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Article

Unexpected Anisotropy of the Electron and Hole Landé g-Factors in Perovskite $\text{CH}_3\text{NH}_3\text{PbI}_3$ Polycrystalline Films

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Abstract: In this work, we studied, at low temperature, the coherent evolution of the localized electron and hole spins in a polycrystalline film of $\text{CH}_3\text{NH}_3\text{PbI}_3$ (MAPI) by using a picosecond-photo-induced Faraday rotation technique in an oblique magnetic field. We observed an unexpected anisotropy for the electron and hole spin. We determined the electron and hole Landé factors when the magnetic field was applied in the plane of the film and perpendicular to the exciting light, denoted as transverse \perp factors, and when the magnetic field was applied perpendicular to the film and parallel to the exciting light, denoted as parallel \parallel factors. We obtained $|g_{e,\perp}| = 2.600 \pm 0.004$, $|g_{e,\parallel}| = 1.604 \pm 0.033$ for the electron and $|g_{h,\perp}| = 0.406 \pm 0.002$, $|g_{h,\parallel}| = 0.299 \pm 0.007$ for the hole. Possible origins of this anisotropy are discussed herein.

Keywords: halide perovskite; $\text{CH}_3\text{NH}_3\text{PbI}_3$; g-factor; polycrystalline film; photo-induced Faraday rotation



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1. Introduction

In the last few years, lead halide perovskites have attracted great interest in the scientific community. This class of materials exhibits many excellent photo-physical properties, such as broad absorption spectra, tunable band gap emission, narrow line-width, high luminescence yields, large charge carrier mobilities, and dielectric constants. They are very useful for their potential applications in photovoltaic, opto-electronic, and electronic devices, such as light-emitting diodes or lasers, photodetectors, photo-sensors, and transistors [1–9]. In addition, these materials are also promising for opto-spintronic devices [10,11] because they show long spin lifetimes [12–15], chiral properties [16–18], and large Rashba splitting [19]. The synthesis of engineered nanostructures [20–22] extends the future applications of these materials to the domain of quantum optics and quantum information processing.

Until now, two main kinds of experiments have been performed to obtain information about g-factors in hybrid perovskite materials. The first includes magneto-luminescence and magneto-absorption experiments, which give access to an effective exciton g-factor that contains electron and hole contributions according to the relation $g_x = g_e + g_h$ [23–28]. The second kind of experiment providing information on Landé factors includes photo-induced Kerr and Faraday rotations [29], both using a pump-probe configuration in reflectivity or transmission mode, respectively. Recently, by using these experimental techniques with a

magnetic field applied in Voigt configuration (the magnetic field direction is perpendicular to the pump beam propagation direction), the g_{\perp} for electrons and holes have been determined in bulk CsPbBr_3 [14], $\text{FA}_{0.9}\text{CsPbI}_{2.8}\text{Br}_{0.2}$ [30], and in $\text{CH}_3\text{NH}_3\text{PbI}_3$ polycrystalline films [13,15].

In this article, we study the coherent evolution of the electronic spin in an oblique magnetic field by using a picosecond-photo-induced Faraday rotation (PFR) technique. We measure an unexpected anisotropy of the electron and hole Landé factors in a polycrystalline film of $\text{CH}_3\text{NH}_3\text{PbI}_3$ (MAPI) and we discuss the possible origins of the observed anisotropy.

2. Materials and Methods

The samples studied here are polycrystalline films of $\text{CH}_3\text{NH}_3\text{PbI}_3$ prepared from a solution of methylammonium iodide (1.5 mmol, 238.5 mg) and lead iodide (1.5 mmol, 691.5 mg) in γ -butyrolactone (2 mL). The new quartz substrate was cleaned in a solution of ethanol in an ultrasonic bath for 10 min and then treated with O_2 plasma for 20 min. In a glove box, 50 μL of MAPI solution was deposited on the substrate by spin coating at 2000 rpm for 30 s. Then, the sample was annealed at 120 $^{\circ}\text{C}$ for 1 min [31]. The thickness of the resulting film was, on average, equal to 170 nm and exhibited a rough surface.

Film morphology was studied with a Zeiss Merlin VP Compact field emission scanning electron microscope (SEM, Zeiss Jena, Germany). The polycrystalline film constituted of grains presenting polydispersity in their sizes, from several tens of nanometers up to 400 nm, as shown in Figure 1.

