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## Voltage-Controlled Magnetic Anisotropy in the Heterostructures with Two-dimensional Magnetic Material

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The advent of two-dimensional (2D) magnetic materials significantly expand the scope of voltage-controlled magnetization-switching scheme as their integration in diverse magnetic tunnel junctions (MTJ) offers a highly attractive perspective for designing future magnetoelectric random access memory (MeRAM). Here, we propose the utilization of 2D ferromagnetic  $Fe<sub>2</sub>I<sub>2</sub>$  to substitute traditional magnetic thin films for assembling energyefficient MTJs combined with Ir caping layers. We find these multilayers exhibit both giant perpendicular magnetic anisotropy (PMA) and voltage controlled magnetic anisotropy (VCMA) efficiency depending strongly on the Ir thickness and epitaxial strain effect. Furthermore, the spin reorientation has also been achieved with increased Ir thickness, and the magnitude and slope of VCMA behavior are dominated by the biaxial strain. We elucidate that the underlying mechanism is the electric-field-induced modifications of the spin-orbit coupling energies of the spin-polarized Ir-*d* orbitals. These findings not only reveal new approaches controlling 2D magnetism, but also pave the way of an alternative strategy for the design of nonvolatile and ultralow power spintronics and magnetic memory storage devices.

Magnetoelectric random access memory (MeRAM) has attracted substantial research interest as a promising memory architecture enabling ultra-low power, highly scalable, nonvolatile and high-speed writing features.[1–3] In contrast to the current-driven magnetization switching scheme for the magnetic RAM (MRAM), the realization of MeRAM relies primarily on the voltage-controlled magnetic anisotropy (VCMA) mechanism which utilizes the electric field to efficiently manipulate spin orientations without Joule heating in magnetic tunnel junctions (MTJs).[4–7] Unfortunately, to date there are still two impedimental factors in optimizing the performance of MeRAM device, namely, the perpendicular magnetic anisotropy (PMA) energy and VCMA coefficient. Typically, large PMA energy ( $>$  2 erg/cm<sup>2</sup>) can assist in maintaining the information bit against thermodynamics fluctuations, while a greater VCMA coefficient  $(> 1000 \text{ fJ/Vm})$ is required to achieve low switching energy  $\left($  < 1 fJ/bit), low write voltage  $(< 1 V)$  and write error rate.[8, 9] In the pursuit of better energy efficiency, various ferromagnetic-insulator interfaces with high tunnel magnetoresistance ratio have been explored such as the FeCo-based, Fe-based and Heusler alloys heterostructures.[10–12] Moreover, the strategy with incorporation of heavy metal (HM) capping layers has been confirmed to push forward the advancement of VCMA behavior reaching few hundreds of fJ/Vm.[13, 14] However, the PMA and VCMA coefficient of these traditional magnetic ultrathin films mainly originate from the complicated interfacial effect and electric field induced charge redistribution.[15–17] Therefore, the qualities of HM/FM and FM/insulators interfaces usually exhibit a decisive impact on their performances.[16] In view of the problem of traditional magnetic ultrathin films in fabricating high-quality samples limited by their dangling

bond and high reactivity,[18] searching alternative ferromagnetic materials would be essential to address the two major challenges for future MeRAM devices.

Meanwhile, the emergence of intrinsic two-dimensional magnetism achieved in experiments further expand the scope of ideal platforms for developing the desired MeRAM.[19– 21] Instead of serving as a tunnel barrier by 2D dielectrics, these layered FM materials offer a new opportunity to substitute traditional magnetic thin films, and can be embedded as free or pinned layer to assemble MTJs.[18, 22–25] Regarding the stacking quality, the covalently bonded 2D surfaces possess the ability to fully decouple from various junctions which in turn can facilitate the high-quality interfaces.[26] Currently, research works have focused on the electron transport properties of various MTJs consisting of  $2D \,\mathrm{CrI}_3$ ,  $VSe_2$ or Fe<sub>3</sub>GeTe<sub>2</sub>, and revealed their giant tunneling magnetoresistance (TMR) ratios ranging from 78% to 3600%.[20, 27– 29] On the other hand, given the unique crystal structures in ultimate thickness dimension, these 2D magnetic materials exhibit a weak electric screening effect and excellent flexibility which allow the electrical control. In fact, the voltage-controlled switching between antiferromagnetic (AFM) and FM states were demonstrated on diverse 2D magnetic materials.[30–34] These results clearly suggest that the choice of 2D FM layers to design MTJ architectures may be a feasible alternative towards higher PMA and VCMA efficiencies.[35]

