


 Cite this: *RSC Adv.*, 2025, 15, 1618

Atomic structures and electronic properties of different contact surfaces for $C_xF_y-SiO_2$ triboelectric nanogenerator based on first-principles investigations

 Baonan Jia, ^{*ab} Jingming Gao, ^b Jiaxiang Zhao, ^b Jiahe Liang, ^a Xinhui Zhang, ^c Wendong Xiao, ^a Xiaoning Guan ^{*b} and Pengfei Lu ^b

Modification of the dielectric friction layer materials is an ideal way to enhance the output performance of a triboelectric nanogenerator (TENG), but current research mostly focuses on the metal–polymer or metal– SiO_2 materials. In this work, we constructed different TENG models based on polymer $C_xF_y-SiO_2$ electret materials, and the electronic properties of the different contact surfaces were investigated using first principles. We found that the charge transfer in $C_xF_y-SiO_2$ materials occurred only at the contact interface, and it was partially affected by the terminal atoms near the SiO_2 interface. The charge transfer of the polymer C_xF_y , that was in contact with the O-terminated SiO_2 achieved a more satisfactory effect. Among them, the II- C_3F_6-O model exhibited the highest amount of charge transfer because of the better hybridization of II- C_3F_6 with the O atoms of SiO_2 layer. Our study showed that instead of adding different types of dielectric friction layers, varying the configurations of the same types of dielectric friction layers is an alternative way to regulate charge transfer. Furthermore, this strategy could provide new ideas for enhancing the performance of TENGs.

Received 12th December 2024

Accepted 6th January 2025

DOI: 10.1039/d4ra08732a

rsc.li/rsc-advances

Introduction

With the continuous advancement of technology, microelectronic devices are increasingly being applied in the fields of self-powered flexible electronics and sensing. However, improving the energy efficiency of these devices to meet the demands of long-term use remains a pressing issue that needs to be addressed.^{1–7} The invention of triboelectric nanogenerators (TENG) marked a significant milestone in the field of self-powered systems, such as multimodal sensors, wearable electronic devices, and real-time human health monitoring, offering a novel approach to efficiently harvest mechanical energy.^{8–10} Current triboelectric nanogenerators still face issues such as insufficient output performance and low energy conversion efficiency,^{11–14} which limit their application in high-power energy generation because the dielectric friction layers cannot store charge for extended periods.^{15–25}

In recent years, the performance enhancement of TENG has been focused on the modification of its dielectric friction layers.^{26–31} Owing to their excellent ability to store charge for long periods, electret materials have become emerging materials for dielectric friction layers. Common electret materials mainly include inorganic electret materials, such as silicon dioxide, and organic electret materials, such as various polymers. In 2018, Wu *et al.*³² first investigated the mechanism of metal–polymer contact charging electrification *via* first-principles calculations and found that the stress on the contact region has a significant effect on the charge transfer. Further, Wu *et al.*³³ investigated the charge transfer mechanism between metal and amorphous polymers and found that the modification of contact materials is an important method to improve the output power of TENG. Subsequently, Al–PE polymer,²⁶ Cu–PVDF polymer,³⁴ Au–PTFE polymer³⁵ and Au–Nylon polymer³⁵ have attracted widespread attention from researchers through first-principles investigations. In 2021, Antony *et al.*³⁶ studied the electronic charge transfer of the hydroxylated metal/ SiO_2 interface using density functional theory and found that the separation distance between the contact surfaces must be small enough to produce electronic states within the apparent insulating bandgap. Owing to the unique physicochemical properties of metals, researchers have also paid significant attention to metal– SiO_2 materials,^{37,38} such as Cu– SiO_2 , Au– SiO_2 , Pd– SiO_2 , Al– SiO_2 and Ag– SiO_2 . In summary, current

^aKey Laboratory of Knowledge Automation for Industrial Processes of Ministry of Education, School of Automation and Electrical Engineering, University of Science and Technology Beijing, Beijing 100083, China. E-mail: jiabaonan@163.com

^bState Key Laboratory of Information Photonics and Optical Communications, Beijing University of Posts and Telecommunications, Beijing 100876, China. E-mail: guanxn@bupt.edu.cn

^cSchool of Science, Xi'an University of Architecture and Technology, Xi'an 710055, China



research on dielectric friction layers mainly focus on single electret materials, such as metal–polymer and metal–SiO₂ materials; however, few studies are reported on polymer–SiO₂ composite electret materials, indicating that there is still room for enhancing the output performance of TENG0073.

On this basis, we constructed polymer–SiO₂ composite electret materials and modified the dielectric friction layers by changing the contact atoms and further altering the charge transfer. In this paper, interface models consisting of polymer C_xF_y ($x = 2, y = 4; x = 3, y = 6$) and SiO₂ layer were considered, and the effect of different contact atoms on the electron transfer was investigated by first-principles calculations. The II-C₃F₆-O model, which consists of II-C₃F₆ and O-terminated SiO₂, shows the best performance of electron transfer because II-C₃F₆ can hybridize more significantly with the O atoms of the SiO₂ layer. Our study suggests that for the design of TENG, in addition to considering the different types of dielectric friction layers, different configurations of the same types can also be considered to enhance the performance of TENG.

