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Tunnel barriers for Fe-based spin valves providing high spin-polarized current

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Employing density functional theory for ground state quantum mechanical calculations and the non-equilibrium Green's function method for transport calculations, we investigate the potential of CdS, ZnS, Cd₃ZnS₄, and Zn₃CdS₄ as tunnel barriers in magnetic tunnel junctions for spintronics. Based on the finding that the valence band edges of these semiconductors are dominated by p_z orbitals and the conduction band edges by s orbitals, we show that Δ₁ symmetry filtering of the Bloch states in magnetic tunnel junctions with Fe electrodes results in high tunneling magnetoresistances and high spin-polarized current (up to two orders of magnitude higher than in the case of the Fe/MgO/Fe magnetic tunnel junction).

1. Introduction

The resistance of an anisotropic magnetoresistance (AMR) device depends on the angle between the electric current and the external magnetic field, being minimal/maximal for perpendicular/parallel orientation.¹ The resistance of a giant magnetoresistance (GMR) device, consisting of a non-magnetic metal (such as Cu and Pt) sandwiched between ferromagnetic electrodes, depends on the relative orientation of the magnetic moments of the electrodes, being minimal/maximal for parallel/antiparallel P/AP orientation.^{2,3} In a tunneling magnetoresistance (TMR) device the non-magnetic metal is replaced with an insulator.⁴ Coherent tunneling through a crystalline barrier results in

$$\text{TMR} = (T_P - T_{AP})/T_{AP}, \quad (1)$$

where T_P/T_{AP} is the transmission for P/AP orientation of the magnetic moments of the electrodes, while incoherent tunneling through an amorphous barrier results in

$$\text{TMR} = 2P_L P_R / (1 - P_L P_R), \quad (2)$$

where P_L/P_R is the density of states at the Fermi energy (E_F) of the left/right electrode.⁵ In the case of coherent tunneling in an epitaxial junction the transverse component k_{\parallel} of the wave vector \mathbf{k} is conserved. The resistance depends not only on the relative orientation of the magnetic moments of the electrodes but also on the orbital angular momentum symmetry of the wave function of the tunneling electrons. While various junctions^{6–8} realize such symmetry filtering, only the combination of a MgO barrier with Fe electrodes is adopted in main-stream

applications, mainly because boron assisted epitaxial growth is used to minimize spin flip and interband scattering.^{9,10}

Achieving a high spin-polarized current is paramount for magnetic random access memories in that switching between the magnetic configurations storing binary information is achieved by spin transfer torque,^{11–13} because the spin transfer torque is proportional to the generating spin-polarized current.^{14,15} The same applies, for example, to magnetic flip-flops in that the magnetic state is switched by spin-polarized current.^{16,17} As it is vital to identify magnetic tunnel junctions (MTJs)

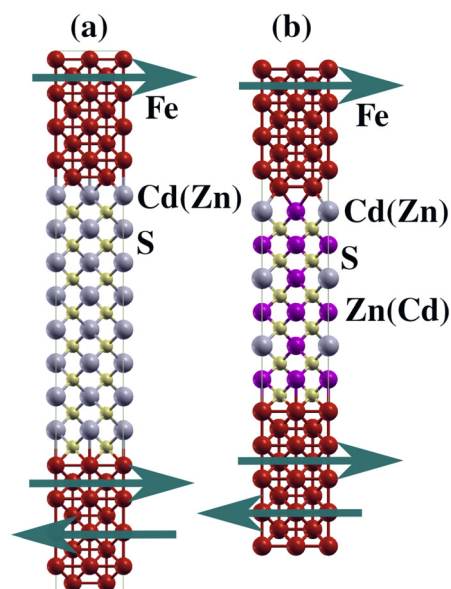


Fig. 1 Scattering regions of the (a) Fe/CdS/Fe (Fe/ZnS/Fe) and (b) Fe/Cd₃ZnS₄/Fe (Fe/Zn₃CdS₄/Fe) MTJs. The arrows represent the magnetic moments of the two Fe electrodes, which can be oriented parallel or antiparallel. Color code: Fe = red; Cd = grey; Zn = magenta; S = yellow.

