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# Edge-controlled half-metallic ferromagnetism and direct-gap semiconductivity in ZrS<sub>2</sub> nanoribbons

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The electronic and magnetic properties of ZrS<sub>2</sub> nanoribbons (NRs) are investigated based on first principles calculations. It is found that the ZrS<sub>2</sub> NRs with armchair edges are all indirect band gap semiconductors without magnetism, and the band gap exhibits odd–even oscillation behavior with the increase of the ribbon width. For the NRs with zigzag edges, those with both edges S-terminated are nonmagnetic direct band gap semiconductors, and the gap decreases monotonically as a function of the ribbon width. However, the NRs with one edge S-terminated and the other edge Zr-terminated are ferromagnetic half-metals, while those with both edges Zr-terminated tend to be ferromagnetic half-metals when the width  $N \geq 9$ . The magnetism of both systems mainly originates from the unsaturated edge Zr atoms. Depending on the different edge configurations and ribbon widths, the ZrS<sub>2</sub> NRs exhibit versatile electronic and magnetic properties, making them promising candidates for applications in electronics and spintronics.

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## 1 Introduction

The electronic and magnetic properties of nanoscale materials have been the subject of extensive research due to their potential applications in electronics and spintronics. Carbon-based systems, graphene for example, are among the most studied low-dimensional materials. Graphene has a high mobility at room temperature, making it a promising candidate for future information technology. However, the lack of an intrinsic band gap has limited its practical applications. Cutting two-dimensional (2D) graphene into one-dimensional (1D) nanoribbons (NRs) can open the band gap in graphene, which was first predicted theoretically<sup>1</sup> and then verified by experiments.<sup>2</sup> Because of the additional edge states unique to the dimensionality, 1D NRs could exhibit rich properties which can be further tuned by modifying their edges. It was found that the narrow zigzag graphene NR is semiconducting with the two edges antiferromagnetically coupled to each other.<sup>1</sup> In addition, the graphene NR can be converted into a half-metal in different ways, such as by applying a homogeneous electrical field<sup>3</sup> or by chemical decoration at the edges.<sup>4,5</sup>

Besides graphene, low-dimensional transition metal dichalcogenides, TMDs (with the formula MX<sub>2</sub>, where M = transition

metal and X = S, Se or Te), are another important material that has received considerable attention. Different from graphene, single layer MX<sub>2</sub> has three atomic layers, with an M-layer sandwiched between two X-layers. As a typical representative of TMDs, single layer MoS<sub>2</sub> is a direct band gap semiconductor.<sup>6,7</sup> An intrinsic 2D MoS<sub>2</sub> monolayer is nonmagnetic. However, distinct electronic and magnetic properties were reported for 1D MoS<sub>2</sub> NRs, that is, armchair MoS<sub>2</sub> NRs are nonmagnetic semiconductors, while zigzag MoS<sub>2</sub> NRs have ferromagnetic (FM) and metallic ground states.<sup>8,9</sup> WS<sub>2</sub> NRs exhibit similar properties to MoS<sub>2</sub> NRs.<sup>10</sup> The zigzag edge-related ferromagnetism was then observed experimentally in MoS<sub>2</sub> (ref. 11–14) and WS<sub>2</sub> (ref. 14 and 15) nanosheets. Furthermore, the electronic and magnetic properties of MoS<sub>2</sub> NRs can be modified by edge passivation<sup>16</sup> or by applying an external strain and/or electric field.<sup>17–20</sup> It was theoretically predicted that by applying a transverse electrical field, the insulator–metal transition occurs and ferromagnetism emerges beyond a critical electric field in the armchair MoS<sub>2</sub> NRs.<sup>19</sup>

Both MoS<sub>2</sub> and WS<sub>2</sub> crystallize in the honeycomb (H) structure. ZrS<sub>2</sub> is another kind of TMD, which crystallizes in the centered honeycomb (T) structure. Different from the MoS<sub>2</sub> monolayer, single layer ZrS<sub>2</sub> is an indirect band gap semiconductor.<sup>21</sup> In experiments, 1D ZrS<sub>2</sub> NRs have been synthesized by the process of chemical vapor transport and vacuum pyrolysis.<sup>22,23</sup> But what different properties do the 1D NRs present? In the present work, we investigate the electronic and magnetic properties of a series of ZrS<sub>2</sub> NRs. The results show that depending on the different edge configurations, the ZrS<sub>2</sub> NRs can be indirect band gap semiconductors, direct band gap

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semiconductors, antiferromagnetic (AFM) metals or FM half-metals, exhibiting versatile electronic and magnetic properties.