Computational details

All the calculations in this paper are based on the Density Functional Theory (DFT) and were performed using the Vienna *ab initio* Simulation Package (VASP) software.^{39,40} Density Functional Theory is a first principles-based theoretical method that can achieve rigorous transformation from many-body problems into single-body problems. The Projector Augmented Wave (PAW) method⁴¹ was used to describe the interactions between the core electrons and the remaining valence electrons. The Perdew–Burke–Ernzerhof (PBE) functional within the generalized gradient approximation (GGA)⁴² was used to describe the exchange correlations of the electrons.

VASP software was used to optimize the atomic structures and calculate the electronic properties of different contact models. The input energy was chosen to be 400 eV to ensure accuracy of calculations. For geometry optimization, all atomic positions and lattice structures were fully relaxed until convergence criteria were reached, with forces less than 0.05 eV per Å per atom and a total energy convergence criterion of 1.0×10^{-5} eV. For electronic properties, the same computational accuracy was used to obtain reliable results. According to previous studies,^{43,44} a vacuum layer of 20 Å was added to the z-axis to eliminate periodicity and avoid interaction between the two periodic units. In addition, considering the large contact models, we used a $10 \times 10 \times 1$ K-point mesh around the gamma point in reciprocal space. The interlayer interaction of all interface models was carried out using Grimme's D3 correction^{45,46} to address the deficiencies in van der Waals forces.

Results and discussion

To better study the microscopic properties of triboelectric nanogenerators, we constructed interface models consisting of polymer C_xF_y (where $x = 2, y = 4; x = 3, y = 6$) and SiO₂ materials.^{47,48} For the C₃F₆ polymer, according to the different atoms

of the main and branched chains of the carbon chain, there are three conformations at the contact interface: CF₂, CF₃ and C₂F₂ atoms, respectively. We constructed three types of contact surface models, which were named I-C₃F₆, II-C₃F₆, and III-C₃F₆, as shown in Fig. 1(a–c), respectively. For the C₂F₄ polymer, on the other hand, the main and branched chains of the carbon chain are the same and there is only one conformation of CF₂ atoms at the contact interface. Therefore, only one C₂F₄ contact surface model was constructed, as shown in Fig. 1(d). Similarly, based on the type of terminal atoms of SiO₂ at the contact interface, we divided them into three configurations, and the atoms at the contact interface were O, Si, and SiO atoms, which were named as SiO₂-O, SiO₂-Si and SiO₂-SiO, as shown in Fig. 1(e–g), respectively. Consequently, considering the different conformations of dielectric friction layers between the polymer C_xF_y and SiO₂ materials, we considered a total of 12 different contact surface structures to investigate related microscopic properties for the nanogenerators.

To determine the stability of the structure and the nature of interaction force between the two dielectric friction layers, the binding energies of interface models with different configurations were calculated as follows

$$E_b = (E_{C_xF_y-SiO_2} - E_{C_xF_y} - E_{SiO_2})/A \quad (1)$$

where $E_{C_xF_y-SiO_2}$, $E_{C_xF_y}$ and E_{SiO_2} denote the total energies of the interfacial model, C_xF_y is the dielectric friction layer and the SiO₂ dielectric friction layer, respectively. A denotes the interface area between the C_xF_y layer and SiO₂ layer of the metal–SiO₂ interface model. Table 1 demonstrates the number of atoms, lattice parameters, layer spacing, and binding energy of the different C_xF_y-SiO₂ interface models.

Analysis of the values in Table 1 indicates that when C₂F₄ and C₃F₆ are combined with SiO₂, the contact area of C₂F₄ is smaller than that of C₃F₆, while the interlayer thickness is larger

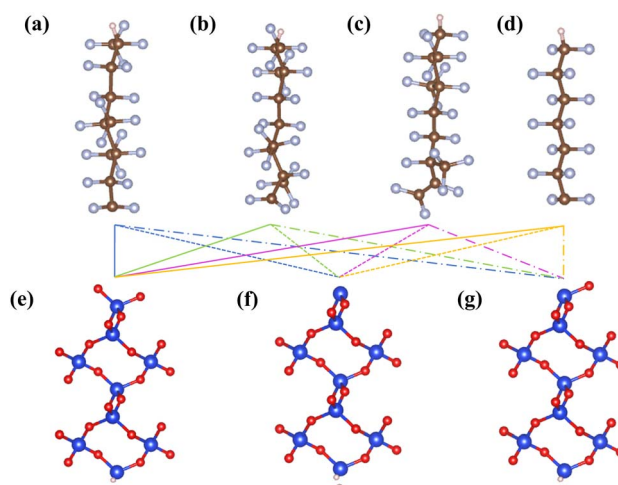


Fig. 1 Side-view geometries of the C_xF_y-SiO₂ interface model: (a–d) I-C₃F₆, II-C₃F₆, III-C₃F₆, and C₂F₄, respectively, and (e–g) O, Si, and SiO termination configurations of SiO₂, respectively. The red, blue, brown, gray, and orange spheres in the figure represent O, Si, C, F, and H atoms, respectively.



