

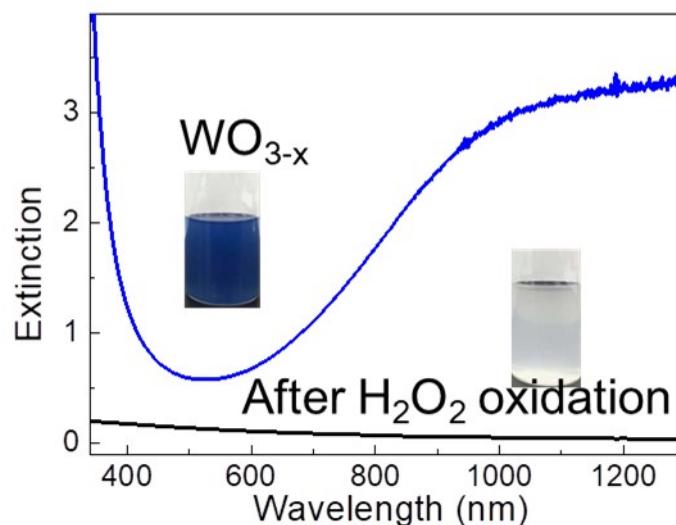
## Supplementary Material

# Self-Z-Scheme Plasmonic Tungsten Oxides Nanowires Boosting Ethanol Dehydrogenation under UV-Visible Light Irradiation

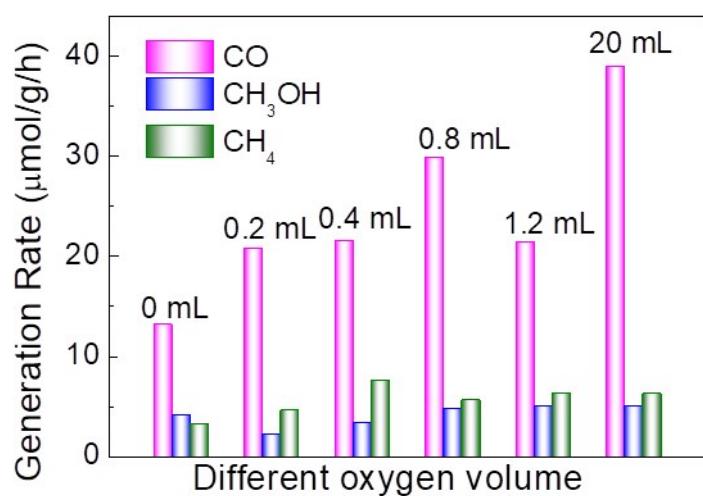
Changhai Lu,<sup># [a]</sup> Juan Li,<sup># [a]</sup> Guanying Chen,<sup>[a]</sup>, Baojun Li<sup>\*[a]</sup> and Zaizhu Lou,<sup>\*[a]</sup>

<sup>a</sup> Institute of Nanophotonics Jinan University Guangzhou 511443, China

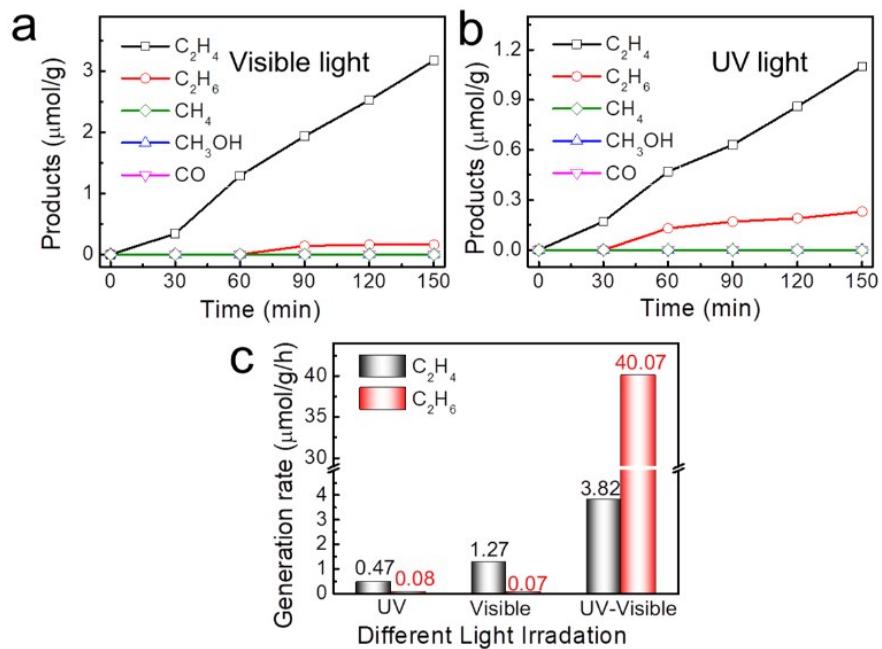
E-mail: [zzlou@jnu.edu.cn](mailto:zzlou@jnu.edu.cn); [baojunli@jnu.edu.cn](mailto:baojunli@jnu.edu.cn).



