnetic field, the degeneracy of the magnetic excitations on the different types of domains is split in external magnetic fields owing to the axial magnetic anisotropy. As the result, the DD as large as 20% is clearly observed free from cancellation among the domains. The magnitude critically depends on each domain, indicating the vital role of the microscopic ME coupling besides macroscopic lifted symmetry.

The lacunar spinel structure of GaV$_4$S$_8$, which has the space group symmetry $F\bar{4}3m$ at room temperature, is shown in Fig. 1(a) [25,26]. It consists of a network of $(V_4S_4)^{5+}$ clusters that form a face centered cubic lattice. Below 42 K, due to cooperative Jahn-Teller distortion, the
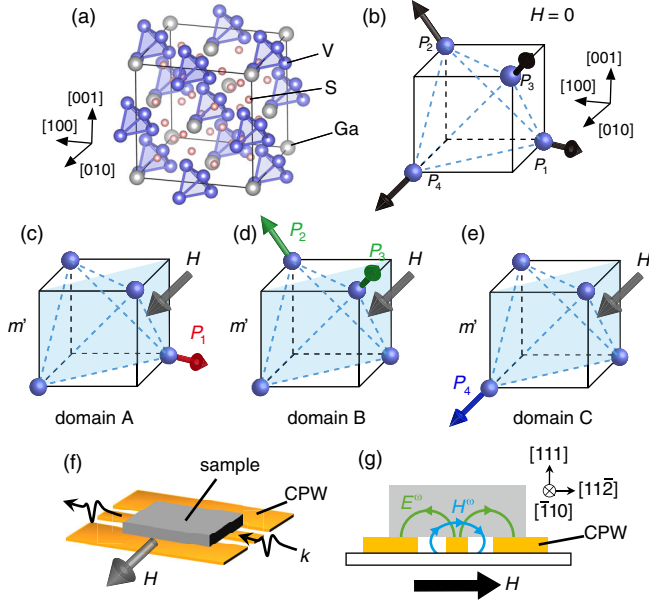
FIG. 1. (a) Crystal structure of GaV_4S_8 at room temperature. (b) The schematic illustration of V_4 clusters. The V ion is displaced along four equivalent cubic $\langle 111 \rangle$ axes below the structural phase transition temperature. [(c)–(e)] Three types of structural domains classified based on the relationship with respect to $H \| [11\bar{2}]$. m' represents a mirror plane combined with time reversal operation as indicated by the blue shaded region. (f) Schematic illustration of the coplanar wave guide (CPW). (g) Cross-sectional view of the CPW; the spatial distribution of microwave electric and magnetic fields is illustrated.

triply degenerate molecular orbitals of V_4 clusters are lifted through the elongation of the lattice along any of the four cubic $\langle 111 \rangle$ axes [Fig. 1(b)]. The crystal structure then becomes polar ($R3m$) with a relatively large pyroelectric polarization, $P \simeq 1\ \mu C/cm^2$ [24]. It should be noted that the opposite-polarization domains may exist due to the noncentrosymmetric nature of the room temperature cubic $F\bar{4}3m$ phase. However, the previous piezoresponse force microscopy (PFM) and static pyro- or magnetocurrent measurements indicate that such inversion domains do not exist in these single crystals [24,27]. Therefore, we analyze the observed DD without considering the coexistence of inversion domains, which would lead to partial cancellation of the DD; i.e., its magnitude characteristic to a single inversion domain would be larger than the observed one. The magnetic transition occurs at $T_c \sim 12.7$ K, well below the structural transition temperature. The cycloidal spin state is stabilized at zero field between 6 and 12.7 K and turns into a collinear field-polarized ferromagnetic state in moderate magnetic fields (H) [28]. The Néel-type skyrmion lattice is observed in a specific temperature and magnetic field region [23,28]. The stability of each magnetic phase critically depends on the magnetic-field direction due to the easy-axis anisotropy with respect to the rhombohedral axis, as discussed later [Figs. 1(c)–1(e)].