For a qualified 2D FM material used in the typical HM/FM/MgO junctions, several critical parameters are required: room-temperature ferromagnetism, suitable PMA energy as well as compatible crystal lattices to MgO substrate. Therefore,  $Fe<sub>2</sub>I<sub>2</sub>$  layer sharing analogous square lattice is one promising candidate FM material as its Curie temperature and strain-modified PMA reach 400K and 1.65 erg/cm<sup>2</sup>, respectively.[19] In terms of the capping layer, our prototype model will focus on the Ir with large SOC which was proved to be easily magnetized in experimental and theoret-

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FIG. 1. (a) Schematic atomic structure of the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir heterostructure. The unit cell is denoted by the black parallelepiped. (b) $\&c)$  shows the stacking configurations of the Fe2I2/Ir and MgO/Fe2I<sup>2</sup> interfaces, respectively.

ical works.[8, 36, 37] Notably, the modulation of PMA and electric field control of magnetism remains unexplored thus far in such kind of multilayers involving 2D magnetic layers.

In this work, we present systematic first-principles calculations for the  $Ir/Fe<sub>2</sub>I<sub>2</sub>/MgO$  heterostructures with the different Ir capping thickness. Both giant PMA and VCMA coefficient have been demonstrated in such multilayers depend on the biaxial strain. Some of the systems show a reorientation of magnetic easy axis in responding to the change of Ir thickness. In addition, we find that the strain effect leads to distinct VCMA responses, which was verified by the change of SOC energy differences. The underlying atomic mechanism mainly lies on electric-field-induced shift of spin-polarized d*<sup>z</sup>* <sup>2</sup> and d*yz* orbitals on the Ir layer. These findings pave the way for the development of high-efficient MTJs based on 2D FM material platform in the next-generation MeRAM devices.

*Methodology:* Our density functional theory (DFT) calculations were performed in conjunction with the projector augmented-wave (PAW) scheme, as implemented in the plane-wave basis Vienna *ab initio* simulation package (VASP).[38] The generalized gradient approximation (GGA) parametrized by Perdew–Burke–Ernzerhof (PBE) was used for exchange and correlation contributions.[39] A kinetic energy cutoff of 500 eV was chosen for the plane-wave expansion of wave functions and the Monkhorst–Pack scheme of  $32 \times 32 \times 1$  k-point sampling was adopted for the integration over the first Brillouin zone.[40] We applied a vacuum space of 15  $\AA$  along the z direction to separate the periodic films and the dipole corrections are also taken along [001]. All structures are fully optimized until the residual forces acting on the ions become less than  $0.01$  eV/ $\AA$ . The convergence criteria for the energy of 10−<sup>6</sup> eV was met. Spin-orbit coupling (SOC) is included in the calculations self-consistently. The DFT+*U* method was employed for the treatment of the strong correlated 3*d* elections on Fe orbitals ( $U_{eff}$  = 2.5 eV).[41] The magnetocrystalline anisotropy (MCA) energy per unit interfacial area is defined as  $E_{MCA} = (E_{[100]} - E_{[001]})/A$ , where  $E_{[100]}$ and  $E_{[001]}$  represent the total energy with in-plane and out-ofplane magnetization, respectively, and *A* is the in-plane area of the unit cell.