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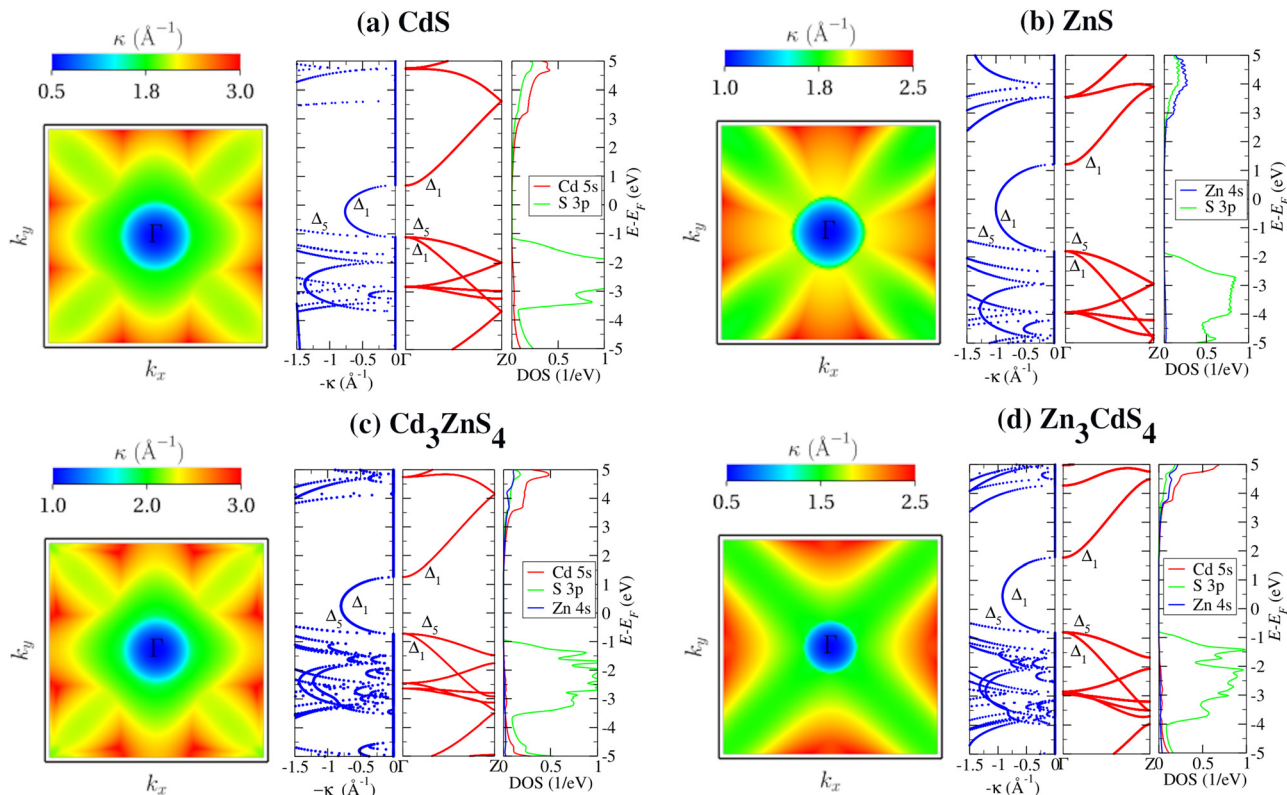


Fig. 2 Electronic structure analysis of (a) CdS, (b) ZnS, (c) Cd₃ZnS₄, and (d) Zn₃CdS₄. The left panels show heat maps (from the minimal to the maximal value on a linear scale) of $\kappa(E_F, \mathbf{k}_{\parallel})$ in the BZ, the middle panels show the real (red) and complex (blue) band structures, and the right panels show the partial densities of states.

based on semiconductors that can be integrated with the current lithographic processes to open new avenues for spintronics, we present here a detailed theoretical analysis of potential barrier materials with zinc blende crystal structure (CdS, ZnS, Cd₃ZnS₄, and Zn₃CdS₄) that are extensively used industrially in light emitting diodes.^{18–20} For the electrodes we use Fe as strong ferromagnet with high Curie temperature.²¹ The bcc structure of Fe shares with the zinc blende structure the 4-fold rotation symmetry about the [001] direction, enabling growth of a MTJ by chemical vapor deposition and molecular beam epitaxy. Being II–VI compounds with direct band gaps, CdS and ZnS are also applied as transparent semiconductors^{22,23} and for photocatalysis.^{24,25} The excitonic properties, particularly the binding energy, can be tuned well by doping.²⁶ High quantum transport is possible due to dispersive bands with small electron effective masses of 0.18 m_0 (CdS)²⁷ and 0.34 m_0 (ZnS),²⁸ where m_0 is the free electron mass.