## 2 Computational details

Our calculations were performed within the framework of density functional theory (DFT),<sup>24</sup> as implemented in the QUANTUM ESPRESSO code.<sup>25</sup> The exchange correlation energy was in the form of the Perdew–Burke–Ernzerhof (PBE) functional<sup>26</sup> with the generalized gradient approximation (GGA). The Brillouin zones were sampled with  $12 \times 1 \times 1$  and  $20 \times 1 \times 1$  Monkhorst–Pack  $k$  meshes for the armchair and zigzag NRs, respectively. The cutoff energy for the plane-wave expansion was set to 40 Ry. Both the  $k$  meshes and cutoff energy were carefully tested so that the electronic structures were converged. The distance between each NR and its periodic image was set to be larger than 10 Å so that they can be treated as independent entities.

## 3 Results and discussion

The  $\text{ZrS}_2$  NRs can be obtained by cutting the monolayer. There are mainly two kinds of  $\text{ZrS}_2$  NR, *i.e.* armchair and zigzag NRs, according to the cutting directions with respect to the monolayer. For the zigzag  $\text{ZrS}_2$  NRs, the edges can be terminated by Zr or S atoms, thus three cases exist. We denote the case in which the two edges are both S atom-terminated as a 1S–1S zigzag NR, and the other two cases are 1Zr–1S (where one edge is a Zr atom and the other edge is an S atom) and 1Zr–1Zr (where both edges are Zr atoms) zigzag NRs. The relaxed structures of the four kinds of  $\text{ZrS}_2$  NR are demonstrated in Fig. 1, where the left and right panels show top and side views, respectively. The width of the NR is defined according to the number of Zr atoms across the ribbon width, as shown in Fig. 1(a) and (c). Compared with the initial structures, the fully relaxed structures of the NRs change slightly. This is mainly due to the edge atoms, as the edge S atoms tend to migrate away from the inner atoms, while the Zr atoms prefer to approach the inner side.

To investigate the stability of the  $\text{ZrS}_2$  NRs, we calculated the binding energy,  $E_b$ , which is defined as  $E_b = (E_{\text{Zr}_n\text{S}_m} - nE_{\text{Zr}} - mE_{\text{S}})/(n + m)$ , where  $E_{\text{Zr}_n\text{S}_m}$ ,  $E_{\text{Zr}}$  and  $E_{\text{S}}$  are the total energies of the  $\text{ZrS}_2$  NRs and the Zr and S atoms, respectively. The larger the binding energy, the more stable the corresponding structure. The NRs with the width  $N = 4$ –14 are considered in this work. The calculated binding energies of the four kinds of  $\text{ZrS}_2$  NR are in the range of 6.19–6.66 eV per atom, indicating that all of the investigated NRs are very stable. Moreover, as shown in Fig. 2, the binding energies of the 1S–1S zigzag NRs are the highest among the four kinds of NR, indicating that this type of NR is the most stable, followed by the armchair NRs, and the 1S–1Zr and 1Zr–1Zr zigzag NRs are relatively less stable. For each kind of NR, the system becomes more and more stable as the ribbon width increases.

To check the possibility of magnetism in  $\text{ZrS}_2$  NRs, both spin-unpolarized and spin-polarized calculations were carried out. For the 1S–1Zr and 1Zr–1Zr zigzag NRs, the total energies of the FM states are lower than those of the nonmagnetic states,