*Results and Discussion:* Fig.1(a) shows the stacking configurations of  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(n)$  trilayer which are modeled by a slab supercell along the [001] direction. These considered multilayers consist of 2D Fe<sub>2</sub>I<sub>2</sub> sandwiched by five MLs of rock-salt MgO and n-MLs fcc Ir (n =  $1~\sim 5$ ). The Ir atoms located in  $Fe<sub>2</sub>I<sub>2</sub>/Ir$  interfaces are placed atop of the I atoms of  $Fe<sub>2</sub>I<sub>2</sub> ML$ , and the MgO/Fe $<sub>2</sub>I<sub>2</sub>$  interfaces are designed by plac-</sub> ing I atoms atop Mg atoms, which are the most stable crystalline structures (See Fig.1(b and c). The lattice constants of Fe<sub>2</sub>I<sub>2</sub> ML and initial bulk Ir are  $3.810$  and  $3.839$  Å, respectively, resulting in a negligible mismatch around 0.76%. It should be noticed that this mismatch ratio is much smaller that these of the FeCo-based junctions, and would promote the fabrication of high-quality interfaces in such heterostructures. Here, the in-plane lattice constants of the  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir$  multilayer are set equal to that of bulk Ir (denoted as  $a_1$ ) and 2D Fe<sub>2</sub>I<sub>2</sub> (denoted as  $a_2$ ), respectively, allowing for the study of the strain modulated effects. In fact, the optimized multilayers in Fig.1(a) indeed represent atomically sharp and chemically clean interfaces without any disorder. The interlayer distances of Fe<sub>2</sub>I<sub>2</sub>/Ir(1) and MgO/Fe<sub>2</sub>I<sub>2</sub> are 2.734 and 3.267 Å, respectively, under the lattice of  $a_1$ . When the lattice constant of the multilayer decreases to  $a_2$ , these interlayer distances become slightly enlarged to  $2.790$  and  $3.298$  Å, respectively. These stacking features also highlight the advantage of 2D materials which can largely decouple from the substrates and preserve their individual properties.

Having substantiated the stacking configurations of  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(n)$  heterostructures, in Fig.2(a) we show the spin-resolved band structure of MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir (n=1) with lattice of al as an example. It can be seen that the  $Fe<sub>2</sub>I<sub>2</sub>$ resolved band dispersion is clearly identifiable compared with our previous results, where the majority-spin channel yields a large band gap and the minority-spin shows linear band crossing.[19] The survived electronic structure of  $Fe<sub>2</sub>I<sub>2</sub>$  ML mainly originates from the clean stacking interfaces. However, we find that the Fermi level shifts downward in energy below the crossing points. This finding suggests that interfacial charge transfer emerges in the interfaces of the multilayers. To gain more insight on the electronic migration, Fig.2(b) depicts the charge difference  $(\Delta_{\rho})$  of the Fe<sub>2</sub>I<sub>2</sub>/Ir interface,  $\Delta \rho = \rho_{MgO/Fe_2I_2/Ir} - \rho_{MgO/Fe_2I_2} - \rho_{Ir}$ , where  $\rho_{MgO/Fe_2I_2/Ir}$ ,  $\rho_{MgO/Fe_2I_2}$  and  $\rho_{Ir}$  refer to the electron densities of the  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir$  system, individual  $MgO/Fe<sub>2</sub>I<sub>2</sub>$  and Ir layers, respectively. We find that a notable electronic redistribution localizes at the  $Fe<sub>2</sub>I<sub>2</sub>/Ir$  interface with an accumulation of charges (red sphere in Fig.2(b)) into the Ir layer originating from the interfacial I atoms (blue sphere in Fig.2(b)). The plane-integrated electron density also confirms that electrons are depleted on the  $Fe<sub>2</sub>I<sub>2</sub>$  ML, giving a variation of 0.482 *e*<sup>−</sup>. These findings are consistent with the mentioned shift of Fermi level for the  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir$  system.



FIG. 2. (a) Atom- and spin-resolved band structures of the MgO/Fe2I2/Ir(1) heterostructure based on the GGA+*U* method. (b) Plane-integrated electron density difference,  $\Delta_{\rho}$ , for the MgO/Fe2I<sub>2</sub>/Ir(1) heterostructure. The inset is the 3D isosurface of the electron density difference where the red and blue areas represent electron accumulation and depletion, respectively. (c) Variation of the MCA for the MgO/Fe2I2/Ir as a function of Ir thickness (n) with different in-pane lattice constants of  $a_1 = 3.839$  Å and  $a_2 = 3.810$  Å, respectively. The horizontal dashed line denotes the shape anisotropy energy of  $K^s$ .