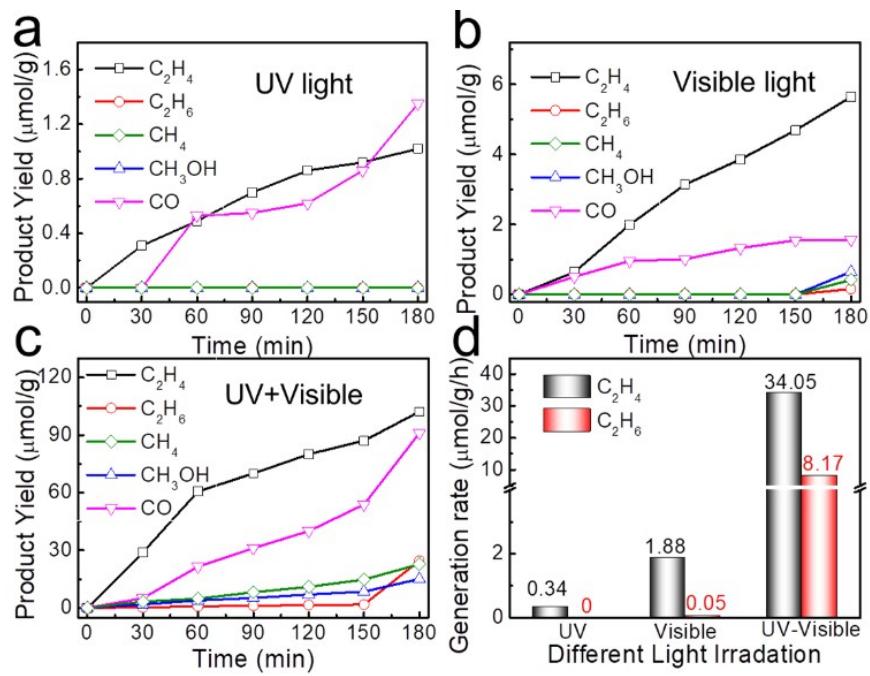
**Fig. S1.** Extinction spectra of plasmonic  $\text{WO}_{3-\text{x}}$  nanowire before and after  $\text{H}_2\text{O}_2$  oxidation, and their solution color



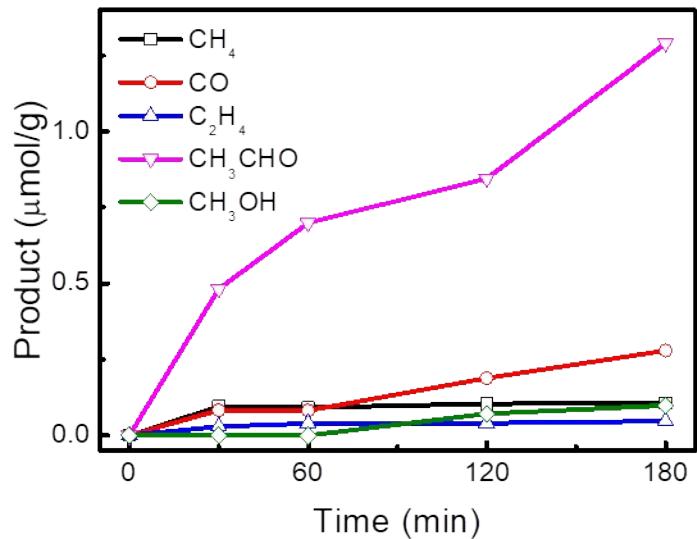
**Fig. S2.** Influence of oxygen on C1 compounds generation over plasmonic  $\text{WO}_{3-x}$  as photocatalysts under UV-visible light irradiation.