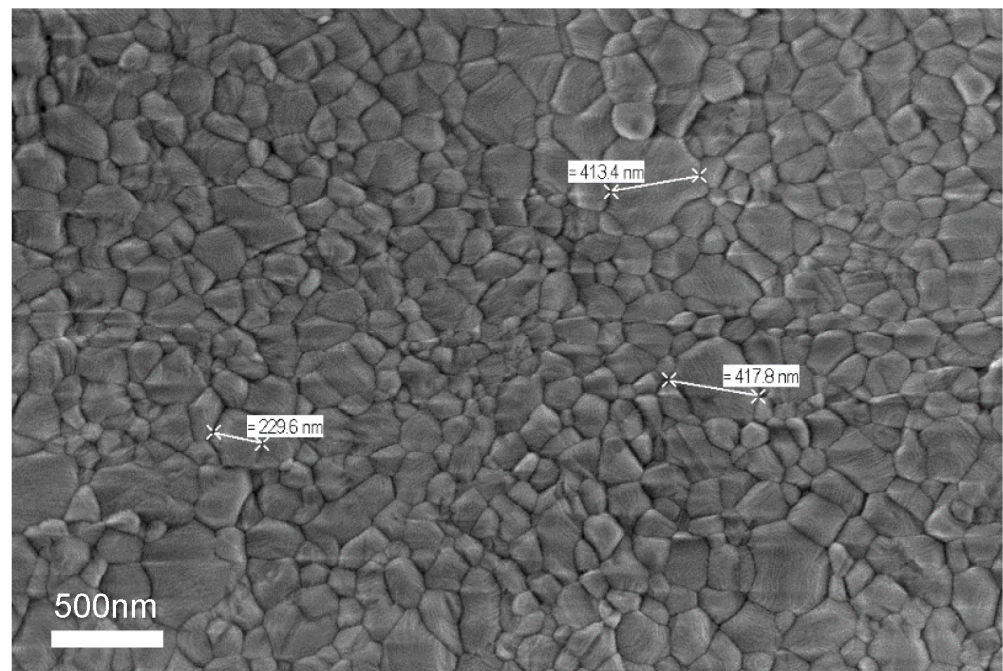


Figure 1. SEM image of MAPI polycrystalline films used in this study.

The absorption spectrum of the MAPI sample at 11 K is shown in the inset of Figure 2a. The evolution in temperature of the lower absorption peak is consistent with the orthorhombic-tetragonal transition shown by other authors [15,31].

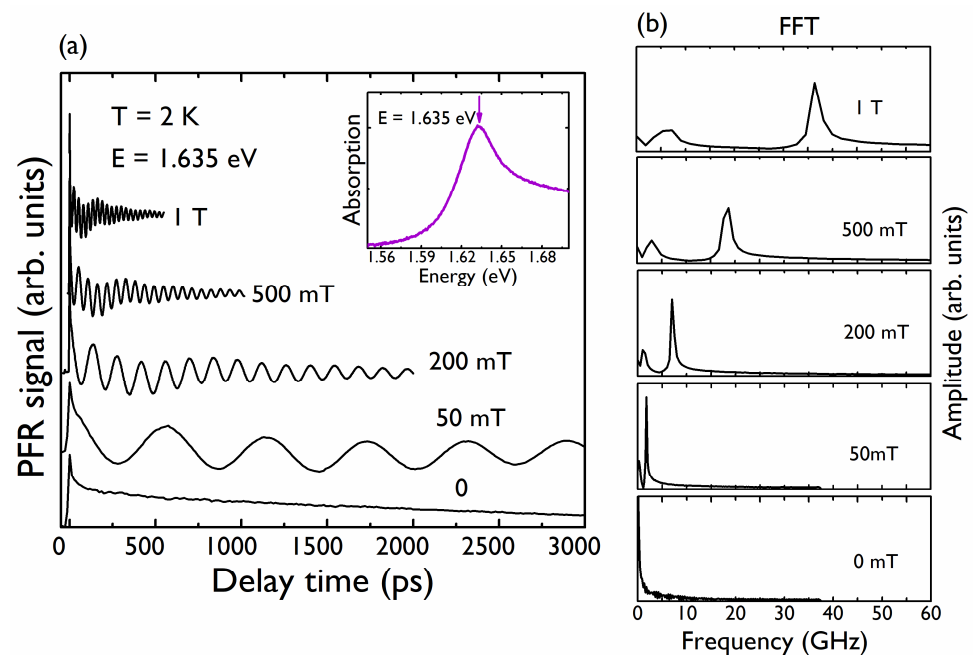


Figure 2. (a) PFR signals at 2 K for different magnetic fields. The curves have been shifted for clarity. The magnetic field is perpendicular to the propagation direction of the pump and probe beams ($\alpha = 0^\circ$). Inset: Absorption spectrum on MAPI sample obtained at 11 K. The arrow denotes the energy at which the PFR measurements were performed; (b) FFT of the PFR signals appearing in (a).

To perform photo-induced Faraday rotation (PFR) experiments, a picosecond Ti:Sapphire laser (76 MHz repetition rate) was split into pump and probe beams. The mean optical powers were 500 μW for the pump beam and 50 μW for the probe beam; the 3-ps duration of the laser pulses allowed a spectral resolution of 1 meV. The pump beam polarization was $\sigma + / \sigma -$ modulated at 500 kHz with an electro-optic modulator in order to avoid nuclear spin polarization. The probe beam was linearly polarized, and its intensity was modulated with an acousto-optic modulator at 3 kHz. After transmission through the sample, the probe beam was spatially resolved into its vertical and horizontal components, and the difference in intensity was measured in a balanced optical bridge. To improve the signal-to-noise ratio, a double lock-in amplifier was used for the measurements; the data presented in this work correspond to one single scan of the pump–probe delay, and thanks to the high signal-to-noise ratio (~ 100), no averaging of several scans is needed. The sample was placed in a cryostat containing superconducting coils. The spot size of the pump (probe) beam on the sample had a diameter of about 30 μm (15 μm), and several crystalline grains (sizes evidenced in Figure 1) on the films were thereby addressed.