Single crystals of GaV_4S_8 were grown with chemical vapor transport and the detail of the growth procedure is described elsewhere [23]. We performed broadband microwave spectroscopy to measure the transmission coefficient of the sample mounted on a CPW as shown in Fig. 1(f). The signal line was designed to be 20 μm in width so that it is much smaller than the sample width; the sample dimension is typically $\sim 2 \times 2 \times 1$ mm^3. The directions of oscillating magnetic and electric field of the microwave, denoted as H^ω and E^ω, respectively, depend on the position in space: the H^ω just above the center of signal line is parallel to the plane of the CPW, while the H^ω between the signal line and the ground is perpendicular to the plane [Fig. 1(g)]. The transmission coefficient for a microwave propagating along k^ω and $-k^\omega$ directions is denoted as S_{12} and S_{21}, respectively, and was recorded with a vector network analyzer (Agilent Technology, E8363C). The absorption spectrum associated with the magnetic excitations, denoted as ΔS_{12} (or ΔS_{21}) for the $+k^\omega$ (or $-k^\omega$) microwave, was obtained by calculating $\Delta S_{12(21)} = -S_{12(21)} + S_{12(21)}(T > T_c)$, where $S_{12(21)}(T > T_c)$ taken at $T > T_c$ does not contain the magnetic signal (0.01–35 GHz) (for more details, see Supplemental Material [29]).

To observe the DD in this material, we focus on the magnetic resonance when $k^\omega \| [1\bar{1}0]$ and $H \| [11\bar{2}]$, as sketched in Fig. 1(g), because the DD can emerge for every type of structural domain from the symmetry point of view, as shown in Figs. 1(c)–1(e). For this choice of the magnetic field, the single mirror plane remains for domain A and C. In contrast, two kinds of domain Bs are interchanged by this mirror reflection combined with time reversal operation. This unique mirror plane combined with the time reversal operation allows the emergence of DD for light beams propagating perpendicular to it but not for beams traveling parallel to the plane [30]. This symmetry is also compatible with a phenomenological toroidal moment $T = P \times M$ pointing perpendicular to the mirror plane [31]. In all of the three kinds of magnetic domains classified in terms of the angle between their $\langle 111 \rangle$-type easy axes and the direction of the magnetic field (90°, 61.9°, and 19.5°), T is finite, although it has different magnitudes in the different domains [Figs. 1(c)–1(e) and Figs. 4(b) and 4(c)].

As revealed in the previous magnetic-resonance study, the resonance frequency critically depends on the magnetic domains, or equivalently, the magnetic-field direction due to the uniaxial anisotropy [32,33]. Thus, we first investigated the evolution of the magnetic resonance with field for the different magnetic domains. Figure 2(a) shows the magnetic resonance spectra at 8 K for $k^\omega \| [1\bar{1}0]$ in various magnetic fields $H \| [11\bar{2}]$. In the cycloidal phase at zero field, three resonance peaks are observed at 5, 18, and 27 GHz, in accord with a previous study [32]. By applying the magnetic field of 100 mT, the intensity of resonance modes at 5 and 18 GHz is weakened and enhanced,
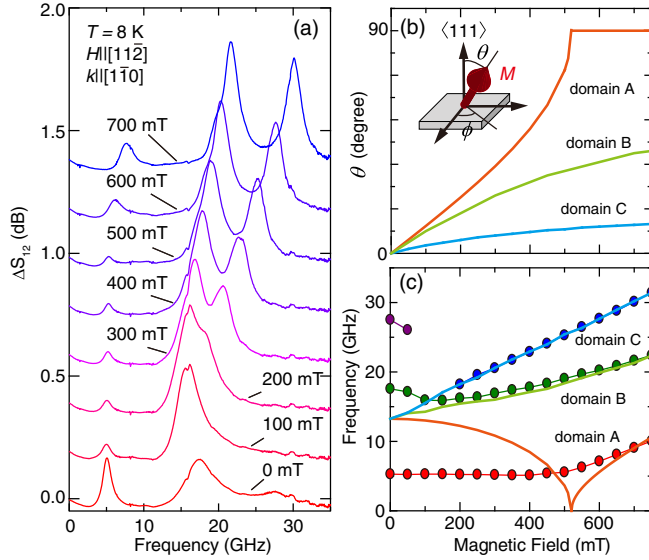
FIG. 2. (a) The magnetic resonance spectra for each magnetic field when $k^\omega\|[1\bar{1}0]$ and $H\|[11\bar{2}]$. The spectra are shifted vertically for clarity. (b) Calculated angle of θ, i.e., the angle between the magnetization direction and polar $\langle 111 \rangle$ axis, numerically calculated based on Eq. (1). (c) The magnetic-field dependence of the resonance frequency (circles). The calculated resonance frequency of ferromagnetic states as a function of the magnetic field (solid lines).