The magnetic moments of Fe vary between 2.975 and 3.069  $\mu$ <sub>B</sub> with different thickness of Ir layers, which is in agreement with the cases of 2D Fe<sub>2</sub>I<sub>2</sub> ( $\sim$ 2.973  $\mu$ <sub>*B*</sub>). Besides, the emergence of local moments on the capped Ir are also confirmed in the  $Ir/Fe<sub>2</sub>I<sub>2</sub>$  interfaces. The single Ir capping layer (n=1) yields the largest magnetic moments of 1.224 and 0.986  $\mu_B$  per atom with the lattice constant of  $a_1$  and  $a_2$ , respectively. These values of local moments are enhanced compared with the strain-induced moments on the free-standing Ir layer, which should originate from the proximity effect. In addition, we noticed that the compressive strain with the lattice constant of *a*<sup>2</sup> result in the reduction of magnetic moments on Ir, consistent with reported monotonic behavior of straindriven magnetism of the free-standing Ir.[42] Therefore, the external strain and proximity effect show the synergistic effects to the induced magnetism of the Ir capping layers. For n  $\geq$  2 the local moments of interfacial Ir decrease dramatically and remain constant around 0.03  $\mu$ <sub>*B*</sub>, which is one order of magnitude smaller than these of FeCo/TM interfaces.[11, 37] On the other hand, the local moments of Ir layers away from the  $Fe<sub>2</sub>I<sub>2</sub>/Ir$  interfaces progressively decrease as the thickness increasing, suggesting a thinner Ir capping layer would trigger more significant influences on the magnetic properties of MgO/Fe2I2/Ir heterostructure.

Given the large spin-orbit-coupling (SOC) of Ir, the induced magnetized Ir caps would directly change the MCA of  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir$  multilayers. In Fig.2(c) we show the calculated MCA for the multilayers with Ir-cap thickness n ranging from 1 to 5 MLs. For the MgO/Fe2I2/Ir system with 1-ML Ir-cap, the MCA is found to be 3.461 and  $6.285$  erg/cm<sup>2</sup> with the lattice constant of  $a_1$  and  $a_2$ , respectively, indicating a preference for out-of-plane magnetization. These MCAs are greatly enhanced by a factor of 4∼8 compared with that of the pristine  $Fe<sub>2</sub>I<sub>2</sub> ML$ . Besides, the strain strategy also exhibits critical effects on the modification of MCA in the capped system since a compressive strain of 0.76% yields a reduction of MCA by a factor of 1.8. Employing the Bruno expression, we have calculated the orbital moment anisotropy ( $\Delta M_o = M_o^{001} - M_o^{100}$ ) of Ir under different lattice constants. Our results show that the  $\Delta M_o$  changes from 0.162 to 0.106  $\mu_B$  as the multilayers undergo compressive strain, correlating with that of the MCA. The dashed line in Fig. 2(c) indicates the shape anisotropy energy,  $K^s = -\mu_0 M_s^2 t/2 = -0.402$  erg/cm<sup>2</sup>, of the magnetic slab, which is about one order of magnitude smaller than its MCA.

Regarding the effect of Ir thickness (n) on the magnetic anisotropy, as shown in Fig.2(c), the MCA declines dramatically below 0.86 erg/cm<sup>2</sup> with 2 to 5 MLs Ir caps (n = 2∼5). Similar behaviors have been reported in the Ir capped FeCo thin films. However, unlike the variation trend of FeCo/Ir, the  $MCA$  for  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir$  system increases monotonically with thicker Ir capping layers, which eventually saturates at ∼0.86  $erg/cm<sup>2</sup>$ . It is noteworthy that the corresponding multilayers with 2- or 3-MLs Ir-cap exhibit negative magnetic anisotropy energies (below the dashed line) upon taking into account  $K^s$ , suggesting an out-of-plane to an in-plane magnetization reorientation. The underlying origin is the diminished magnetic and orbital moments on the Ir, yielding a negative contribution to the MCA. When the Ir thickness reaches 5 MLs, the Fe atoms with  $\Delta M$ <sup>*o*</sup> around 0.032∼0.043 μ<sub>*B*</sub> dominantly contribute the MCA of the MgO/Fe $_2$ I<sub>2</sub>/Ir system, leading to the MCA ranging from 0.847 to 0.861 erg/cm<sup>2</sup> similar to that of  $2D Fe<sub>2</sub>I<sub>2</sub>$  (orange stripe area). These findings demonstrate that the Ir cap layer can achieve the efficient modification of the magnetic properties of 2D  $Fe<sub>2</sub>I<sub>2</sub>$ , which is essential for future spintronic applications.