After introducing our computational methodology, we will analyze the properties of Fe/CdS/Fe, Fe/ZnS/Fe, Fe/Cd₃ZnS₄/Fe, and Fe/Zn₃CdS₄/Fe MTJs based on the real and complex band structures of the barrier and electrode materials, to prepare for a detailed discussion of the transmission and TMR. A summary of the main findings is provided in the conclusion section.

II. Computational details

The ground state electronic properties of CdS, ZnS, Cd₃ZnS₄, and Zn₃CdS₄ are obtained from Kohn–Sham density functional

theory using the SIESTA engine²⁹ with non-relativistic norm-conserving Troullier–Martins pseudopotentials and localized atomic orbitals as basis set. The exchange–correlation functional is treated in the Ceperley–Alder parametrization of the local density approximation. Double- ζ basis functions are used for the s, p, and d orbitals of Fe, Cd, and Zn, while double- ζ plus polarization basis functions are used for the s and p orbitals of S. The atomic self-interaction correction (ASIC) is employed to correct the band gaps.^{30,31} We use a grid spacing equivalent to a plane wave cutoff of 400 Ry and a Monkhorst–Pack $8 \times 8 \times 8$ \mathbf{k} -point mesh. Complex band structures are obtained through the secular equation for $\mathbf{k}_{\parallel} = 0$.

The experimental lattice parameters of zinc blende CdS and ZnS are 5.83 Å³² and 5.41 Å,³³ respectively, and the optimized lattice parameters of Cd₃ZnS₄ and Zn₃CdS₄ are found to be 5.76 Å and 5.51 Å. We use a 2×2 supercell of body-centered cubic Fe to create MTJs and absorb the lattice mismatch by altering the lattice parameter of Fe (experimental value: 2.86 Å³⁴), which hardly modifies the electronic properties. The thickness of the barrier material is chosen as 26.75 Å (3 unit cells), 40.24 Å (7 unit cells), 20.69 Å (3 unit cells), and 19.79 Å (3 unit cells) for CdS, ZnS, Cd₃ZnS₄, and Zn₃CdS₄, respectively. Due to the larger band gap of ZnS an increased thickness is chosen to enhance the TMR. The atomic positions in the MTJs (inequivalent interfaces, see Fig. 1) are relaxed by the conjugate gradient method until all atomic forces stay below 0.01 eV Å⁻¹.



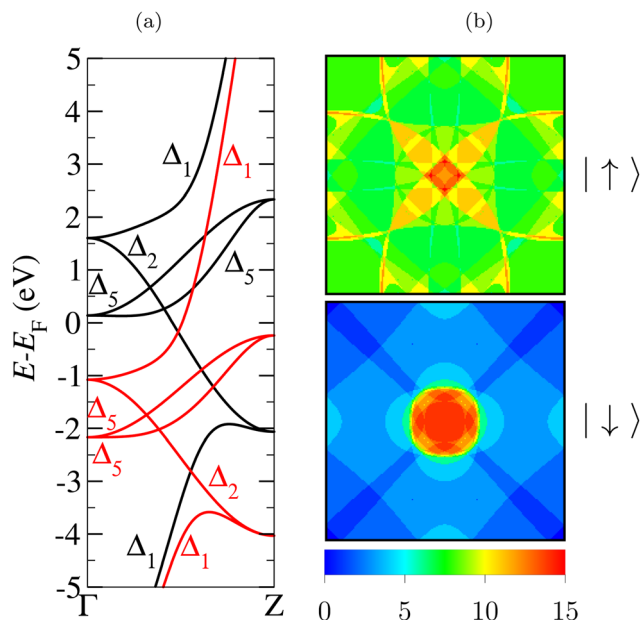


Fig. 3 (a) Real band structure with the Bloch symmetry indicated and (b) numbers of majority and minority spin channels in the BZ (simulation cell with 16 atoms) of body-centered cubic Fe.