3. Results

Figure 2a shows the PFR signal obtained at 2 K; the pump–probe energy is tuned to the maximum of the lowest absorption band of MAPI, centered at 1.635 eV. A magnetic field B , was applied in the Voigt configuration, i.e., perpendicular to the pump and probe propagation directions. The PFR signal contained two oscillatory contributions, which were revealed by the Fast Fourier Transform (FFT), as shown in Figure 2b. Recently, the controversy regarding the identification of these two signals has been resolved in favor of the localization of electrons and holes in $\text{CH}_3\text{NH}_3\text{PbI}_3$ at spatially separated locations [15]. This interpretation has also been adopted and demonstrated in other halide perovskite bulk materials [14,15,30]. At zero magnetic field, the PFR curves can be described by the sum of two exponential contributions. The initial fast decay (\approx tens of ps) is likely related to the spin-polarized population of excitons bound to donors and/or acceptors and neglected in the fitting procedure (Figure 2a). The long decay of the PFR signal is related to electrons and/or holes bound to donors and acceptors. In the presence of an applied magnetic field

and with a weak excitation density, the PFR signal can be well described by the sum of two damped oscillations associated with localized electrons and holes, respectively:

$$\theta_F = A_e e^{-t/T_{2,e}^*} \cos(2\pi t f_e) + A_h e^{-t/T_{2,h}^*} \cos(2\pi t f_h), \tag{1}$$

where f_e, f_h are the electron and hole frequencies, respectively, identified in the FFT, and $T_{2,e}^*, T_{2,h}^*$ are the corresponding dephasing times at the given magnetic field. $T_{2,e}^*, T_{2,h}^*$ shorten with an increasing magnetic field due to the inhomogeneities of g values [29].

We fit Equation (1) to all the PFR curves. The inset in Figure 3 shows the dependence of the two extracted frequencies $\omega_i = 2\pi f_i$ ($i = e, h$) on the applied magnetic field B . The two frequencies increase linearly with the magnetic field, following the expression:

$$\omega_i = \frac{|g_{i,\perp}| \mu_B}{\hbar} B, \tag{2}$$

where μ_B is the Bohr magneton, \hbar the reduced Planck constant, $g_{i,\perp}$ is the Landé factor of the corresponding species, and \perp means that the magnetic field is applied in the plane of the film and perpendicular to the exciting light. We underline that in the full magnetic range, even for the smallest applied magnetic fields, the electron and hole Zeeman splittings show a linear behavior with the field. We conclude then that the Rashba effect is negligible in our case. From the slopes of the two linear fits, we obtained two Landé factors $|g_{e,\perp}| = 2.600 \pm 0.004$ and $|g_{h,\perp}| = 0.406 \pm 0.002$. We did not detect any difference in the measured Landé factors when the magnetic field was applied along two perpendicular directions in the plane of the film. These values are very close to the values $|g_{e,\perp}| = 2.63$ and $|g_{h,\perp}| = 0.33$ obtained in MAPI films in a recent work [13].

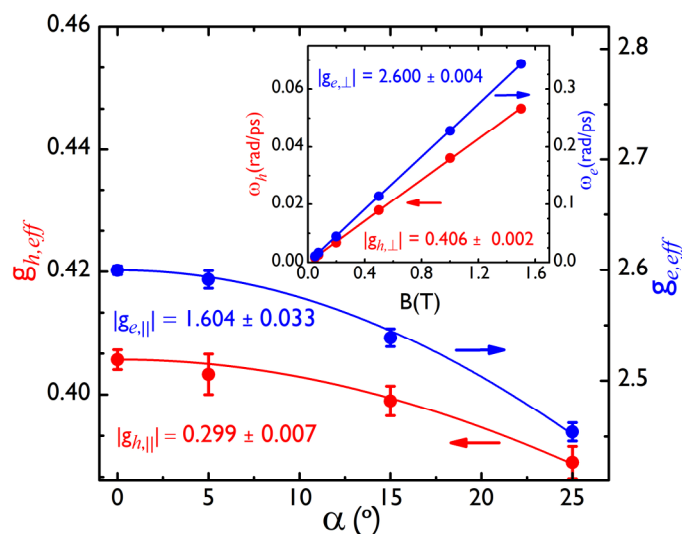


Figure 3. Effective Landé factors for the hole (left, red color) and the electron (right, blue color) versus the angle α . The points represent the experimental data, and the solid lines are fits to Equation (3). Inset: Larmor precession frequencies of the hole (left, red line) and the electron (right, blue line) versus the magnetic field, when the sample is perpendicular to pump and probe beams ($\alpha = 0^\circ$) with an in-plane magnetic field.