FIG. 3. The MCA of the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir (n = 1) heterostructure as a function of electric field in MgO with lattice constant of (a)  $a_1 =$ 3.839 Å and (b)  $a_2 = 3.810$  Å, respectively.

Next, to further probe the VCMA coefficient, an electric field is applied perpendicular to the  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1)$  heterostructure as an example in view of that 1ML Ir induces the highest PMA in the proposed multilayers. Figure 3 shows the variation of the MCA as a function of the electric field in MgO under different lattice constant. The VCMA coefficient (β) can be expressed in the linear regime as β = ∆*EMCA*/*E<sup>I</sup>* , where ∆*EMCA* is the electric-field-induced change of MCA, and  $E_I$  is the electric field in the insulator. We find that the epitaxial strain give rise to a dramatic effect on the magnitude and sign of the VCMA efficiency. More specifically, the Ir-capped multilayer for the  $a_1$  lattice constant shows an asymmetric *V*-shape field behavior with  $β$  values of 1791 (-881) fJ/Vm above (below) the critical field  $E_I = -0.185$  V/nm. Here, the left part of the *V*-shape field behavior was confirmed by another point of MCA of 6.046 erg/cm<sup>2</sup> under the E-field of -0.370 V/nm. In contrast, the VCMA behavior for the *a*<sup>2</sup> lattice constant exhibits a linear response with the β value of -1956 fJ/Vm. These optimized values are comparable to the TM-based FeCo or Mn<sub>3</sub>Ga junctions, which are about one order of magnitude higher than these of most experimentally reported values of FeCo-based MTJs with  $β$  in the range of 20∼300 fJ/Vm. Furthermore, we determined the influence of the Ir cap thickness on the VCMA coefficients, where the  $\beta$ values for the 1ML- and 3MLs Ir caps are summarized in Table I. We find that increasing the Ir overlayer thickness leads to the reduction in the VCMA coefficient, which shows asymmetric Λ-shape E-field behavior with a sign reversal. Here, the maximum  $|\beta|$  for MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(3) decreases to 593 and 701 fJ/Vm under  $a_1$  and  $a_2$  lattice constant, respectively. Therefore, these findings demonstrate that the 1ML-Ir-capped  $Fe<sub>2</sub>I<sub>2</sub>$ multilayers could service as excellent material platforms for magnetoelectric spintronics, since the predicted giant VCMA coefficient values fulfills the requirement to achieve a switching bit energy below 1 fJ in the next-generation of MeRAM. In addition, the feasibility of controlling the VCMA behavior

TABLE I. The VCMA coefficient,  $\beta$  (in fJ/Vm), of the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir multilayers versus Ir cap thickness (in ML) under  $a_1 = 3.839$  Å and  $a_2$ =3.810 Å.

$\beta$ (fJ/Vm)	$1ML-Ir$			$3ML-Ir$
$a_1$	-881	1791	593	$-525$
$a_2$		$-1956$	186	-701

via strain engineering was also demonstrated to design desired MeRAM devices.

In order to elucidate the microscopic origin of the giant VCMA efficiency in the MgO/Fe $_2I_2/Ir(1)$  multilayer, we have also calculated the atom-resolved SOC energy difference  $\Delta E_{SOC} = (E_{SOC}^{100} - E_{SOC}^{001})/2$ .[43] Here,  $E_{SOC}^{100}$  and  $E_{SOC}^{001}$  represent the SOC energies with in-plane and out-of-plane magnetization, given by

$$
E_{\rm soc} = \frac{\hbar^2}{2m^2c^2} \left\langle \frac{1}{r} \frac{dV}{dr} \vec{L} \cdot \vec{\hat{S}} \right\rangle, \qquad (1)
$$