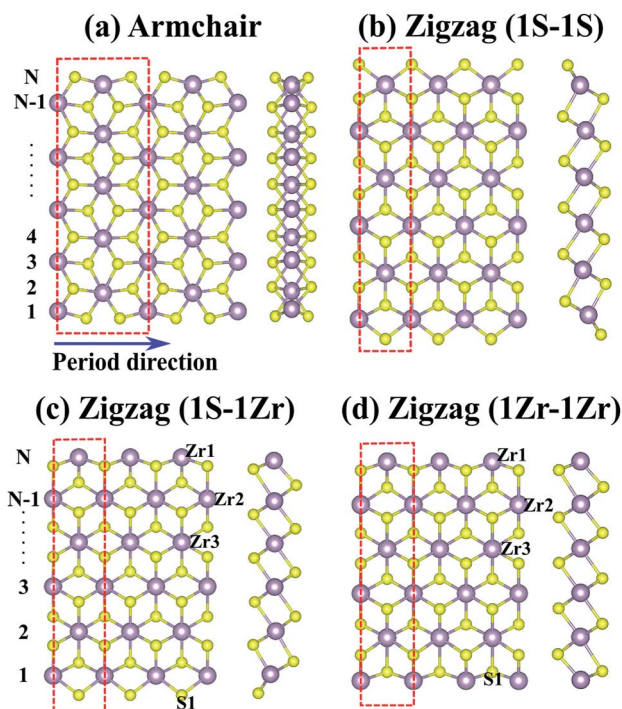


Fig. 1 Top and side views of the four different configurations of  $\text{ZrS}_2$  NRs: (a) armchair, (b) 1S–1S zigzag, (c) 1S–1Zr zigzag and (d) 1Zr–1Zr zigzag NRs. The red dashed squares indicate the unit cell.

indicating that these two kinds of NR have a magnetic ground state. However, the other two cases, *i.e.* the armchair and 1S–1S zigzag NRs, are nonmagnetic. Firstly, we focus on the electronic properties of the nonmagnetic armchair and 1S–1S zigzag  $\text{ZrS}_2$  NRs. The band structures show that all the investigated armchair  $\text{ZrS}_2$  NRs are indirect band gap semiconductors (see Fig. 3(a)). The band gap exhibits interesting odd–even oscillation as the ribbon width increases (Fig. 3(d)), that is, the even-numbered NRs have relatively larger band gaps than the neighboring odd-numbered NRs. Moreover, the band gap of the

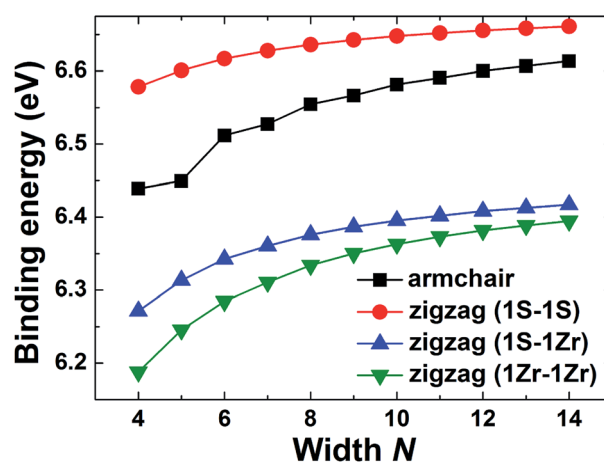


Fig. 2 Binding energies of the four kinds of  $\text{ZrS}_2$  NR as a function of ribbon width.