To obtain a configuration for which an oblique magnetic field was applied to the sample, the sample was rotated around the axis perpendicular to the plane defined by the applied magnetic field and the pump–probe propagation direction. The rotation angle α was measured between \vec{B} and the MAPI film plane. β is the angle (when $\alpha \neq 0$) between the normal direction of the MAPI film and the direction of the oriented electronic spin, given by \vec{S} as seen in the inset of Figure 4. For an anisotropic g tensor, the spin precession

vector $\vec{\Omega}$ is generally non-collinear with \vec{B} . The vector $\vec{\Omega}$ is tilted away from the direction of \vec{B} by the angle γ given by $g_{\perp} \tan \gamma = g_{\parallel} \tan \alpha$ with g_{\parallel} the Landé factor for a magnetic field applied parallel to the exciting light and perpendicular to the film. Figure 4 shows PFR measurements at $B = 500$ mT for $\alpha = 0^{\circ}, 5^{\circ}, 15^{\circ}, 25^{\circ}$. The period of the two contributions to each PFR signal increases as α increases. In Figure 4, this is especially clear for the high-frequency contribution (electron contribution), as shown by the red dashed line. Figure 5a shows the FFT of PFR signals shown in Figure 4. Due to the smaller anisotropy of the hole Landé factor, in comparison with the one found for the electron, it is more difficult to appreciate the decrease in the frequency (increase of the period) for the hole component in the PFR signal at only one value of the magnetic field; that is why we show, in Figure 5b, the FFT of PFR signals for different values of α at 1.5 T.

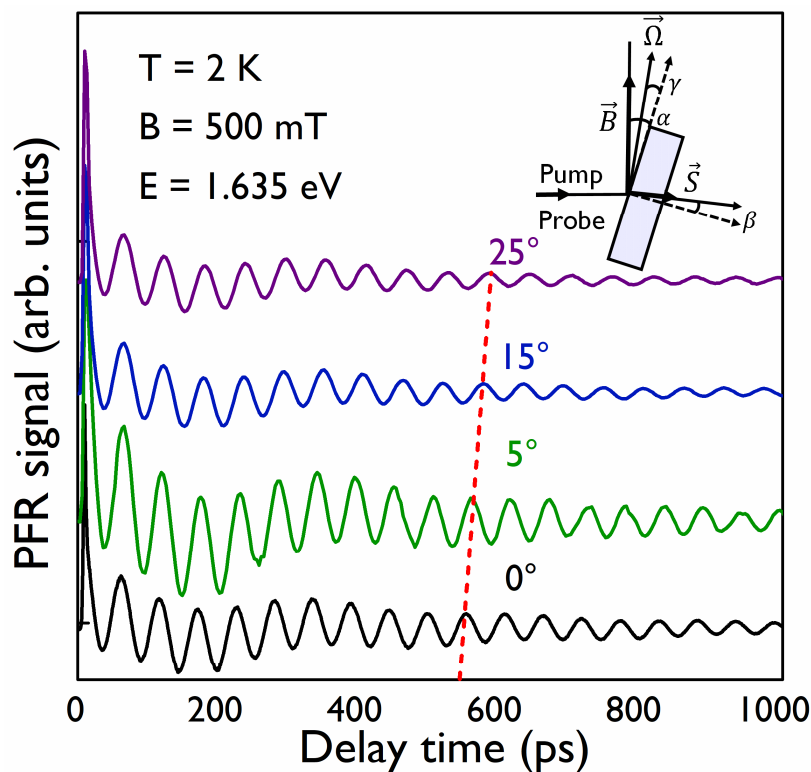


Figure 4. PFR signals obtained at 2 K and at $B = 500$ mT for different values of α angle. The red dashed line follows the 10-th maximum peak of the high-frequency component, showing the increase of the period as the angle α increases.

The effective $g_{i,eff}$ ($i = \text{electron or hole}$) factors for a tilted magnetic field, when $\alpha \neq 0$, are given by:

$$|g_{i,eff}| = \sqrt{g_{i,\perp}^2 \cos^2 \alpha + g_{i,\parallel}^2 \sin^2 \alpha} \tag{3}$$

We determined $|g_{i,eff}|$ at each explored value of α , thanks to a linear fit of the oscillation frequency versus the magnetic field, as we have done for $|g_{i,\perp}|$. The values found for the effective Landé factor for the electron and hole versus the tilted angle, α , are plotted in Figure 3. We finally obtain, through a fit to Equation (3), $|g_{h,\parallel}| = 0.299 \pm 0.007$ for the hole parallel g-factor and $|g_{e,\parallel}| = 1.604 \pm 0.033$ for the parallel electron Landé factor. Very comparable results were observed on two different samples and on different points of each sample, making the results very reproducible.