The SMEAGOL engine,^{35–37} which is interfaced with the SIESTA engine, is used to execute non-equilibrium Green's function transport calculations. In this formalism the complex self-energy

$$\Sigma_\lambda^\sigma(E, V) = H_\lambda^{\sigma\dagger}(E, V)g_\lambda^\sigma(E, V)H_\lambda^\sigma(E, V), \quad (3)$$

where $\sigma = \uparrow/\downarrow$ is the spin, $H_\lambda^\sigma(E, V)$ is the coupling Hamiltonian between the electrode and scattering region, $g_\lambda^\sigma(E, V)$ is the surface Green's function of the electrode, and $\lambda = L/R$ denotes the left/right electrode, is added to the Hamiltonian $H_C^\sigma(E, V)$ of the scattering region to obtain the effective Green's function

$$G_C^\sigma(E, V) = \lim_{\eta \rightarrow 0} [E + i\eta - H_C^\sigma(E, V) - \Sigma_L^\sigma(E, V) - \Sigma_R^\sigma(E, V)]^{-1}. \quad (4)$$

$G_C^\sigma(E, V)$ is finite and Hermitian with density matrix

$$\rho^\sigma(V) = \frac{1}{2\pi i} \int_{-\infty}^{\infty} dE G_C^{\sigma<}(E, V), \quad (5)$$

where

$$G_C^{\sigma<}(E, V) = iG_C^\sigma(E, V) \left[\sum_\lambda f_\lambda^\sigma(E, V) \Gamma_\lambda^\sigma(E, V) \right] G_C^{\sigma\dagger}(E, V) \quad (6)$$

is the lesser Green's function, $f_\lambda^\sigma(E, V)$ is the Fermi–Dirac distribution function, and

$$\Gamma_\lambda^\sigma(E, V) = i(\Sigma_\lambda^{\sigma+}(E, V) - \Sigma_\lambda^{\sigma-}(E, V)) \quad (7)$$

is the coupling matrix between the electrode and scattering region (retarded/advanced part of the self-energy denoted by superscript +/–).

Using the spin-dependent transmission

$$T^\sigma(E, V) = \text{Tr}[\Gamma_L^\sigma(E, V)G_C^{\sigma-}(E, V)\Gamma_R^\sigma(E, V)G_C^{\sigma+}(E, V)], \quad (8)$$

the SMEAGOL engine calculates the voltage-dependent spin-polarized current in the Landauer–Büttiker formalism as

$$I^\sigma(V) = \frac{e}{h} \int dE T^\sigma(E, V) [f_L^\sigma(E - \mu_L) - f_R^\sigma(E - \mu_R)], \quad (9)$$

where e is the electron charge, h is the Planck constant, and $\mu_{L/R} = E_F \pm eV/2$ is the chemical potential. In the barrier the normal component of the wave vector becomes complex, $k_\perp(E, \mathbf{k}_\parallel) = i\kappa(E, \mathbf{k}_\parallel)$, with

$$\kappa(E, \mathbf{k}_\parallel) = \sqrt{2m(U - E)/\hbar^2 + \mathbf{k}_\parallel^2 - \langle \phi | \left(\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} \right) | \phi \rangle} / \langle \phi | \phi \rangle, \quad (10)$$

where m is the effective mass, U is the barrier height, and the last term takes into account that the decay in the barrier is stronger when the transverse angular momentum of the wave function is higher. Since the junction is translationally invariant in the transverse plane, we can write

$$T^\sigma(E, V) = \frac{1}{\Omega_{\text{BZ}}} \int_{\text{BZ}} d\mathbf{k}_\parallel T_0^\sigma(E, V) e^{-2\kappa(E, \mathbf{k}_\parallel)d}, \quad (11)$$

where Ω_{BZ} is the volume of the two-dimensional Brillouin zone (BZ), the \mathbf{k}_\parallel -dependent transmission $T_{\mathbf{k}_\parallel}^\sigma(E, V) = T_0^\sigma(E, V) e^{-2\kappa(E, \mathbf{k}_\parallel)d}$ is determined by the atomic composition and structure of the junction, and d is the thickness of the barrier. We calculate $T(E) = T^\uparrow(E, 0) + T^\downarrow(E, 0)$ for both the P and AP orientations to obtain the TMR through eqn (1). The transport calculations are performed after converging the charge density matrix with a tolerance of $10^{-4} e$. We show in the following results obtained for a $50 \times 50 \times 1$ \mathbf{k} -point mesh, while no relevant change is observed in test calculations for a $100 \times 100 \times 1$ \mathbf{k} -point mesh.