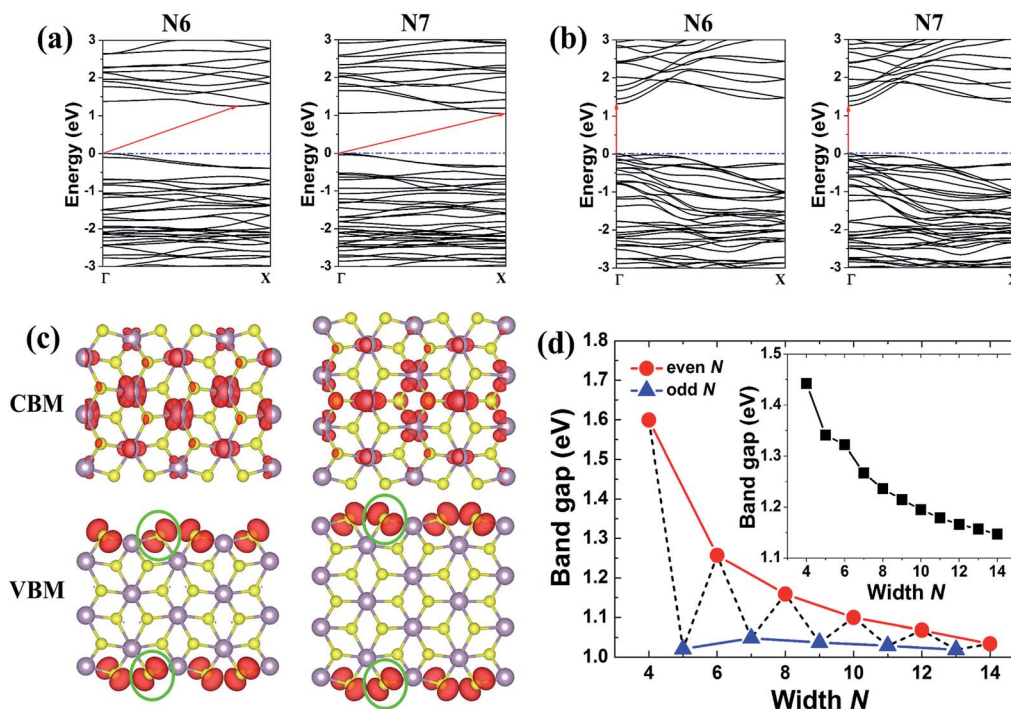


Fig. 3 Band structures of the two kinds of semiconducting  $\text{ZrS}_2$  NRs: (a) armchair and (b) 1S–1S zigzag NRs with widths  $N = 6$  and 7. (c) The band-decomposed charge densities for the conduction band minimum (CBM) and valence band maximum (VBM) of the armchair NRs with width  $N = 6$  (left panel) and  $N = 7$  (right panel). (d) Width-dependent band gap of the armchair  $\text{ZrS}_2$  NRs. The inset is the result for the 1S–1S zigzag NRs.

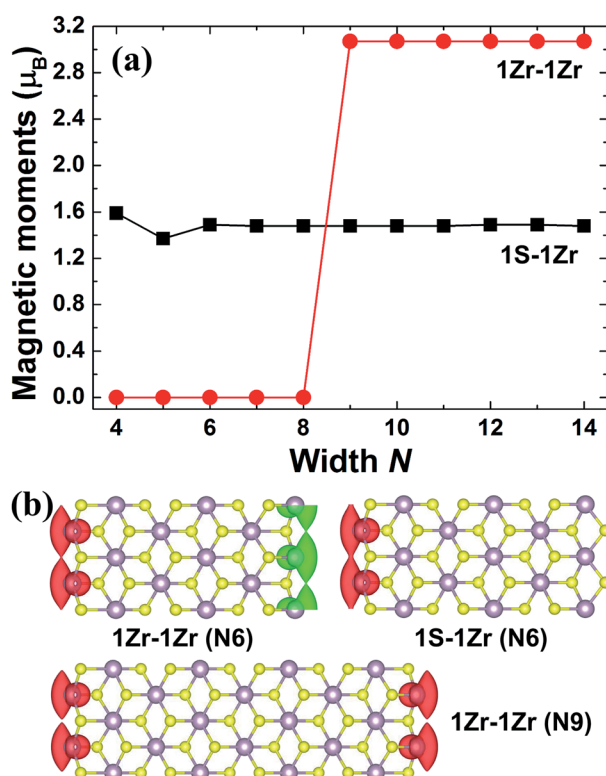


Fig. 4 (a) Total magnetic moments of the 1S–1Zr and 1Zr–1Zr zigzag  $\text{ZrS}_2$  NRs. (b) Spin densities ( $\rho_{\uparrow} - \rho_{\downarrow}$ ) for 1Zr–1Zr ( $N = 6$ ), 1S–1Zr ( $N = 6$ ) and 1Zr–1Zr ( $N = 9$ ) NRs.

even-numbered NRs decreases as a function of the width, while it changes slightly for the odd-numbered NRs. On the other hand, all the 1S–1S zigzag NRs are semiconducting as well but have direct band gaps (Fig. 3(b)). The band gap decreases monotonically as the ribbon width increases (see the inset of Fig. 3(d)). Although the  $\text{ZrS}_2$  monolayer is an indirect band gap semiconductor,<sup>21</sup> which makes it investigated less than  $\text{MoS}_2$ , the indirect-direct band gap transition can be obtained by cutting it into a 1D zigzag NR with both edges terminated by S atoms.

To investigate the origin of the odd–even oscillation of the band gap, we plot in Fig. 3(c) the band-decomposed charge densities for armchair  $\text{ZrS}_2$  NRs with widths  $N = 6$  and 7. We can see from Fig. 3(a) that for the armchair NRs, the valence band maximum (VBM) locates at the  $\Gamma$  point for both the even- and odd-numbered NRs, while the conduction band minimum (CBM) locates at the X points for the odd-numbered NRs and at the position between the  $\Gamma$  and X points for the even-numbered ones. The corresponding charge densities for the CBM and VBM are plotted in the upper and lower panels in Fig. 3(c), respectively. We can see that the CBM is mainly determined by the Zr atoms located in the inner part of the NRs, and there are much smaller contributions coming from the edge atoms. Therefore, the edge configurations have little impact on the electronic properties of the CBM. In contrast, the VBM is mainly controlled by the S atoms at the edge of the NRs. The structures of the odd-numbered NRs are symmetric with respect to the central line, thus the two nearest-neighboring S atoms at the two edges (which are circled by green lines) are just opposite