Table 1 Number of atoms, lattice parameters (a , b), layer spacing, and binding energies for the interfacial configurations of C_2F_4 , I- C_3F_6 , II- C_3F_6 , and III- C_3F_6 terminated with different atoms of SiO_2 , respectively

Material	Atomic number	$a/\text{\AA}$	$b/\text{\AA}$	Interlayer thickness/ \AA^2	Binding energy $eV \text{\AA}^{-2}$
C_2F_4 - SiO_2 -O	45	5.46	4.87	2.719	0.034
C_2F_4 - SiO_2 -Si	43	5.46	4.87	2.740	0.011
C_2F_4 - SiO_2 -SiO	44	5.46	4.87	2.369	0.053
I- C_3F_6 -O	54	6.37	4.89	2.024	0.038
I- C_3F_6 -Si	52	6.37	4.89	2.093	0.032
I- C_3F_6 -SiO	53	6.37	4.89	2.362	0.008
II- C_3F_6 -O	57	6.37	4.89	1.568	0.046
II- C_3F_6 -Si	55	6.37	4.89	1.378	0.024
II- C_3F_6 -SiO	56	6.37	4.89	2.038	0.012
III- C_3F_6 -O	56	6.37	4.89	1.118	0.019
III- C_3F_6 -Si	54	6.37	4.89	1.426	0.002
III- C_3F_6 -SiO	55	6.37	4.89	1.356	0.002

than that of C_3F_6 . Consequently, it is easier for C_3F_6 to form a contact interface model with SiO_2 than C_2F_4 . Among the interface models consisting of SiO_2 with different C_3F_6 configurations, the I- C_3F_6 model always has the largest interlayer thickness, whereas the II- C_3F_6 model has a suitable interlayer thickness. For the interface models consisting of C_3F_6 with different SiO_2 configurations, the binding energies of the C_3F_6 -O configurations are consistently larger than those of the C_3F_6 -Si configurations and larger than those of the C_3F_6 -SiO configurations. Based on previous studies,^{49,50} we screened the interface models with van der Waals interactions using 20–40 meV \AA^{-1} as the range of binding energies. When the polymer C_3F_6 layer contacted with SiO-terminated SiO_2 , the binding energy is less than 0.02 eV \AA^{-2} . When the SiO_2 layer contacted with II- C_3F_6 , the formed configuration has the largest binding

energy, of which the II- C_3F_6 -O configuration has the most suitable binding energy of 0.046 eV \AA^{-2} .

In order to better understand the output performance, we analyzed the plane-averaged charge density difference of these 12 interface models and attempted a preliminary screening based on the amount and direction of charge transfer between the contact surfaces. It can be inferred from Fig. 2 that when the polymer C_xF_y is contacted with O-terminated SiO_2 to form the C_xF_y -O interface model, the values of average charge density difference all reach up to $6 \times 10^{-3} e \text{\AA}^{-1}$. In contrast, when it is contacted with Si-terminated and SiO-terminated SiO_2 to form the C_xF_y -Si and C_xF_y -SiO interface models, respectively, the values are much less likely to reach the same level in both dielectric friction layers.

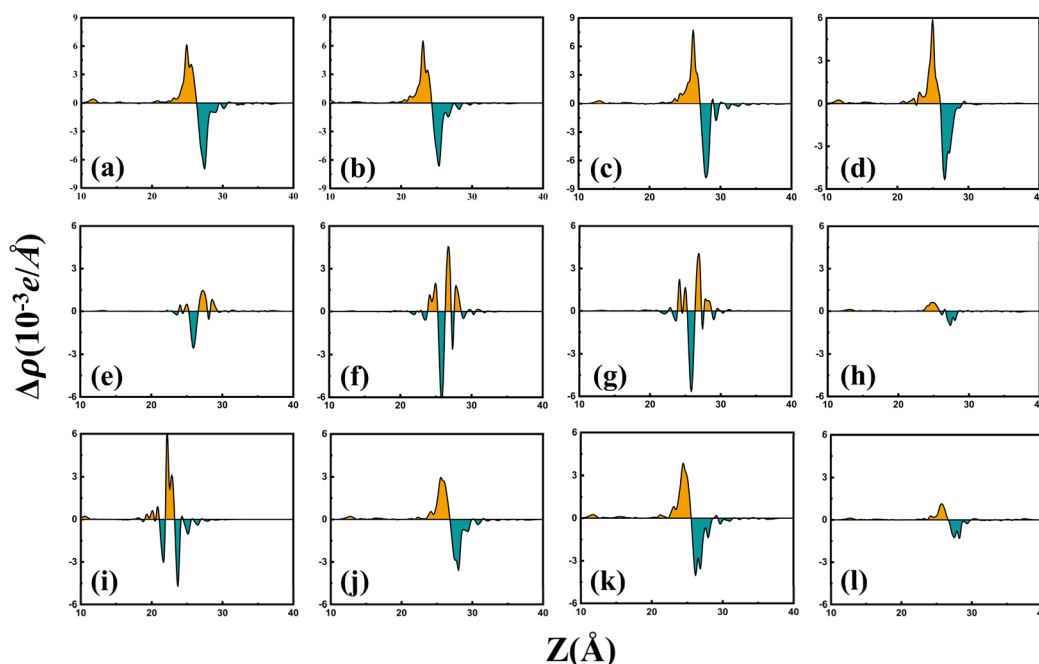


Fig. 2 Plane-averaged charge density difference at the interface of C_xF_y and SiO_2 : (a, e and i) C_2F_4 (b, f and j) I- C_3F_6 , (c, g and k) II- C_3F_6 , and (d, h and l) III- C_3F_6 , where (a–d) denote O-contacting surfaces, (e–h) denote Si-contacting surfaces, and (i–l) denote SiO-contacting surfaces.