where  $V(r)$  is the spherical part of the effective potential within the PAW sphere. In view of the failure of perturbation theory due to the strong SOC of Ir ( $\xi$  = 451 meV), in Fig.4(a) we display the E-field induced changes of ∆*ESOC* and MCA for the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1) with lattice constant of  $a_1 = 3.839$  Å as an example, where  $\delta E_{SOC} = \Delta E_{SOC} (E_I^1) - \Delta E_{SOC} (E_I^2)$  and  $\delta(MCA) = MCA(E_I^1) - MCA(E_I^2)$ . It should be noted that the electric field-induced variation of δ*ESOC* has been found to correlate well to that of  $\delta(MCA)$  sharing the similar asymmetric *V*-shape field behavior. Overall, the results reveal that Ir-resolved  $\delta E_{SOC}$  yields the dominant contribution to the total  $\delta E_{SOC}$  under the perpendicular electric field, indicating the giant VCMA coefficient mainly originate from the magnetized Ir. Similar conclusions have also been reported for 5*d*-TM modified magnetic thin films. However, our findings emphasize the feasibility to optimize 2D magnetism by stacking ultrathin 5*d*-TM layers with high-quality interfaces, giving rise to giant PMA and VCMA efficiency for MeRAMs.

The electric-field-driven modulation of MCA via the capped Ir can also be understood from the second-order perturbation theory, which assumes the MCA modifications are induced by changes of coupling strength between occupied and unoccupied *d*-states through the orbital angular momentum operators  $\hat{L}_z$  and  $\hat{L}_x$ , given by:[42, 44]

$$
MCA \approx \sum_{I, lmm'\sigma\sigma'} \sum_{o,u} \sigma\sigma' \xi_{Il}^2 \frac{P_u^{Ilm\sigma} P_o^{Ilm'\sigma'}}{E_u^{\sigma} - E_o^{\sigma'}} \Delta L_{l,mm'}, \qquad (2)
$$

where

$$
\Delta L_{l,mm'} = |\langle lm|\hat{L}_z\left|lm'\right\rangle|^2 - |\langle lm|\hat{L}_x\left|lm'\right\rangle|^2. \tag{3}
$$

Here,  $\hat{\xi}$  refers to the diagonal matrix containing the SOC strength;  $\hat{L}_z(\hat{L}_x)$  denotes the z(x) component of the orbital angular momentum operator;  $P_n^{Ilm\sigma} = |\langle \Psi_n^{\sigma} | Ilm \rangle|^2$  is the Bloch wave amplitude projected on Ir atoms, orbital index *lm*, spin



FIG. 4. (a) The electric-field-induced changes of SOC energies and MCA, i.e.  $\delta E_{SOC}$  and  $\delta(MCA)$ , relative to zero-field  $(E_I^2=0)$  for the  $MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1)$  heterostructures. (b) The spin-projected density of states (DOS) of  $d_{yz}$  and  $d_{z^2}$  orbitals for Ir under  $E_I$ . Here,  $E_I > 0$  ( $E_I$  $<$  0) denotes the applied E-field of +0.278 (-0.185) V/nm in MgO. Energy and k-resolved distributions of the minority-spin d-orbital of Ir under the external E-field of (c) -0.185 V/nm and (d) +0.278 V/nm, respectively.

index  $\sigma$ ,  $\Psi_n^{\sigma}(E_n^{\sigma})$  is the one-electron occupied and unoccupied spin-polarized Bloch states (energies) of band index *n* and wave vector *k* (omitted for simplicity). Analysis of the orbital-resolved density of states (DOS) of the capped Ir atoms shows that the spin-minority occupied  $d_{z^2}$  and unoccupied  $d_{yz}$ are primarily responsible for the linear VCMA behavior under the E-field ranging from -0.185 to 0.278 V/nm. As shown in Fig.4(b) the electric field has a larger effect on  $d_{z^2}$ -derived Ir projected DOS, which shift downward in energy as the *E<sup>I</sup>* changes from negative to positive. In view of the unchanged  $d_{yz}$  states, this would cause a weaker of SOC coupling between  $d_{z^2}$  and  $d_{yz}$  through the  $\hat{L}_x$  operator, which in turn reduces the negative term of  $\langle d_{\mathcal{A}}^{\downarrow}$  $\int_{z^2}^{1} \left| \hat{L}_x \right| d_{yz}^{\downarrow}$  and eventually enhances the MCA.