III. Results and discussion

While the experimental band gaps of CdS and ZnS are 2.6 eV²⁴ and 3.7 eV,²² respectively, we obtain appropriate ASIC values of 1.9 eV and 3.0 eV. For Cd₃ZnS₄ and Zn₃CdS₄ we obtain ASIC band gaps of 2.0 eV and 2.5 eV, respectively. Note that the size of the band gap scales $T_{\mathbf{k}_\parallel}^\sigma(E, V)$ but does not change its distribution in the BZ. The heat maps of $\kappa(E_F, \mathbf{k}_\parallel)$ in Fig. 2 show a four-fold rotation symmetry. Mainly the regions around the Γ -point and along the diagonals of the BZ contribute to $T^\sigma(E_F, V)$ due to low values of $\kappa(E_F, \mathbf{k}_\parallel)$, which also turns out to be true for $E \neq E_F$. As the tunneling electrons experience minimal $\kappa(E, \mathbf{k}_\parallel)$ at the Γ -point, that is, when they approach the barrier in the normal direction (minimal effective barrier thickness), the following analysis focusses on the Γ -point. However, note that this provides only a qualitative picture, because $T^\sigma(E, V)$, while being dominated by the Γ -point, generally depends on the entire BZ. $T^\sigma(E, V)$ of the MTJ will be high (and therefore the current will be high) if the symmetries of the



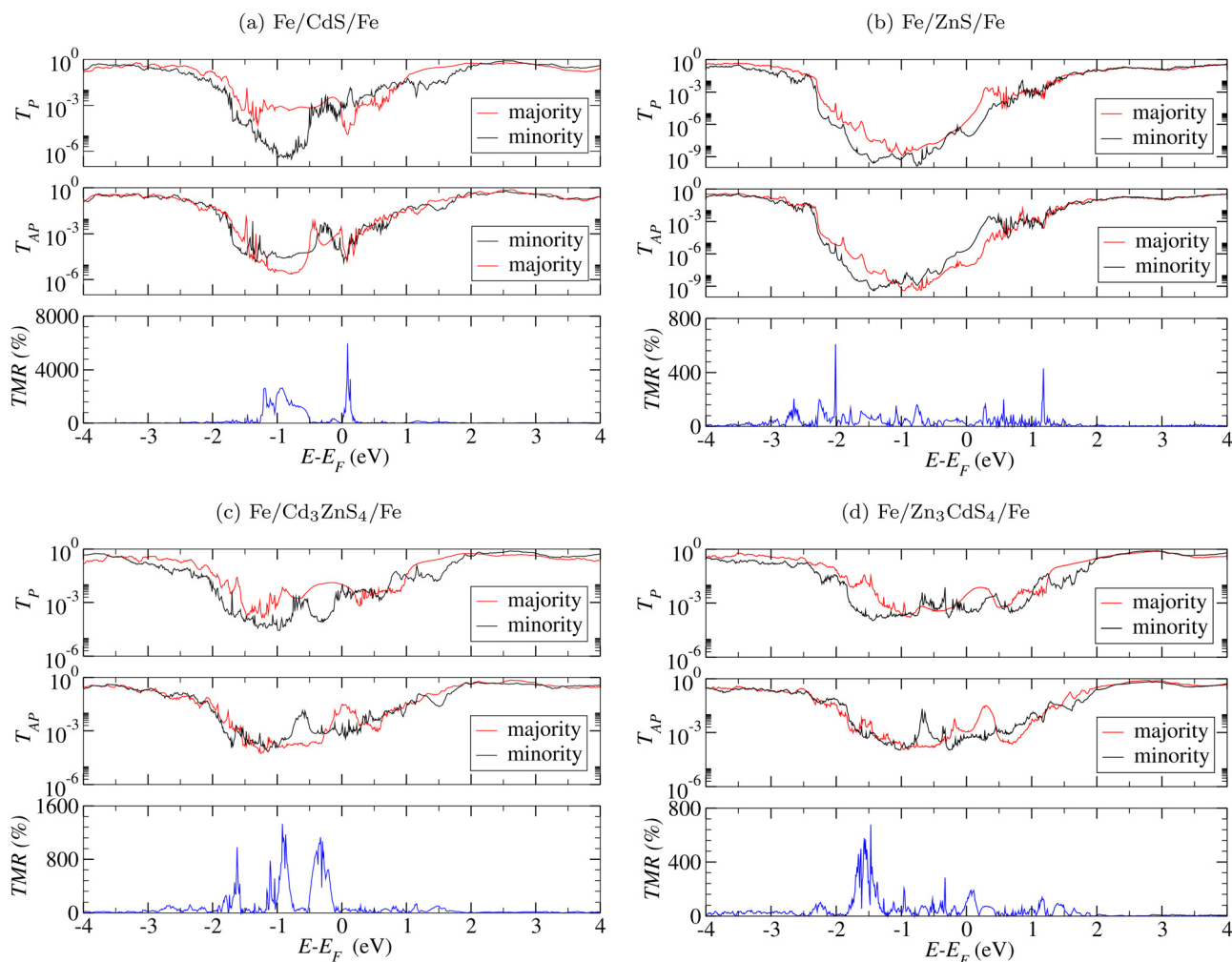


Fig. 4 Zero-bias transmission and TMR of the (a) Fe/CdS/Fe, (b) Fe/ZnS/Fe, (c) Fe/Cd₃ZnS₄/Fe, and (d) Fe/Zn₃CdS₄/Fe MTJs as functions of energy.