respectively, while the resonance peak discerned at 27 GHz immediately disappears. The absorption peak observed at 27 GHz, indicated as the purple dots in Fig. 2(c), is therefore the resonance mode characteristic of the cycloidal spin structure. With further increasing of the field, the lower-lying mode slightly shifts towards the higher-frequency region and the higher-lying mode splits into two modes. In domains B and C the cycloidal phase is suppressed and the field-polarized ferromagnetic state is reached by fields below ~ 100 mT. In contrast, in domain A the transverse conical state formed in finite fields is robust to ~ 500 mT, around which a spin-flop transition occurs to the ferromagnetic state. In the following we focus on the magnetic resonances and their DD in the ferromagnetic state, where a simple phenomenological model can be used to describe the observed DD.

In general, the resonance frequency of the ferromagnetic state can be calculated on the basis of the Smith-Suhl formula [34], $(\omega/\gamma)^2 = [1/(M^2\sin^2\theta)]\{[(\partial^2 E)/(\partial\theta^2)] [(\partial^2 E)/(\partial\phi^2)] - [(\partial^2 E)/(\partial\theta\partial\phi)]^2\}$, where γ, M, and E are the gyromagnetic ratio, magnetization, and free energy, respectively. θ and ϕ are the polar angle and azimuthal angle of the M, respectively, in a spherical coordinate system [Fig. 2(b), inset]. In the present system, the free energy is given by

$$E = -M \cdot H - K(Mz/M)^2, \qquad (1)$$

where K and z represent the uniaxial anisotropy and unit vector along the [111] axis, respectively. The resonance frequency is thus calculated as

$$\frac{\omega}{\gamma} = \sqrt{H^2\sin^2\theta_H + \frac{H^2\cos\theta\sin 2\theta_H}{2\sin\theta} + \frac{2KH\sin\theta_H\cos 2\theta}{M\sin\theta}}, \qquad (2)$$

where θ_H is the polar angle of the H from the z axis. On the basis of Eq. (1), θ can be numerically calculated for each domain so as to minimize E, as shown in Fig. 2(b). By substituting this result into Eq. (2), it is found that the theory well reproduces the field dependence of the resonances in the ferromagnetic state with $K/M = 260$ mT [Fig. 2(c)]. Here we used $g = 1.82$ estimated in Ref. [33] and the deduced K/M value almost coincides with the value determined in Ref. [33]. Thus, the low-, intermediate- and high-frequency modes observed at 700 mT are attributed to resonance modes in the domain A, B, and C, respectively. Note that the discrepancy between the observed and calculated results for the domain A in the lower field region is due to the subsisting transverse conical state up to 500 mT, not the ferromagnetic state assumed in the calculation.

Figures 3(a) and 3(b) show the absorption spectra when the microwave propagates along $+k^\omega$ and $-k^\omega$ directions at the field of ± 700 mT, where the magnetic state is ferromagnetic for all the domains. We found a clear signature of the DD for the intermediate- and high-frequency modes: The resonance peaks are higher in the ΔS_{21} spectrum than that in the ΔS_{12} spectrum at $+700$ mT for both modes. This relationship is reversed in -700 mT, further verifying the nonreciprocal nature of the transmission. The DD spectra change systematically by applying magnetic fields, as shown in Fig. 3(c). The magnitude of the DD, which is defined as $\{(\Delta S_{12} - \Delta S_{21})/[(\Delta S_{12} + \Delta S_{21})/2]\}$, increases monotonically for the higher-lying mode and is basically unchanged for the intermediate mode as increasing the fields [Fig. 3(d)]. The magnitude of the DD amounts to approximately 20% at $+750$ mT, which is the largest value among those reported for multiferroics in the microwave frequency range to the best of our knowledge [18–21].