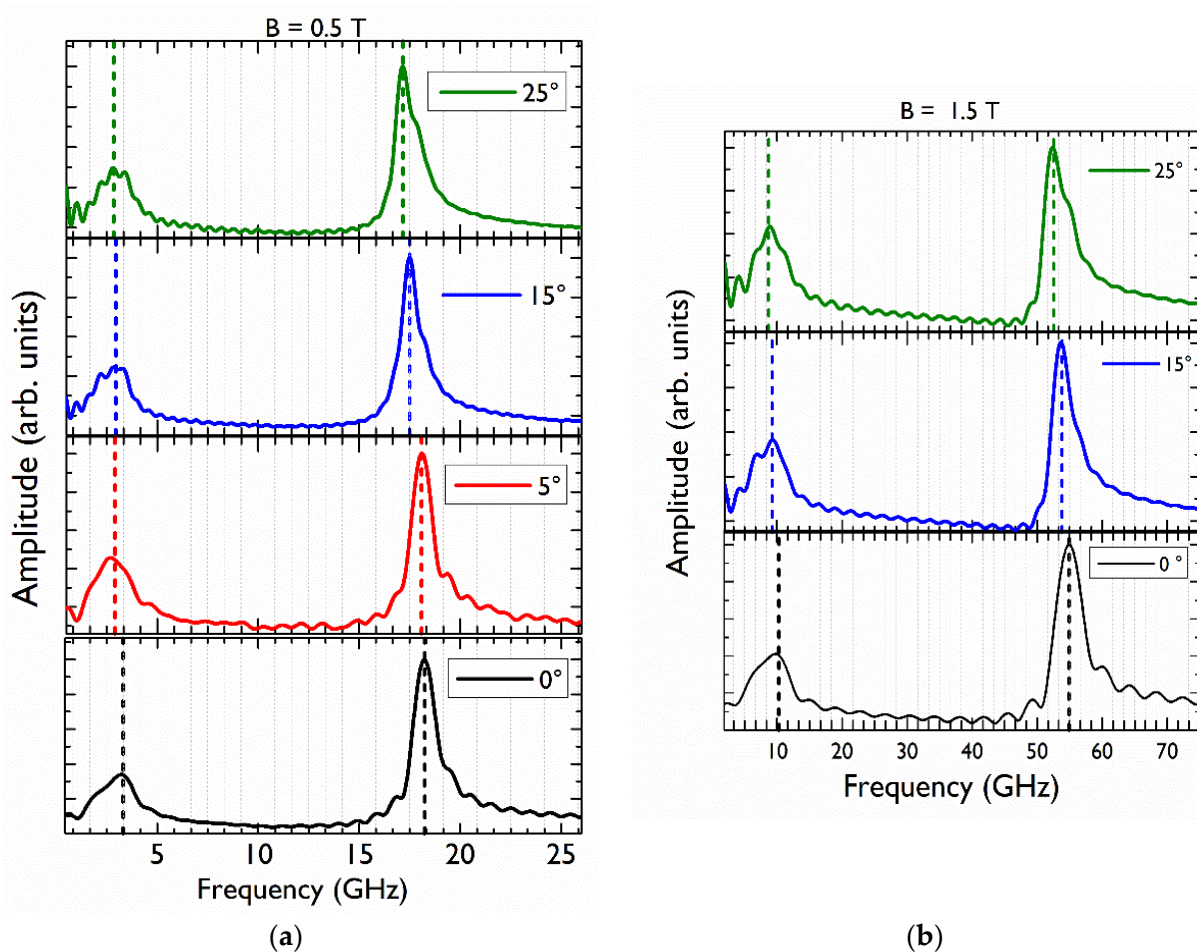


Figure 5. (a) FFT of the PFR signals appearing in Figure 4 obtained at a fixed magnetic field $B = 0.5$ T and different angles 0° (black), 5° (red), 15° (blue), 25° (green). The curves have been vertically shifted for clarity. The vertical dashed lines indicate the two identified frequencies corresponding to the hole and electron in each case. (b) FFT of the PFR signals obtained at $B = 1.5$ T. The decrease in the frequencies as the angle increases is indicated with the dashed lines.

4. Discussion

The method used to obtain the experimental values for the Landé factors does not allow us to give the sign of the g -factors, but $\mathbf{k}\cdot\mathbf{p}$ calculations [32] and recent experimental results [33] allow us to attribute a negative sign to the hole Landé factor and a positive sign to the electron factor.

The anisotropy ratio, defined as $\frac{|g_{i,\perp} - g_{i,\parallel}|}{|g_{i,\perp} + g_{i,\parallel}|}$ and measured here, is equal to 24% for the electron and 15% for the hole. This anisotropy was fully unexpected for a polycrystalline film if all the c axis of the crystals were randomly oriented.

There are very few studies reporting experimental g -factor values for electrons and holes in halide perovskite materials [13–15,30,33]. Very recently [33], experimental results on MAPI single crystals were obtained showing a strong 43% anisotropy for the hole g -factor ($-0.71 \leq g_h \leq -0.28$) and a smaller 10% anisotropy for the electron g -factor $2.46 \leq g_e \leq 2.98$. However, using these results as an experimental reference for MAPI single crystals is not straightforward, as the authors underlined that the main axes of the g -factor tensor are tilted with respect to cubic axes; they gave different explanations for this experimental evidence, including the possibility of a transition to a monoclinic phase.