Bloch states at E_F in the electrodes and at the band edges in the barrier are the same, and a high TMR is expected if only one spin channel (either majority or minority) is present in the ferromagnetic electrodes. The absence of an abrupt change in the Hartree potential at the interface of an epitaxial junction results in a weak Rashba field, which minimizes the spin flip and interband scatterings to assist coherent quantum tunneling.

According to Fig. 2, the four barrier materials under investigation exhibit direct band gaps and similar band dispersions along Γ -Z. No complex ghost bands due to the non-orthogonal orbital basis set are observed.³⁸ The valence band edges are dominated by p_z orbitals and the conduction band edges are dominated by s orbitals. These orbitals couple with the Bloch states of the electrodes. The symmetry Δ of a Bloch state is assigned by projecting the wavefunction to the atomic orbitals and analyzing the angular momentum along the transport direction. The s , p_z , and $d_{3z^2-r^2}$ orbitals have zero angular momentum along the transport direction and thus map to Δ_1 symmetry. Similarly, the p_x , p_y , d_{xz} , and d_{yz} orbitals map to Δ_5 symmetry, the d_{xy} orbital maps to Δ_2' symmetry, and the $d_{x^2-y^2}$

orbital maps to Δ_2 symmetry. The results of the symmetry assignment are shown in Fig. 2. Analysis of the complex band structures in Fig. 2 at the Γ -point suggests that all four barrier materials filter Bloch states of Δ_1 symmetry. In each case, a complex band of Δ_1 symmetry smoothly connects the valence band maximum to the conduction band minimum. On the other hand, the complex band of Δ_5 symmetry present at the valence band edge connects to deep inside the conduction band and thus hardly contributes to $T^\sigma(E, V)$, unless E_F of the MTJ is pinned close to valence band edge (implying that hopping dominates over tunneling and the MTJ behaves like a GMR device).

In the case of the epitaxial Fe/MgO/Fe MTJ the Δ_1 -symmetric majority spin states of Fe match exactly with the Δ_1 -symmetric states forming the valence and conduction band edges of MgO (at the Γ -point), resulting in a high TMR of $\sim 10\,000\%$.^{39,40} The band structure of body-centered cubic Fe along Γ -Z is shown in Fig. 3 together with heat maps of the numbers of majority and minority spin channels in the BZ (reflecting the four-fold rotation symmetry of the body-centered cubic structure). Both the majority and minority spin channels are concentrated



Table 1 Symmetry analysis

MTJ	Configuration	Energy range ($E - E_F$)	Dominant symmetry of majority spin channel	Dominant symmetry of minority spin channel
Fe/CdS/Fe	P	-3.0 eV to -0.5 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_1, Δ_2
	P	+0.0 eV to +1.0 eV	Δ_1	Δ_2, Δ_5
	P	+1.0 eV to +2.1 eV	Δ_1	$\Delta_1, \Delta_2, \Delta_5$
Fe/CdS/Fe	AP	-1.7 eV to -1.5 eV	Δ_2, Δ_5	Δ_2
	AP	-1.3 eV to -0.5 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_2
Fe/ZnS/Fe	P	-2.4 eV to -0.3 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_2
	P	-0.1 eV to +0.6 eV	Δ_1, Δ_5	Δ_2, Δ_5
Fe/ZnS/Fe	AP	-2.4 eV to -1.1 eV	Δ_2, Δ_5	Δ_2
	AP	-1.0 eV to +0.6 eV	Δ_1, Δ_5	Δ_2, Δ_5
Fe/Cd ₃ ZnS ₄ /Fe	P	-3.0 eV to -0.7 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_1, Δ_2
	P	-0.5 eV to +0.0 eV	Δ_1, Δ_5	Δ_2, Δ_5
	P	+0.7 eV to +1.0 eV	Δ_1	Δ_2, Δ_5
	P	+1.0 eV to +2.0 eV	Δ_1	$\Delta_1, \Delta_2, \Delta_5$
Fe/Cd ₃ ZnS ₄ /Fe	AP	-1.0 eV to -0.3 eV	Δ_1, Δ_5	Δ_2, Δ_5
	AP	-0.2 eV to +0.3 eV	Δ_1, Δ_5	Δ_2, Δ_5
	AP	+0.3 eV to +0.6 eV	Δ_1	Δ_2, Δ_5
	AP	+1.3 eV to +1.9 eV	Δ_1	$\Delta_1, \Delta_2, \Delta_5$
Fe/Zn ₃ CdS ₄ /Fe	P	-2.0 eV to -1.0 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_2
	P	-1.0 eV to +0.0 eV	$\Delta_1, \Delta_2, \Delta_5$	Δ_2, Δ_5
Fe/Zn ₃ CdS ₄ /Fe	AP	-1.0 eV to -0.5 eV	Δ_1, Δ_5	Δ_2
	AP	-0.5 eV to +0.3 eV	Δ_1, Δ_5	Δ_2, Δ_5