For the C_xF_y-O , $I-C_3F_6-SiO$, and $II-C_3F_6-SiO$ configurations, as shown in Fig. 2(a)–(d), (j) and (k), the six configurations have similar charge transfer modes. The difference values between the C_xF_y and SiO_2 layers are almost equal, indicating a nearly unidirectional electron transfer. The configurations terminated with O atoms show a significantly larger charge transfer than those terminated with SiO atoms. In particular, the $II-C_3F_6-O$ configuration has better charge transfer capacity with a larger amount of charge transfer, and the values of the average charge density difference reach up to $7.6 \times 10^{-3} e \text{ \AA}^{-1}$, which represents an increase of 26.7% compared to the previously mentioned average value. For the C_xF_y-Si and C_2F_4-SiO configurations, as shown in Fig. 2(e)–(i), the frequent charge transfer at the interface implies that the electrons of these models do not simply transfer between the C_xF_y and SiO_2 layers; some electrons accumulate near the interface instead of moving unidirectionally. For the $III-C_3F_6-Si$ and $III-C_3F_6-SiO$ configurations, as shown in Fig. 2(h) and (i), the charge transfers are too small and not considered in the subsequent calculations. In summary, among the above 12 interface conformations, the $II-C_3F_6-O$ model has the most charge transfer and unidirectional electrons transfer.

In order to understand the effect of different polymer friction layers on the interfacial properties, the charge density difference $\Delta\rho$ at the interfaces was computed to further investigate the charge transfer and charge redistribution between the C_xF_y and SiO_2 interfaces.⁵¹

$$\Delta\rho = \rho_{C_xF_y-SiO_2} - \rho_{C_xF_y} - \rho_{SiO_2} \quad (2)$$

where $\rho_{C_xF_y-SiO_2}$ is the total charge density of the $C_xF_y-SiO_2$ interfacial system while $\rho_{C_xF_y}$ and ρ_{SiO_2} are the charge densities of the individual C_xF_y and SiO_2 layers, respectively. $\Delta\rho$ denotes the change in charge density due to the formation of chemical bonds between the C_xF_y and SiO_2 layers. Based on the analysis of the plane-averaged charge density difference, we identified 6

correlated structures with superior charge transfer properties, namely, C_2F_4-O , $I-C_3F_6-O$, $II-C_3F_6-O$, $III-C_3F_6-O$, $II-C_3F_6-Si$, $II-C_3F_6-SiO$, and further analyzed the charge transfer at the contact surfaces by the differential charge densities.

It can be inferred from Fig. 3 that the yellow area indicates the charge depletion region and the blue area indicates the charge aggregation region. Charge transfer at the contact interface can be visualized more intuitively through the differential charge density map. We can judge the level of charge density based on the size of the area where the charge accumulates. The charge aggregation and depletion mainly occur at the contact interface between the polymer C_xF_y and SiO_2 electret materials. There is more charge transfer occurring between the C_xF_y and O-terminated SiO_2 , no significant charge transfer between the C_xF_y and Si-terminated SiO_2 , and less charge transfer between the C_xF_y and SiO-terminated SiO_2 . For the polymer C_xF_y contacting the SiO_2-O interface, as shown in Fig. 3(a–d), C_xF_y acquires electrons from the SiO_2 interface, forming a charge depletion region at the SiO_2 layer and a charge aggregation region at the C_xF_y layer. As shown in Fig. 3(c), $II-C_3F_6$ contact with the SiO_2-O interface has a greater charge density. This indicates that the $II-C_3F_6-O$ model has a strong charge transfer capability as the electret materials of the triboelectric nanogenerator. For the polymer $II-C_3F_6$ contacting different termination atoms of SiO_2 , as shown in Fig. 3(c, e and f), based on the different contact surfaces as well as the different sizes of the charge transfer region, it can be further judged that the $II-C_3F_6-O$ structure has a better charge transfer ability and is more suitable to be the electret material for triboelectric nanogenerator.