Notably, although k^ω, P, and M are perpendicular to each other in the domain A satisfying the necessary condition to observe the DD [6–10,12,16–18,21], the DD in the domain A is negligibly small. On the other hand, the domain C exhibits a strong DD, although for this type of domain P and M are nearly antiparallel; i.e., toroidal moment $T = P \times M$ is smaller as compared to the domain A [Figs. 4(b) and 4(c)]. Moreover, domains A and C are related to each other by rotating 70.5° around the $k^\omega\|[1\bar{1}0]$. These facts highlight the crucial role of the direction of the M in the ME coupling. It should be noted that, although the Damon-Eshbach mode or magnetostatic surface wave (at nonzero wave number) may show nonreciprocal propagation, it
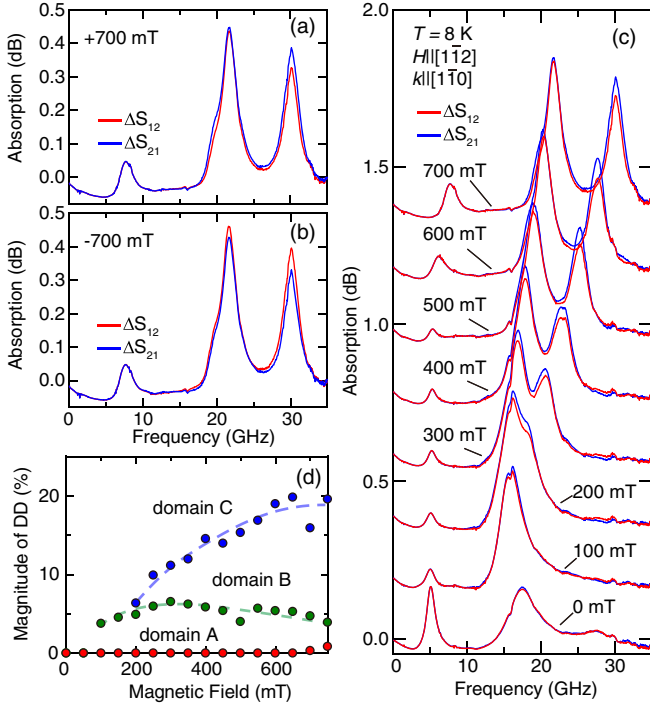
FIG. 3. The magnetic resonance spectra for $+\boldsymbol{k}^{\omega}$ and $-\boldsymbol{k}^{\omega}$ directions at $+700$ (a) and -700 mT (b). (c) The $+\boldsymbol{k}^{\omega}$ and $-\boldsymbol{k}^{\omega}$ spectra for each magnetic field. The spectra are shifted vertically for clarity. (d) The magnetic-field dependence of the magnitude of the directional dichroism. The red, green, and blue circles correspond to the lower-, intermediate-, and higher-frequency modes, respectively.
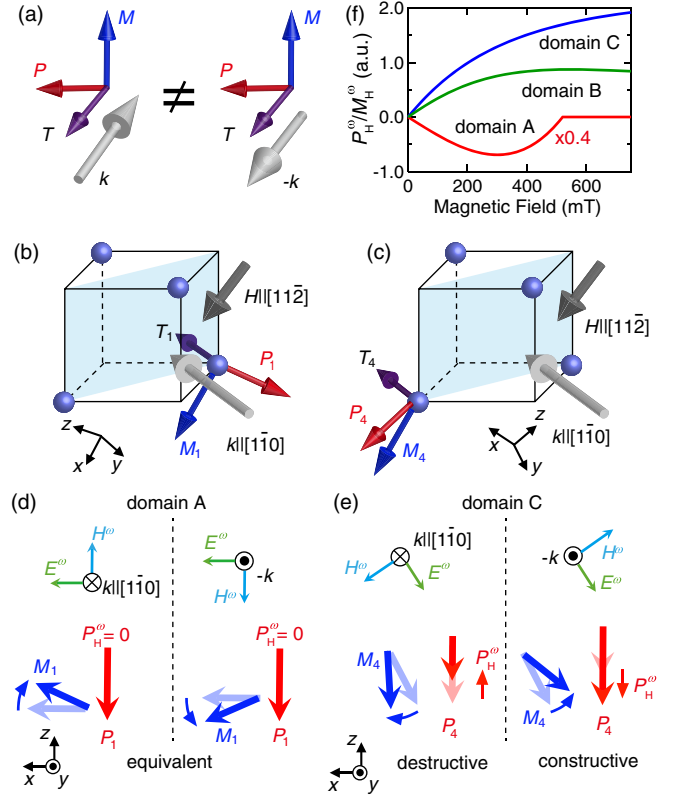