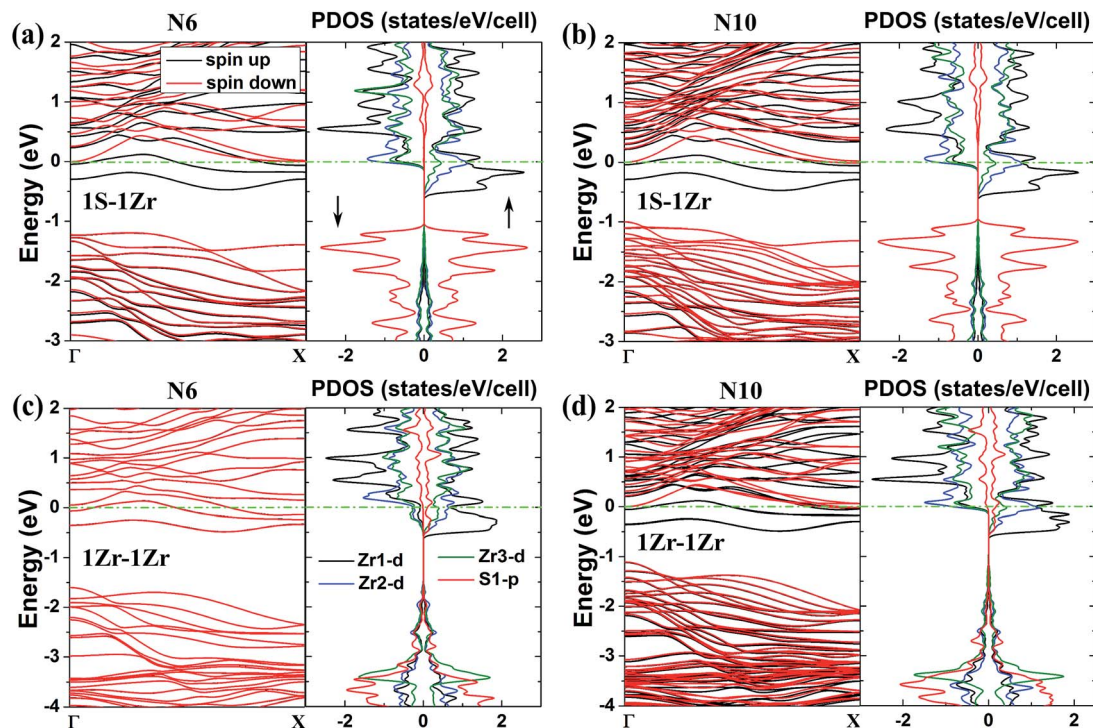


Fig. 5 Band structures and partial density of states (PDOS) of the two kinds of magnetic  $\text{ZrS}_2$  NR: (a) 1S-1Zr ( $N = 6$ ), (b) 1S-1Zr ( $N = 10$ ), (c) 1Zr-1Zr ( $N = 6$ ) and (d) 1Zr-1Zr ( $N = 10$ ) NRs. The black and red lines denote the spin-up and spin-down states, respectively. For the PDOS, the results of four kinds of atom (denoted in Fig. 1(c) and (d)), the edge S atom (S1), edge Zr atom (Zr1) and two inner Zr atoms near the edge (Zr2 and Zr3), are shown.

each other, making their distance shortest for the two edge atoms. However, for the even-numbered NRs, the two nearest-neighbor S atoms at the two opposite edges are staggered. Their distance is therefore enlarged, making interactions between the two edges relatively smaller. As a result, the VBM is lowered for the even-numbered NRs and their band gaps are enlarged accordingly compared with those of the neighboring odd-numbered NRs.