The electrostatic potential difference between the two dielectric friction layers can be used to understand the amount of charge transfer and the time of storing charge. All configurations were analyzed by the value of differential charge density, and the variation of electrostatic potential along the z-direction was plotted, as shown in Fig. 4.⁵² The electrostatic potential

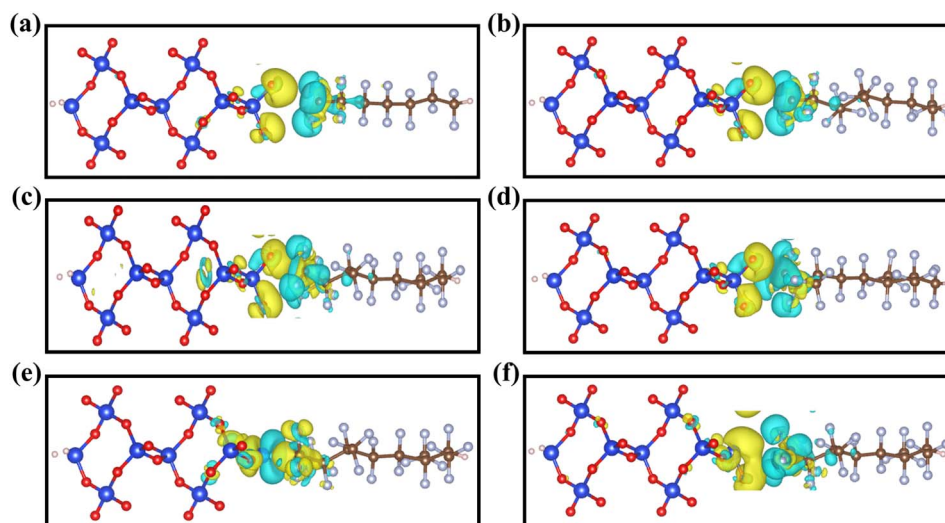


Fig. 3 Differential charge density of (a) C_2F_4-O , (b) $I-C_3F_6-O$, (c) $II-C_3F_6-O$, (d) $III-C_3F_6-O$, (e) $II-C_3F_6-Si$, and (f) $II-C_3F_6-SiO$. The blue and yellow regions indicate the electron depletion and accumulation regions, respectively. The size of the isosurface was set to $0.001 e \text{ \AA}^{-3}$.



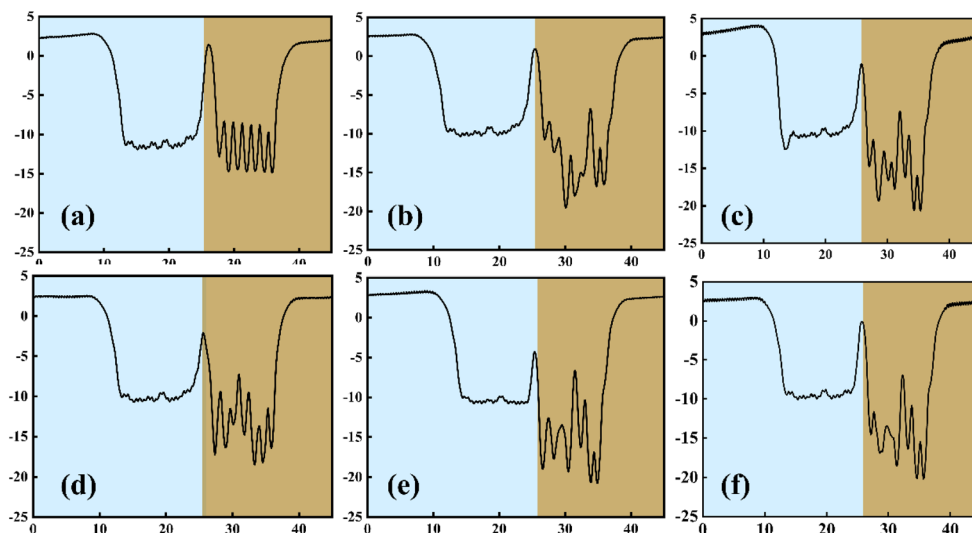


Fig. 4 Electrostatic potential plots of (a) C_2F_4-O , (b) $I-C_3F_6-O$, (c) $II-C_3F_6-O$, (d) $III-C_3F_6-O$, (e) $II-C_3F_6-Si$, and (f) $II-C_3F_6-SiO$, with the horizontal coordinates denoting the distances in the z -direction, and the vertical coordinates denoting the electrostatic potentials. The blue area in the figure indicates the SiO_2 layer and the brown area indicates the C_xF_y layer.

differences between C_xF_y and SiO_2-O , $II-C_3F_6$ and SiO_2-Si , $II-C_3F_6$ and SiO_2-SiO configurations were investigated. The electrostatic potential of the SiO_2 layer is shown on the left side of the figure and that of the polymer layer is on the right side. By quantitatively analyzing the electrostatic potential, for the configurations of the C_xF_y contacting the O-terminated SiO_2 , as shown in Fig. 4(a–d), the difference in electrostatic potentials of the C_3F_6-O interface is significantly larger than that of the C_2F_4-O interface, which indicated that the charge transfer between the dielectric friction layers of the C_3F_6 and SiO_2-O interfaces is closer, among which the $II-C_3F_6$ configuration performs

optimally. For the configurations of $II-C_3F_6$ contacting different termination atoms of SiO_2 , as shown in Fig. 4(c, e and f), they possess almost the same trend of electrostatic potential change at the same distance. However, the difference in the electrostatic potential between the $II-C_3F_6$ and SiO_2-O interfaces is the largest, as shown in Fig. 4(c), which indicates that the $II-C_3F_6-O$ model possesses more charge transfer and has better power generation performance.