FIG. 4. (a) Schematic illustration of the DD and toroidal moment. [(b) and (c)] The experimental configurations for domain A (b) and domain C (c). [(d) and (e)] Instantaneous response of $\boldsymbol{P}$ and $\boldsymbol{M}$ for the P_1 domain (d) and P_4 domain (e). (f) The magnetic-field dependence of the dynamical polarization induced by the microwave magnetic field in domains A, B, and C.

does not lead to nonreciprocal absorption [34]. Moreover, the Damon-Eshbach mode cannot be clearly discerned in the present experiment, thus allowing us to exclude its contribution to the presently observed DD.

To understand the role of the ME coupling, we consider the instantaneous responses induced by $\boldsymbol{H}^{\omega}$ and $\boldsymbol{E}^{\omega}$ in the same manner as adopted in Ref. [19]. From the viewpoint of the symmetry, the $\boldsymbol{P}$ can be described by

$$P_x = aM_xM_z - bM_xM_y,$$
$$P_y = -bM_x^2 + bM_y^2 + aM_yM_z,$$
$$P_z = cM_z^2 + dM^2, \quad (3)$$

where x, y, and z represent $[11\bar{2}]$, $[\bar{1}10]$, and $[111]$ axes, respectively. The terms that contain H such as $P_x = a'H_xH_z$ are also allowed and have the same form as M but we omit them here for clarity. Given that the $+\boldsymbol{k}^{\omega}(\|-y)$ microwave in the CPW contains $\boldsymbol{H}^{\omega}\|z$ and $\boldsymbol{E}^{\omega}\|x$ [Fig. 1(g)] and that the $\boldsymbol{M}$ points to the x direction in the domain A with P_1, the magnetic resonance is induced by $\boldsymbol{H}^{\omega}\|z$ and produces the dynamical magnetization, $\boldsymbol{M}_H^{\omega}$, along the z direction [Fig. 4(d)]. $\boldsymbol{M}_H^{\omega}$ simultaneously induces dynamical polarization, $\boldsymbol{P}_H^{\omega}$, along the x direction through the ME coupling given by Eq. (3);

$P_{H,x}^{\omega} = aM_xM_{H,z}^{\omega}$. This $P_{H,x}^{\omega}$ may interfere with the dynamical polarization ($\boldsymbol{P}_E^{\omega}$) induced by $\boldsymbol{E}^{\omega}(\|x)$ and lead to the DD. In the present case, however, the DD is experimentally not observed in the domain A and therein the coefficient a should be negligibly small.

In the domain C with P_4, the $\boldsymbol{M}$ is slanted from both x and z axes and therefore $\boldsymbol{M}$ and $\boldsymbol{M}_H^{\omega}$ have both finite x and z components [Fig. 4(e)]. As the result, $\boldsymbol{P}_H^{\omega}$ is generated along the z direction, which can be described as $2cM_zM_{H,z}^{\omega}$. The resulting $P_{H,z}^{\omega}$ is antiparallel to $P_{E,z}^{\omega}$ for the $+\boldsymbol{k}^{\omega}$ microwave whereas they are parallel for the $-\boldsymbol{k}^{\omega}$ microwave, as schematically illustrated in Fig. 4(e); here, we note that $\boldsymbol{H}^{\omega}$ and $\boldsymbol{E}^{\omega}$ also have both x and z components. Therefore, the interference between $P_{H,z}^{\omega}$ and $P_{E,z}^{\omega}$ occurs destructively for the $+\boldsymbol{k}^{\omega}$ microwave and constructively for the $-\boldsymbol{k}^{\omega}$ microwave. This difference may lead to the DD, which is indeed observed in Fig. 3. Hence, the DD observed here originates from the form of the ME coupling, $P_z = cM_z^2$, which is in accord with the nature of the static ME effect reported in Ref. [24]. The microscopic origin of the ME coupling is therefore attributed either to the anisotropic exchange mechanism or to the single-site ME effect.