In Fig.4(c and d) we also show the minority-spin bands of the Ir-derived  $d_{z^2}$  and  $d_{yz}$  states with E-field of -0.185 and 0.278 V/nm, respectively. The results reveal that the SOC matrix element of  $\langle d_{\vec{x}}^{\downarrow}$  $\frac{1}{z^2} \left| \hat{L}_x \right| d_{yz}^{\downarrow}$  mainly yield negative contributions around  $\Gamma$ -X-M. Except for the rigid shift of the occupied  $d_{z^2}$  (gray area), it should be noted that the  $d_{z^2}$  states around Γ between -0.4 to 0 eV were eliminated when the E-field changes from -0.185 to 0.278 V/nm. Consequently, the vanished negative contributions around  $\Gamma$  further increases

the MCA of the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1) junction, yielding a positive VCMA coefficient. These findings in the electric-fieldinduced shift of the Ir-derived DOS are similar to other TMcaped traditional magnetic thin films, while the electric-field mainly changes the atomic positions along the z-direction and thus influences the charge accumulation and the out-of-plane *d* states.



FIG. 5. The *d* orbital-projected contributions of Ir to the electricfield-induced change of SOC energy difference, δ*ESOC*, for the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1) multilayer with lattice constant of (a)  $a_1 = 3.839$  Å and (b)  $a_2 = 3.810 \text{ Å}$ , respectively. Here,  $E_I^1$  and  $E_I^2$  are 0.278 (0.278) and -0.185(-0.278) V/nm in the left (right) panel, respectively.

In order to understand the sign reversal of the VCMA coefficient from 1791 to -1956 fJ/Vm with decreasing lattice constant shown in Fig.3(a) and (b), we display the corresponding *d* orbital-projected contributions of Ir atoms to the electric-field-induced change of SOC energy difference in Fig.5. For the larger lattice constant, Fig.5(a) reveals that the spin parallel SOC matrix element,  $\left\langle d_{z^2}^{\sigma} \right| \hat{L}_x | d_{yz}^{\sigma} \rangle$ , yields the dominant contribution to the increase of  $\delta E_{SOC}$ . This finding indicates that the negative value of the SOC between  $d_{z^2}^{\sigma}$ and  $d_{yz}^{\sigma}$  through in-plane orbital angular-momentum operator  $(\hat{L}_x)$  becomes weaker when the  $E_I$  changes from -0.185 to 0.278 V/nm, resulting in a positive VCMA coefficient. Furthermore, for the smaller lattice constant, the monotonic decrease of MCA originates primarily from the spin-parallel SOC of the  $\left\langle d_{z^2}^{\sigma} \Big| \hat{L}_x | d_{yz}^{\sigma} \right\rangle$  and  $\left\langle d_{yz}^{\sigma} \Big| \hat{L}_x \Big| \frac{d_{x^2-y^2}}{dx^2} \right\rangle$  matrix elements shown in Fig.5(b). In contrast to the results for  $a_1$ = 3.839 Å, the electric-field-induced changes of  $\left\langle d_{z^2}^{\sigma} \right| \hat{L}_x | d_{yz}^{\sigma} \right\rangle$ and  $\langle d_{yz}^{\sigma} | \hat{L}_x | d_{x^2-y^2}^{\sigma} \rangle$  are negative, indicating these spin parallel SOC terms become more negative as the electric field increases leading in turn to the VCMA coefficient sign reversal.

*Conclusion:* In summary, we propose a new general approach to assemble high-efficient voltage-controlled MTJ based on the 2D magnetic  $Fe<sub>2</sub>I<sub>2</sub>$  layer, instead of the traditional ferromagnetic thin films, with Ir capping layers. Employing *ab initio* electronic structure calculations, the key parameters, such as the Ir capping layer thickness and epitaxial strain, have been investigated to optimize the MCA and VCMA coefficient for the MgO/Fe $_2I_2/Ir$  stacking architectures. We find that the MCA magnitude and spin orientation as well as the VCMA coefficients exhibit strong dependence on Ir thickness. Notably, reducing the Ir thickness down to monolayer can significantly enhance the MCA of the multilayers, and spin reorientation has been achieved in the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(2/3) heterostructure. These primarily arise from the change of magnetic and orbital moments in Ir layers, which dominated by the biaxial strain and proximity effect. In addition, we predict extremely-high VCMA coefficients with a sign reversal behavior for the MgO/Fe<sub>2</sub>I<sub>2</sub>/Ir(1) system exceeding the required 1000 fJ/Vm threshold, where the VCMA magnitude and slope depend dramatically on biaxial strain. Our results reveal that the underlying origin of the VCMA behavior lies on the electric-field-induced modifications of SOC

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