In order to gain insight into the nature of charge transfer between the dielectric friction layers composed of polymer C_xF_y and SiO_2 electret materials, we calculated the total density of

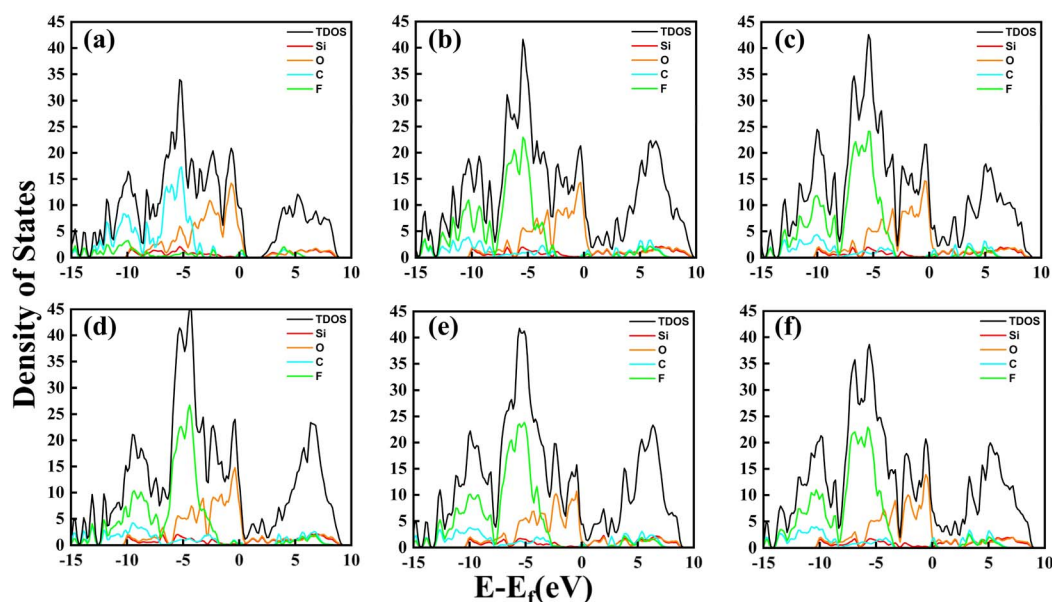


Fig. 5 PDOS plots of (a) C_2F_4-O , (b) $I-C_3F_6-O$, (c) $II-C_3F_6-O$, (d) $III-C_3F_6-O$, (e) $II-C_3F_6-Si$, and (f) $II-C_3F_6-SiO$. The black, red, orange, blue, and green colors in the plots indicate the orbital occupancy of TDOS, Si, O, C, and F, respectively.



states (TDOS) and the projected density of states (PDOS) of the atoms for these interface models, as shown in Fig. 5. Due to the difference in the C_2F_4 and C_3F_6 extra-nuclear electronic structures, the TDOS peaks of the C_2F_4 structure in Fig. 5(a) are smaller than those of the other C_3F_6 structures. Also, there is a region with no density of states near 1 eV that is also different from the other C_3F_6 structures, a characteristic that is not conducive to electron transfer between dielectric friction layers.

In the interface models, as shown in Fig. 5(b)–(d), one layer of the dielectric friction layers is O-terminated SiO_2 and the other layer contacts different C_3F_6 structures. It can be seen that above the Fermi energy level, for the I- C_3F_6 structure, the first peak of the O atom is close to the C and F atoms and shows hybridization in both contour and value. For the II- C_3F_6 structure, the hybridization of the O atom with C and F atom also changes obviously, and the change in the O atom is more obvious than that of the Si atom, which makes it reasonable to assume that the II- C_3F_6 -O structure has better orbital hybridization and charge transfer. However, for the III- C_3F_6 structure, the first peak does not show significant hybridization with O atoms, which shows that the charge transfer of the III-structure is not as good as that of the I- and II-structures.

In the interface models, as shown in Fig. 5(c), (e) and (f), one layer of dielectric friction layers is the polymer II- C_3F_6 material, the other layer contacts different termination atoms of SiO_2 . For the SiO_2 -O model, the first peak on the O, C and F atoms shows obvious hybridization above the Fermi energy level. For the SiO_2 -Si model, the first peak also shows obvious hybridization but is numerically smaller than the O-terminated SiO_2 model. For the SiO_2 -SiO model, the first peak has no obvious orbital hybridization compared with the other two models. Therefore, it is considered that the II- C_3F_6 -O interface model, constituted by II- C_3F_6 in contact with the O-terminated SiO_2 , has better output performance, which also explains the charge transfer phenomenon between the dielectric friction layers.