The ME coupling discussed above reproduces the magnetic-field dependence of the DD in the ferromagnetic state as well: Because the H^ω circulates around the signal line and is perpendicular to the [110] axis in the present experimental setup, we calculated $P^\omega_{H,z} = 2cM_zM^\omega_{H,z}$, which is proportional to the magnitude of the DD, for each domain by averaging all the contributions induced by H^ω lying in the (110) plane. The calculation qualitatively reproduces the experiment except for the low-field region for domain A, where the transverse conical state, not the ferromagnetic state presumed in the calculation, is formed: The DD for domain A is negligibly small, for domain B it shows a broad maximum, and for domain C it keeps increasing with the magnetic field [see Figs. 3(d) and 4(f)].

It has been believed that the DD should be maximized when k^ω, P, and M are perpendicular to each other, whereas the present observation unambiguously demonstrates the crucial role of the microscopic ME coupling as well as the macroscopic symmetry: The macroscopic symmetry defines the necessary conditions for the existence of the DD, as illustrated in Fig. 4(a), but leaves freedom for different ME mechanisms to govern the magnitude of the DD; as is clear from the comparison of the magnetization and polarization dynamics for domain A and C, the DD can emerge even when k^ω, P, and M are not totally perpendicular to each other.

In summary, we have investigated the microwave DD in the polar ferromagnetic state of GaV$_4$S$_8$. The ferromagnetic resonance is separated for the structural domains with different directions of the electric polarization due to uniaxial anisotropy and we clearly observed the DD as large as 20% for a specific domain without cancellation among the multidomain states. Our findings widen the class of materials that can potentially show the large DD even in the presence of the multidomain states.

[1] T. Kimura, T. Goto, H. Shintani, K. Ishizaka, T. Arima, and Y. Tokura, Nature (London) **426**, 55 (2003).

[2] W. Eerenstein, N. D. Mathur, and J. F. Scott, Nature (London) **442**, 759 (2006).

[3] S.-W. Cheong and M. Mostovoy, Nat. Mater. **6**, 13 (2007).

[4] Y. Tokura, S. Seki, and N. Nagaosa, Rep. Prog. Phys. **77**, 076501 (2014).

[5] R. V. Pisarev, B. B. Krichevtsov, and V. V. Pavlov, Phase Transitions **37**, 63 (1991).

[6] G. L. Rikken and E. Raupach, Nature (London) **390**, 493 (1997).

[7] M. Kubota, T. Arima, Y. Kaneko, J. P. He, X. Z. Yu, and Y. Tokura, Phys. Rev. Lett. **92**, 137401 (2004).

[8] M. Saito, K. Taniguchi, and T. Arima, J. Phys. Soc. Jpn. **77**, 013705 (2008).

[9] I. Kézsmárki, N. Kida, H. Murakawa, S. Bordács, Y. Onose, and Y. Tokura, Phys. Rev. Lett. **106**, 057403 (2011).

[10] Y. Takahashi, R. Shimano, Y. Kaneko, H. Murakawa, and Y. Tokura, Nat. Phys. **8**, 121 (2012).

[11] S. Bordács, I. Kézsmárki, D. Szallar, L. Demkó, N. Kida, H. Murakawa, Y. Onose, R. Shimano, T. Rõõm, U. Nagel, S. Miyahara, N. Furukawa, and Y. Tokura, Nat. Phys. **8**, 734 (2012).

[12] Y. Takahashi, Y. Yamasaki, and Y. Tokura, Phys. Rev. Lett. **111**, 037204 (2013).

[13] I. Kézsmárki, N. Szaller, S. Bordács, H. Murakawa, Y. Tokura, H. Engelkamp, T. Rõõm, and U. Nagel, Nat. Commun. **5**, 3203 (2014).