The already cited $\mathbf{k}\cdot\mathbf{p}$ calculations [32] determined anisotropic hole and electron Landé factors for the MAPI tetragonal crystal phase as follows: $g_{h,\parallel} = -0.472$, $g_{h,\perp} = -0.354$, and $g_{e,\parallel} = 1.67$, $g_{e,\perp} = 2.281$. In this case, the anisotropy is a consequence of the anisotropic

crystal phase, and \parallel (\perp) means that the magnetic field is parallel (perpendicular) to the crystal c-axis of the tetragonal phase. This calculated anisotropy is very close to (smaller than) the experimentally observed hole (electron) g-factor anisotropy in the MAPI polycrystalline sample studied here, but the origin of the latter anisotropy in our samples has not yet been found. Among the possible causes of this anisotropy, we cite the non-random distribution of the crystallographic axes of MAPI crystals in the film or the residual strains.

A first explanation may be found in a preferential orientation of the polycrystals. Indeed, T. M. Brenner et al. [34] evidenced the preferential growth and crystallographic orientation of MAPI grains obtained from PbI_2 crystal-precursor by using a “two-step” synthesis method. L. Oesinghaus et al. [35] also demonstrated a preferential orientation of MAPI crystals in a polycrystalline film by using wide-angle X-ray scattering. They found a weaker preferential orientation in films obtained by “one-step” methods than in films resulting from the use of a “two-step” synthesis process.

On the other hand, recent studies of polycrystalline films of halide perovskites using X-ray diffraction techniques [36] evidenced the presence of lattice strains in these films due to the thermal expansion mismatch between the perovskite material and the substrate for a large variety of substrates. Cheng Zhu et al. [37], by using depth-dependent grazing incident X-ray diffraction measurements, also showed the existence of a gradient distribution of the in-plane strain component perpendicular to the substrate of halide perovskite films. Beyond the effect of these residual strains on the stability of perovskite films under illumination or on the carrier dynamics, they may also be at the origin of an anisotropy of the magnetic properties via the anisotropy of g-factors.

In conclusion, our PFR experimental study of the electron and hole spin dynamics in polycrystalline MAPI films, performed at 2 K, made it possible to measure the absolute values of the corresponding Landé factors in the orthorhombic phase. By controlling the tilt angle of the sample in the direction of the magnetic field, we observed a sharp and intriguing anisotropy of these Landé factors: 24% and 15% for the electron and the hole, respectively. At present, the origin of this anisotropy has not been identified. Further and methodical studies on X-ray diffraction must be done to clearly establish a correlation between the morphology of the polycrystalline films and the anisotropy of g-factors.

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References

1. Stoumpos, C.C.; Malliakas, C.D.; Peters, J.A.; Liu, Z.; Sebastian, M.; Im, J.; Chasapis, T.C.; Wibowo, A.C.; Chung, D.Y.; Freeman, A.J.; et al. Crystal Growth of the perovskite semiconductor CsPbBr_3 : A material for high energy radiation detection. *Cryst. Growth Des.* **2013**, *13*, 2722–2727. [[CrossRef](#)]
2. Lee, M.M.; Teuscher, J.; Miyasaka, T.; Murakami, T.N.; Snaith, H.J. Efficient hybrid solar cells based on mesosuperstructured organometal halide perovskites. *Science* **2012**, *338*, 643–647. [[CrossRef](#)]