As discussed above, the other two kinds of  $\text{ZrS}_2$  NR, *i.e.* 1S-1Zr and 1Zr-1Zr zigzag NRs, are magnetic. The results of the spin density calculations (Fig. 4(b)) demonstrate that the magnetism of both systems mainly originates from the unsaturated edge Zr atoms. This is different from the H-structured zigzag  $\text{MoS}_2$  NRs, whose magnetic moments are concentrated on both the edge Mo and S atoms.<sup>8</sup> For narrow graphene NRs, the spins of the two edges are antiparallel to each other, *i.e.* the two edges are antiferromagnetically coupled.<sup>1</sup> Since the 1Zr-1Zr  $\text{ZrS}_2$  NRs have both edges terminated by Zr atoms, we also investigate the case in which the Zr atoms at the two edges are antiferromagnetically coupled. When the width  $N = 4-8$ , the energies of the AFM states are relatively smaller than those of the FM states, thus the ground states of the NRs are AFM. When the width increases up to 9, the system tends to be FM. A similar property was also observed in zigzag graphene NRs, for which switching from AFM to FM states may occur when the ribbon width is larger than 7 nm, due to FM inter-edge coupling.<sup>27,28</sup> For both the 1S-1Zr and 1Zr-1Zr zigzag NRs, the magnetic moments for the FM states are nearly width-independent, as

shown in Fig. 4(a). Since both edges of the 1Zr-1Zr NR are terminated by Zr atoms, while for the 1S-1Zr NR there is only one edge of Zr atoms, the total magnetic moments of the 1Zr-1Zr NRs are twice those of the 1S-1Zr systems.

In Fig. 5, we plot the band structures and partial density of states (PDOS) for the 1S-1Zr and 1Zr-1Zr zigzag NRs with ribbon widths  $N = 6$  and 10. The spin-up and spin-down channels are drawn in black and red lines, respectively. For the PDOS, the results of four kinds of atoms (denoted in Fig. 1(c) and (d)), including the edge S atom (S1), edge Zr atom (Zr1) and two inner Zr atoms near the edge (Zr2 and Zr3), are shown. For the 1S-1Zr  $\text{ZrS}_2$  NRs (Fig. 5(a) and (b)), the spin-up channels are metallic while the spin-down channels are semiconducting, thus the NRs are 100% spin-polarized and the systems are FM half-metals. The results of the PDOS show that for S1, the spin-up and spin-down channels are symmetric, so it has no contribution to the magnetic moment. For Zr1, Zr2 and Zr3, the spin-up and spin-down channels are asymmetric. The edge Zr atom (Zr1) contributes most to the magnetic moment, and from the edge to the inner sites, the contribution becomes smaller and smaller. Therefore, the magnetic moments mainly originate from the unsaturated Zr atoms located at the edge, which is consistent with the results of the spin density calculations (Fig. 4(b)). For the 1Zr-1Zr  $\text{ZrS}_2$  NRs, both the spin-up and spin-down states are metallic when  $N \leq 8$  (Fig. 5(c)), thus the systems are AFM metals. However, the system tends to be a FM half-metal when  $N$  increases up to 9 and the magnetic moments mainly come from the Zr atoms at the two edges.



Compared with the H-structured MoS<sub>2</sub> NRs, the T-structured MX<sub>2</sub> NRs have been investigated less. Reyes-Retana *et al.* reported that zigzag NiSe<sub>2</sub> NRs are metallic and the armchair systems are semiconducting, but both of these are nonmagnetic.<sup>29</sup> Our results demonstrate that for the 1S–1Zr and 1Zr–1Zr zigzag ZrS<sub>2</sub> NRs, 1D electrical current with complete spin polarization can be realized along the Zr edges of the systems. The intrinsic half-metallicity predicted in the ZrS<sub>2</sub> NRs is highly desirable for applications in spintronics.

## 4 Conclusion

In conclusion, we have investigated the electronic and magnetic properties of ZrS<sub>2</sub> NRs. Armchair ZrS<sub>2</sub> NRs are indirect band gap semiconductors and the gap exhibits odd–even oscillation as the ribbon width increases. For zigzag ZrS<sub>2</sub> NRs, the 1S–1S NRs are direct band gap semiconductors and the gap decreases as the width increases. Both the armchair and 1S–1S zigzag ZrS<sub>2</sub> NRs are nonmagnetic. However, the 1S–1Zr and 1Zr–1Zr ( $N \geq 9$ ) zigzag NRs are found to be FM half-metals and the magnetism mainly comes from the edge Zr atoms. Our results indicate that by tuning the edge configurations, ZrS<sub>2</sub> NRs could exhibit rich electronic and magnetic properties. The properties of direct gap and half-metallic ferromagnetism make ZrS<sub>2</sub> NRs promising candidates for future applications in electronics and spintronics.

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