Conclusions

In this study, considering the different contact surfaces between polymer C_xF_y and SiO_2 electret materials, we constructed $C_3F_6(C_2F_4)$ - SiO_2 interface models as the dielectric friction layers. The charge transfer between the contact surfaces was investigated through first principles to study the intrinsic reasons of TENG modification. It was found that a better charge transfer effect was achieved when the polymer C_xF_y was in contact with the O-terminated SiO_2 . Among the interface models, the II- C_3F_6 configuration was likely to contact with the O atoms of the SiO_2 layer, which not only had the largest amount of charge transfer but also possessed unidirectional charge transfer. Notably, the II- C_3F_6 -O model possesses the largest electrostatic potential difference, indicating that this TENG structure has the best output performance. By calculating and analyzing the density of states, we found that the II- C_3F_6 configuration will hybridize more significantly with the O atoms of SiO_2 compared to the other polymer configurations. This study reveals the intrinsic properties of TENG composed of polymer C_xF_y and SiO_2 electret materials and provides

theoretical guidance for modifying dielectric friction layers through the different configurations of the same materials. How to improve the charge storage capability of the dielectric friction layers through material composition design is currently a hot topic of concern. In future, dielectric friction layers composed of composite electret materials need to be expanded.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Author contributions

Baonan Jia: writing – original draft, investigation, methodology, formal analysis. Jingming Gao: investigation, data curation, software. Jiaxiang Zhao: investigation, data curation, software. Guoying Qin: investigation, data curation, software. Xiaoning Guan: writing – review & editing, formal analysis. Xinhui Zhang: investigation, formal analysis. Wendong Xiao: writing – review & editing, conceptualization. Pengfei Lu: investigation, conceptualization.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Fundamental Research Funds for the Central Universities (No. 00007845), Foundation of Laboratory of Computational Physics (No. 6142A05QN22017) and project funded by China Postdoctoral Science Foundation (No. 2022M720516). We thank for the helpful discussion with Prof. Pengfei Guan and the computational support from the Beijing Computational Science Research Center (CSRC).

Notes and references

- 1 G. Ruhl, S. Wittmann, M. Koenig and D. Neumaier, *Beilstein J. Nanotechnol.*, 2017, **8**, 1056–1064.
- 2 J. Yang, K. Liu, X. Chen and D. Shen, *Prog. Polym. Sci.*, 2022, **83**, 100397.
- 3 A. I. Inamdar, S. Kamal, M. Usman, M.-H. Chiang and K.-L. Lu, *Coord. Chem. Rev.*, 2024, **502**, 215596.
- 4 X. Cao, Y. Xiong, J. Sun, X. Xie, Q. Sun and Z. L. Wang, *Nano-Micro Lett.*, 2023, **15**, 14.
- 5 X. Tao, X. Chen and Z. L. Wang, *Energy Environ. Sci.*, 2023, **16**, 3654–3678.
- 6 H. Xiang, L. Peng, Q. Yang, Z. L. Wang and X. Cao, *Sci. Adv.*, 2024, **10**, eads2291.
- 7 Y. Jiang, X. Liang, T. Jiang and Z. L. Wang, *Engineering*, 2024, **33**, 204–225.
- 8 F.-R. Fan, Z.-Q. Tian and Z. L. Wang, *Nano Energy*, 2012, **1**, 328–334.
- 9 J. Luo and Z. L. Wang, *EcoMat*, 2020, **2**, e12059.