[14] V. Kocsis, K. Penc, T. Rõõm, U. Nagel, J. Vít, J. Romhanyi, Y. Tokunaga, Y. Taguchi, Y. Tokura, I. Kézsmárki, and S. Bordács, Phys. Rev. Lett. **121**, 057601 (2018).

[15] S. Bordács, V. Kocsis, Y. Tokunaga, U. Nagel, T. Rõõm, Y. Takahashi, Y. Taguchi, and Y. Tokura, Phys. Rev. B **92**, 214441 (2015).

[16] I. Kézsmárki, U. Nagel, S. Bordács, R. S. Fishman, J. H. Lee, Hee Taek Yi, S.-W. Cheong, and T. Rõõm, Phys. Rev. Lett. **115**, 127203 (2015).

[17] S. Yu, B. Gao, J. W. Kim, S.-W. Cheong, M. K. L. Man, J. Madéo, K. M. Dani, and D. Talbayev, Phys. Rev. Lett. **120**, 037601 (2018).

[18] Y. Okamura, F. Kagawa, M. Mochizuki, M. Kubota, S. Seki, S. Ishiwata, M. Kawasaki, Y. Onose, and Y. Tokura, Nat. Commun. **4**, 2391 (2013).

[19] Y. Okamura, F. Kagawa, S. Seki, M. Kubota, M. Kawasaki, and Y. Tokura, Phys. Rev. Lett. **114**, 197202 (2015).

[20] Y. Nii, R. Sasaki, Y. Iguchi, and Y. Onose, J. Phys. Soc. Jpn. **86**, 024707 (2017).

[21] Y. Iguchi, Y. Nii, and Y. Onose, Nat. Commun. **8**, 15252 (2017).

[22] D. Khomskii, Physics **2**, 20 (2009).

[23] I. Kézsmárki, S. Bordács, P. Milde, E. Neuber, L. M. Eng, J. S. White, H. M. Rønnow, C. D. Dewhurst, M. Mochizuki, K. Yanai, H. Nakamura, D. Ehlers, V. Tsurkan, and A. Loidl, Nat. Mater. **14**, 1116 (2015).

[24] E. Ruff, S. Widmann, P. Lunkenheimer, V. Tsurkan, I. Kézsmárki, and A. Loidl, Sci. Adv. **1**, e1500916 (2015).

[25] R. Pocha, D. Johrendt, and R. Pöttgen, Chem. Mater. **12**, 2882 (2000).

[26] A. V. Powell, A. McDowall, I. Szkoda, K. S. Knight, B. J. Kennedy, and T. Vogt, Chem. Mater. **19**, 5035 (2007).

[27] Á. Butykai, S. Bordács, I. Kézsmárki, V. Tsurkan, A. Loidl, J. Döring, E. Neuber, P. Milde. S. C. Kehr, and L. M. Eng, Sci. Rep. **7**, 44663 (2017).

[28] J. S. White, Á. Butykai, R. Cubitt, D. Honecker, C. D. Dewhurst, L. F. Kiss, V. Tsurkan, and S. Bordács, Phys. Rev. B **97**, 020401 (2018).

[29] See Supplemental Material at http://link.aps.org/supplemental/10.1103/PhysRevLett.122.057202 for details of data and the analysis procedure.

[30] D. Szaller, S. Bordács, and I. Kézsmárki, Phys. Rev. B **87**, 014421 (2013).

[31] N. A. Spaldin, M. Fiebig, and M. Mostovoy, J. Phys. Condens. Matter **20**, 434203 (2008).

[32] D. Ehlers, I. Stasinopoulos, V. Tsurkan, H.-A. Krug von Nidda, T. Fehér, A. Leonov, I. Kézsmárki, D. Grundler, and A. Loidl, Phys. Rev. B **94**, 014406 (2016).

[33] D. Ehlers, I. Stasinopoulos, I. Kézsmárki, T. Fehér, V. Tsurkan, H.-A. Krug von Nidda, D. Grundler, and A. Loidl, J. Phys. Condens. Matter **29**, 065803 (2017).

[34] A. G. Gurevich and G. A. Melkov, *Magnetization Oscillations and Waves* (CRC Press, Boca Raton, 1996).