3. Manser, J.S.; Christians, J.A.; Kamat, P.V. Intriguing optoelectronic properties of metal halide perovskites. *Chem. Rev.* **2016**, *116*, 12956–13008. [[CrossRef](#)] [[PubMed](#)]
4. Li, G.; Rocca Rivarola, F.W.; Davis, N.J.J.L.; Bai, S.; Jellicoe, T.C.; de la Peña, F.; Hou, S.; Ducatin, C.; Gao, F.; Friend, R.H.; et al. Highly efficient perovskite nanocrystals light-emitting diodes enabled by a universal crosslink method. *Adv. Mater.* **2016**, *28*, 3528–3534. [[CrossRef](#)] [[PubMed](#)]
5. Cho, H.; Jeong, S.H.; Park, Y.H.; Kim, M.H.; Wolf, C.; Lee, C.L.; Heo, J.H.; Sadhanala, A.; Myoung, N.S.; Yoo, S.; et al. Overcoming the electroluminescence efficiency limitations of perovskite light-emitting diodes. *Science* **2015**, *350*, 1222–1225. [[CrossRef](#)] [[PubMed](#)]
6. Dursun, I.; Shen, C.; Parida, M.R.; Pan, J.; Sarmah, S.P.; Priante, D.; Alyami, N.; Liu, J.; Saidaminov, M.I.; Alias, M.S.; et al. Perovskite nanocrystals as a color converter for visible light communication. *ACS Photonics* **2016**, *3*, 1150–1156. [[CrossRef](#)]
7. Jana, S.; Carlos, E.; Panigrahi, S.; Martins, R.; Fortunato, E. Toward stable solution-processed high-mobility p-type thin film transistors based on halide perovskites. *ACS Nano* **2020**, *14*, 14790–14797. [[CrossRef](#)]
8. Panigrahi, S.; Jana, S.; Calmeiro, T.; Nunes, D.; Martins, R.; Fortunato, E. Imaging the anomalous charge distribution inside CsPbBr₃ perovskite quantum dots sensitized solar cells. *ACS Nano* **2017**, *11*, 10214–10221. [[CrossRef](#)]
9. Panigrahi, S.; Jana, S.; Calmeiro, T.; Nunes, D.; Deuermeier, J.; Martins, R.; Fortunato, E. Mapping the space charge carrier dynamics in plasmon-based perovskite solar cells. *J. Mater. Chem.* **2019**, *7*, 19811–19819. [[CrossRef](#)]
10. Kepenekian, M.; Robles, R.; Katan, C.; Saponi, D.; Pedesseau, L.; Even, J. Rashba and Dresselhaus effects in hybrid organic-inorganic perovskites: From basics to devices. *ACS Nano* **2015**, *9*, 1157–11567. [[CrossRef](#)]
11. Wang, J.; Zhang, C.; Liu, H.; McLaughlin, R.; Zhai, Y.; Vardeny, S.R.; Liu, X.; McGill, S.; Semenov, D.; Guo, H.; et al. Spin opto-electronic devices based in hybrid organic-inorganic trihalide perovskites. *Nat. Commun.* **2019**, *10*, 129. [[CrossRef](#)]
12. Giovanni, D.; Ma, H.; Chua, J.; Grätzel, M.; Ramesh, R.; Mhaisalkar, S.; Mathews, N.; Sum, T.C. Highly spin-polarized carrier dynamics and ultralarge photoinduced magnetization in CH₃NH₃PbI₃ perovskite thin films. *Nano Lett.* **2015**, *15*, 1553–1558. [[CrossRef](#)]
13. Odenthal, P.; Talmadge, W.; Gundlach, N.; Wang, R.; Zhang, C.; Sun, D.; Yu, Z.G.; Vardeny, Z.V.; Li, Y.S. Spin-polarized carrier dynamics and ultralarge photoinduced magnetization in CH₃NH₃PbI₃ perovskite thin films. *Nature Phys* **2017**, *13*, 894–899. [[CrossRef](#)]
14. Belykh, V.V.; Yakovlev, D.R.; Glazov, M.M.; Grigoryev, P.S.; Hussain, M.; Rautert, J.; Dirin, D.N.; Kovalenko, M.V.; Bayer, M. Coherent spin dynamics of electrons and holes in CsPbBr₃ perovskite crystals. *Nat. Commun.* **2019**, *10*, 673. [[CrossRef](#)]
15. Garcia-Arellano, G.; Trippé-Allard, G.; Legrand, L.; Barisien, T.; Garrot, D.; Deleporte, E.; Bernardot, F.; Testelin, C.; Chamarro, M. Energy tuning of electronic spin coherent evolution in methylammonium lead iodide perovskites. *J. Phys. Chem. Lett.* **2021**, *12*, 8272–8279. [[CrossRef](#)]
16. Billing, D.G.; Lemmerer, A. Bis [(S)-β-phenethylammonium] tribromoplumbate (II). *Acta Cryst.* **2003**, *59*, m381–m383.
17. Ma, J.; Chen, C.; Jin, L.; Wang, J.; Wang, S.; Tang, J.; Li, D. Chiral 2D perovskites with a high degree of circularly polarized luminescence. *ACS Nano* **2019**, *13*, 3659–3665. [[CrossRef](#)]
18. Chen, C.; Gao, L.; Gao, W.; Ge, C.; Du, X.; Li, Z.; Yang, Y.; Niu, G.D.; Tang, J. Circularly polarized light detection using chiral hybrid perovskites. *Nat. Commun.* **2019**, *10*, 1927–1934. [[CrossRef](#)]
19. Niesner, D.; Wilhelm, M.; Levchuk, I.; Osvet, A.; Shrestha, S.; Batentschuk, M.; Brabec, C.; Fauster, T. Giant Rashba splitting in CH₃NH₃PbBr₃ organic-inorganic perovskites. *Phys. Rev. Lett.* **2016**, *117*, 126401. [[CrossRef](#)]
20. Bekenstein, Y.; Koscher, B.A.; Eaton, S.W.; Yang, P.; Alivisatos, A.P. Highly luminescent colloidal nanoplatelets of perovskite cesium lead halide and their oriented assemblies. *J. Am. Chem. Soc.* **2015**, *137*, 16008–16011. [[CrossRef](#)]
21. Gonzalez-Carrero, S.; Galian, R.E.; Pérez-Prieto, J. Maximizing the emissive properties of CH₃NH₃PbBr₃ perovskite nanoparticles. *J. Mat. Chem. A* **2015**, *3*, 9187–9193. [[CrossRef](#)]
22. Steinmetz, V.; Ramade, J.; Legrand, L.; Barisien, T.; Bernardot, F.; Lhuillier, E.; Bernard, M.; Vabre, M.; Saïdi, I.; Ghribi, A.; et al. Anisotropic shape of CsPbBr₃ colloidal nanocrystals: From 1D to 2D confinement effects. *Nanoscale* **2020**, *12*, 18978–18986. [[CrossRef](#)] [[PubMed](#)]
23. Miyata, A.; Mitioglu, A.; Plochocka, P.; Portugall, O.; Tse-Wei Wang, J.; Stranks, S.D.; Snaith, H.J.; Nicholas, R.J. Direct measurement of the exciton binding energy and effective masses for charge carriers in an organic-inorganic tri-halide perovskite. *Nat. Phys.* **2015**, *11*, 582–587. [[CrossRef](#)]
24. Baranowski, M.; Plochocka, P. Excitons in metal halide perovskites. *Adv. Energy Mat.* **2020**, *10*, 1903659. [[CrossRef](#)]
25. Hirasawa, M.; Ishihara, T.; Goto, T.; Uchida, K.; Miura, N. Magnetoabsorption of the lowest exciton in perovskite-type compound (CH₃NH₃)PbI₃. *Phys. B* **1994**, *201*, 427–430. [[CrossRef](#)]
26. Tanaka, K.; Takahashi, T.; Ban, T.; Kondo, T.; Uchida, K.; Noboru, M. Comparative study on the excitons in lead-halide-based perovskite-type crystals CH₃NH₃PbBr₃ CH₃NH₃PbI₃. *Solid State Comm.* **2003**, *127*, 619–623. [[CrossRef](#)]
27. Zhang, C.; Su, D.; Yu, Z.G.; Sheng, C.X.; McGill, S.; Semenov, D.; Vardeny, Z.V. Field induced spin splitting and anomalous photoluminescence circular polarization in CH₃NH₃PbI₃ films at high magnetic field. *Phys. Rev. B* **2018**, *97*, 134412. [[CrossRef](#)]
28. Yang, Z.; Surrente, A.; Galkowski, K.; Bruyant, N.; Maude, D.K.; Haghighirad, A.A.; Snaith, H.J.; Plochocka, P.; Nicholas, R.J. Unraveling the exciton binding energy and the dielectric constant in single-crystal methylammonium lead triiodide perovskite. *J. Phys. Chem. Lett.* **2017**, *8*, 1851–1855. [[CrossRef](#)]