- 10 J. Tao, L. Wang, K. Kong, M. Hu and Z. Dai, *Biomimetics*, 2022, **7**, 216.
- 11 Y. Li, Y. Luo, H. Deng, S. Shi, S. Tian, H. Wu, J. Tang, C. Zhang, X. Zhang and J. W. Zha, *Adv. Mater.*, 2024, **36**, 2314380.
- 12 X. Zhu, Y. Hao, J. Yang, W. Su, H. Zhang, Y. Qin, C. Zhang and X. Li, *Appl. Mater. Today*, 2024, **41**, 102492.
- 13 K. Xiao, W. Wang, K. Wang, H. Zhang, S. Dong and J. Li, *Adv. Funct. Mater.*, 2024, **34**, 2404744.
- 14 W. Peng and S. Du, *IEEE Trans. Circuits Syst. I Regul. Pap.*, 2023, **70**, 3049–3062.
- 15 W. Sun, G. Ji, J. Chen, D. Sui, J. Zhou and J. Huber, *Nano Energy*, 2023, **108**, 108248.
- 16 H. Zhang, P. Zhang, P. Li, L. Deng, W. Zhang, B. Liu and Z. Yang, *Nano Res.*, 2022, **15**, 7163–7171.
- 17 Q. M. Saqib, M. Y. Chougale, M. U. Khan, R. A. Shaikat, J. Kim, J. Bae, H. W. Lee, J.-I. Park, M. S. Kim and B. G. Lee, *Nano Energy*, 2021, **89**, 106458.
- 18 Y. Zhu, Y. Zhao, L. Hou and P. Zhang, *Micromachines*, 2022, **13**, 2053.
- 19 P. Zhang, Y. Ma, H. Zhang and L. Deng, *ACS Appl. Energy Mater.*, 2023, **6**, 6598–6606.
- 20 L. Shooshtari, N. Rafiefard, M. Barzegar, S. Fardindoost, A. Irajizad and R. Mohammadpour, *ACS Appl. Nano Mater.*, 2022, **5**, 17123–17132.
- 21 X. Liu, Y. Wang, G. Wang, Y. Ma, Z. Zheng, K. Fan, J. Liu, B. Zhou, G. Wang and Z. You, *Matter*, 2022, **5**, 4315–4331.
- 22 W. Liu and J. Shi, *Nano Energy*, 2021, **89**, 106479.
- 23 S.-N. Lai, C.-K. Chang, C.-S. Yang, C.-W. Su, C.-M. Leu, Y.-H. Chu, P.-W. Sha and J.-M. Wu, *Nano Energy*, 2019, **60**, 715–723.
- 24 Y. Liu, G. Liu, T. Bu and C. Zhang, *Mater. Today Energy*, 2021, **20**, 100686.
- 25 K. Xia and Z. Xu, *Smart Mater. Struct.*, 2020, **29**, 095016.
- 26 L. Li, X. Wang, P. Zhu, H. Li, F. Wang and J. Wu, *Nano Energy*, 2020, **70**, 104476.
- 27 X. Chen, F. Wang, Y. Zhao, P. Wu, L. Gao, C. Ouyang, Y. Yang and X. Mu, *Research*, 2022, **2022**, 9765634.
- 28 G. Yinben, C. Zixi, W. Hongzhi and Z. Qinghong, *J. Inorg. Mater.*, 2021, **36**, 919–928.
- 29 C. Xu, B. Zhang, A. C. Wang, W. Cai, Y. Zi, P. Feng and Z. L. Wang, *Adv. Funct. Mater.*, 2019, **29**, 1903142.
- 30 G. Khandelwal, A. Chandrasekhar, N. P. Maria Joseph Raj and S.-J. Kim, *Adv. Energy Mater.*, 2019, **9**, 1803581.
- 31 G. Khandelwal, M. K. Ediriweera, N. Kumari, N. P. Maria Joseph Raj, S. K. Cho and S.-J. Kim, *ACS Appl. Mater. Interfaces*, 2021, **13**, 18887–18896.
- 32 J. Wu, X. Wang, H. Li, F. Wang, W. Yang and Y. Hu, *Nano Energy*, 2018, **48**, 607–616.
- 33 J. Wu, X. Wang, H. Li, F. Wang and Y. Hu, *Nano Energy*, 2019, **63**, 103864.
- 34 L. Li, X. Wang, Y. Hu, Z. Li, C. Wang and Z. Zhao, *Adv. Funct. Mater.*, 2022, **32**, 2109949.
- 35 D. Kang, J.-H. Hwang, Y.-J. Kim, P. Zhao, H. Y. Lee, J. Kim, M. S. Shin, S. Jeon, S. Kim and S.-W. Kim, *Mater. Today*, 2024, **72**, 109–116.
- 36 A. C. Antony, D. Thelen, N. Zhelev, K. Adib and R. G. Manley, *J. Appl. Phys.*, 2021, **129**, 065304.
- 37 T.-R. Shan, B. D. Devine, S. R. Phillpot and S. B. Sinnott, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2011, **83**, 115327.
- 38 J. Huang, E. Tea, G. Li and C. Hin, *Appl. Surf. Sci.*, 2017, **406**, 128–135.
- 39 V. Wang, N. Xu, J.-C. Liu, G. Tang and W.-T. Geng, *Comput. Phys. Commun.*, 2021, **267**, 108033.
- 40 Y. Y. Liang, D. M. Chen and J. F. Tong, *Key Eng. Mater.*, 2012, **512**, 490–493.
- 41 P. E. Blöchl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953.
- 42 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- 43 F. Withers, O. Del Pozo-Zamudio, A. Mishchenko, A. P. Rooney, A. Gholinia, K. Watanabe, T. Taniguchi, S. J. Haigh, A. Geim and A. Tartakovskii, *Nat. Mater.*, 2015, **14**, 301–306.
- 44 A. K. Geim and I. V. Grigorieva, *Nature*, 2013, **499**, 419–425.
- 45 W. Liu, C.-L. Yang, Y.-T. Zhu and M.-S. Wang, *J. Phys. Chem. C*, 2008, **112**, 1803–1811.
- 46 R. Chawla and S. Sharma, *Compos. Sci. Technol.*, 2017, **144**, 169–177.
- 47 K. Zhao, W. Sun, S. Li, Z. Song, M. Zhong, D. Zhang, B.-N. Gu, M.-J. Liu, H. Fu and H. Liu, *Discover Nano*, 2023, **18**, 69.
- 48 D. L. Vu, C. D. Le and K. K. Ahn, *Polymers*, 2022, **14**, 960.
- 49 T. Björkman, A. Gulans, A. V. Krasheninnikov and R. M. Nieminen, *Phys. Rev. Lett.*, 2012, **108**, 235502.
- 50 Y. Chen, X. Guan, L. Yang, B. Jia, H. Zhao, L. Han, P. Guan and P. Lu, *Appl. Surf. Sci.*, 2023, **611**, 155679.
- 51 M. T. As-Samee, M. S. Hasan Khan, K. Kubra, M. R. Islam and M. T. Hasan, *AIP Adv.*, 2023, **13**, 065004.
- 52 Q. Campbell, *J. Electrochem. Soc.*, 2023, **170**, 031506.