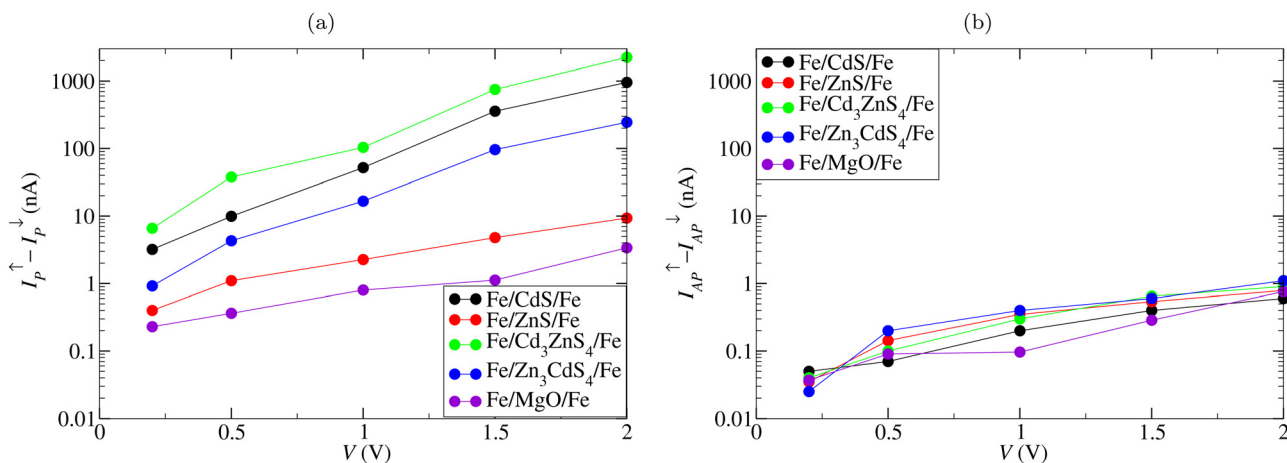


Fig. 5 Spin-polarized current in the (a) P and (b) AP configurations.

around the Γ -point. A majority spin Δ_1 band crosses E_F with high dispersion, while the corresponding minority spin Δ_1 band appears at higher energy due to the magnetic exchange field, resulting in 100% spin polarization in the Δ_1 channel. The Δ_2 and Δ_5 bands crossing E_F give only small contributions to $T^\sigma(E, V)$ due to mismatch with the Bloch state symmetry of the barrier (Δ_1).

Fig. 4 shows $T(E)$ for both the P and AP configurations separately for the majority and minority spin channels. The obtained dominant symmetries of the two spin channels at the Γ -point are summarized in Table 1. The Fe/CdS/Fe MTJ shows in the P configuration almost pure majority spin transmission in the range $-1.5 \text{ eV} < E - E_F < -0.5 \text{ eV}$. At E_F the TMR is 30% and at $E_F + 0.1 \text{ eV}$ the highest TMR of 5000% is obtained. The Fe/ZnS/Fe MTJ shows in both the P and AP configurations almost pure majority spin transmission in the range $-2.4 \text{ eV} < E - E_F < -1.1 \text{ eV}$ and in the P (AP) configuration almost