29. Chamarro, M.; Bernardot, F.; Testelin, C. Spin decoherence and relaxation processes in zero-dimensional semiconductor nanostructures. *J. Phys. Condens. Matter* **2007**, *19*, 445007. [[CrossRef](#)]
30. Kirstein, E.; Yakovlev, D.R.; Glazov, M.M.; Evers, E.; Zhukov, E.A.; Belykh, V.V.; Kopteva, D.; Kudlacik, N.E.; Nazarenko, O.; Dirin, D.N.; et al. Lead-dominated hyperfine interaction impacting the carrier spin dynamics in halide perovskites. *Adv. Mater.* **2021**, *34*, 2105263. [[CrossRef](#)]
31. Diab, H.; Allard-Trippé, G.; Lédée, F.; Jemli, K.; Vilar, C.; Bouchez, G.; Jacques, V.L.R.; Tejeda, A.; Even, J.; Lauret, J.-S.; et al. Narrow linewidth excitonic emission in organic-inorganic lead iodide perovskite single crystals. *J. Phys. Chem. Lett.* **2016**, *7*, 5093–5100. [[CrossRef](#)]
32. Yu, Z.G. Effective mass model and magneto-optical properties in hybrid perovskites. *Sci. Rep.* **2016**, *6*, 28576. [[CrossRef](#)]
33. Kirstein, E.; Yakovlev, D.R.; Glazov, M.M.; Zhukov, E.A.; Kudlacik, D.; Kalitukha, I.V.; Sapaga, V.F.; Dimitriev, G.S.; Semina, M.A.; Nestoklon, M.O.; et al. The Lande factors of electrons and holes in lead halide perovskites: Universal dependence on the band gap. *arXiv* **2021**, arXiv:2112.15384v1.
34. Brenner, T.M.; Rakita, Y.; Orr, Y.; Klein, E.; Feldman, I.; Elbaum, M.; Cahen, D.; Hodes, G. Conversion of single crystalline PbI_2 to $\text{CH}_3\text{NH}_3\text{PbI}_3$: Structural relations and transformation dynamics. *Chem. Mater.* **2016**, *28*, 6501–6510. [[CrossRef](#)]
35. Oesinghaus, L.; Schlipf, J.; Giesbrecht, N.; Song, L.; Hu, Y.; Bein, T.; Docampo, P.; Müller-Buschbaum, P. Towards tailored film morphologies: The origin of crystal orientation of hybrid perovskite thin films. *Adv. Mater. Interfaces* **2016**, *3*, 1600403. [[CrossRef](#)]
36. Zhao, J.; Deng, Y.; Wei, H.; Zheng, X.; Yu, Z.; Shao, Y.; Shield, J.E.; Huang, J. Strained hybrid perovskite thin films and their impact on the intrinsic stability of perovskite solar cells. *Sci. Adv.* **2017**, *3*, 5616. [[CrossRef](#)]
37. Zhu, C.; Niu, X.; Fu, Y.; Li, N.; Hu, C.; Chen, Y.; He, X.; Na, G.; Liu, P.; Zai, H.; et al. Strain engineering in perovskite solar cells and its impacts on carrier dynamics. *Nat. Commun.* **2019**, *10*, 815. [[CrossRef](#)]