pure majority (minority) spin transmission in the range $-0.1 \text{ eV} < E - E_F < 0.6$. At E_F the TMR is 10% and at $E_F - 2.0 \text{ eV}$ the highest TMR of 600% is obtained. The Fe/Cd₃ZnS₄/Fe MTJ shows in the P configuration almost pure majority spin transmission in the range $-0.5 \text{ eV} < E - E_F < 0.0 \text{ eV}$. At E_F the TMR is 40% and at $E_F - 0.9 \text{ eV}$ the highest TMR of 1000% is obtained. The Zn₃CdS₄ MTJ shows in the P configuration almost pure majority spin transmission in the range $-2.0 \text{ eV} < E - E_F < -1.0 \text{ eV}$, in the AP configuration almost pure minority spin transmission in the range $-1.0 \text{ eV} < E - E_F < -0.5 \text{ eV}$, and in both the P and AP configurations almost pure majority spin transmission in the range $0.0 \text{ eV} < E - E_F < 0.3 \text{ eV}$. At E_F the TMR is 110% and at $E_F - 1.5 \text{ eV}$ the highest TMR of 600% is obtained.

The spin-polarized current $I^\uparrow - I^\downarrow$ is shown in Fig. 5 for both the P and AP configurations in comparison to that of a Fe/MgO/Fe MTJ with a barrier thickness of 15.44 Å. At $V = 2 \text{ V}$, for



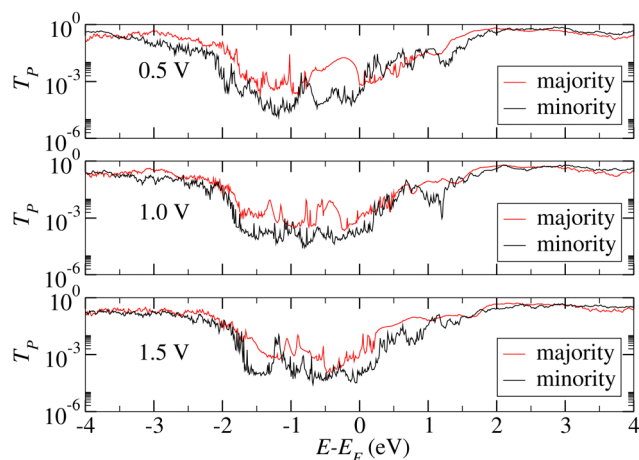


Fig. 6 Bias-dependence of the transmission of the Fe/Cd₃ZnS₄/Fe MTJ in the P configuration.

example, the spin-polarized current density of the P configuration amounts to 3.0×10^8 A cm⁻² for the Fe/CdS/Fe MTJ, 6.0×10^6 A cm⁻² for the Fe/ZnS/Fe MTJ, 4.4×10^8 A cm⁻² for the Fe/Cd₃ZnS₄/Fe MTJ, and 5.0×10^7 A cm⁻² for the Fe/Zn₃CdS₄/Fe MTJ, exceeding the value of 2.4×10^6 A cm⁻² obtained for the Fe/MgO/Fe MTJ by up to two orders of magnitude. The bias-dependence of the transmission of the Fe/Cd₃ZnS₄/Fe MTJ with the highest spin-polarized current density of the P configuration is shown in Fig. 6.

IV. Conclusion

We have explored the possibility of using the opto-electronic materials CdS, ZnS, Cd₃ZnS₄, and Zn₃CdS₄ as tunnel barriers in next generation MTJs. Fe-based MTJs adopting these semiconductors are found to have excellent potential in hybrid memory/logic components, displays with large viewing angle, and quantum-bit manipulators, for example, due to the provided high spin-polarized current. We have identified the areas of the two-dimensional BZ where $\kappa(E_F, \mathbf{k}_{\parallel})$ is minimal and $T(E_F)$ therefore is maximal, finding in each case dominant transport at the Γ -point, where the valence band edges are dominated by p_z orbitals and the conduction band edges are dominated by s orbitals. By complex band analysis we have established Δ_1 symmetry filtering of the Bloch states at the Γ -point. The highest TMR is found to be 5000% at $E_F + 0.1$ eV for the Fe/CdS/Fe MTJ, 600% at $E_F - 2.0$ eV for the Fe/ZnS/Fe MTJ, 1000% at $E_F - 0.9$ eV for the Fe/Cd₃ZnS₄/Fe MTJ, and 600% at $E_F - 1.5$ eV for the Fe/Zn₃CdS₄/Fe MTJ.

Data availability

The article includes all the data generated in this study.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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