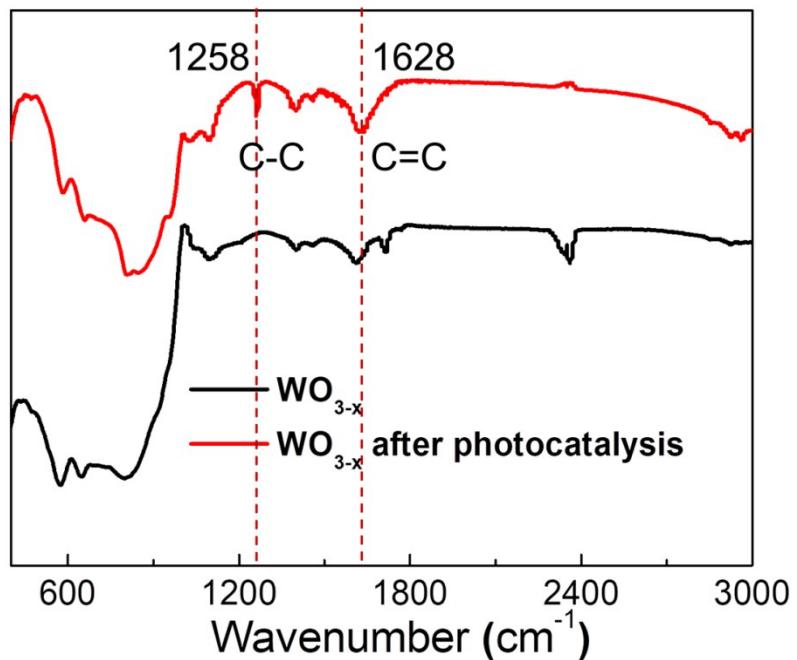
**Fig. S3 .** Products generation over plasmonic  $\text{WO}_{3-x}$  nanowire in nitrogen under visible (a), UV (b) and UV-visible, respectively. Ethylene and ethane generation rates (c) over plasmonic  $\text{WO}_{3-x}$  nanowire in nitrogen under visible, UV and UV-visible light irradiation, respectively.



**Fig. S4.** Products generation over plasmonic  $\text{WO}_{3-x}$  nanowire in optimal oxygen (0.4 mL) under UV (a), visible (b) and UV-visible (c) light irradiation, respectively. Ethylene and ethane generation rates (d) over plasmonic  $\text{WO}_{3-x}$  nanowire in optimal oxygen (0.4 mL) under visible, UV and UV-visible light irradiation, respectively.



**Fig. S5.** Products of methanol dehydrogenation reaction with absence of plasmonic  $\text{WO}_{3-\text{x}}$  under UV-visible light irradiation.



**Fig. S6.** IR spectra of plasmonic  $\text{WO}_{3-\text{x}}$  before and after photocatalysis.

Catalysts	T (K)	Products	Conversion (%)	Selectivity (%)	Byproducts	Ref.
Cu/C/SiO <sub>2</sub>	533	acetaldehyde	83.0	95.1	ethyl acetate, butanal, butanol	[1]
Cu/SiO <sub>2</sub>	533	acetaldehyde	85.4	79.4	ethyl acetate, butanal, butanol	[1]
MoO <sub>2</sub>	573	acetaldehyde, ethane	42.9	47.1, 45.6	ethylene	[2]
Cu/MC	553	acetaldehyde	73.0	94	ethyl acetate, acetone, acetone	[3]
Vo <sub>x</sub> /TiO <sub>2</sub> /SiO <sub>2</sub>	333	acetaldehyde	66.0	99	ethylene, carbon dioxide	[4]
Pt/TiO <sub>2</sub>	333	acetaldehyde	69.0	90	ethylene, carbon dioxide crotonaldehyde	[5]
<b>WO<sub>3-x</sub></b>	<b>303</b>	<b>acetaldehyde</b>	<b>84.2</b>	<b>91</b>	<b>ethylene, carbon monoxides</b>	<b>This work</b>

Table. S1 Comparison of the catalytic performance over WO<sub>3-x</sub> with the reported catalysts for the ethanol dehydrogenation

## References

- [1] Q. N. Wang, L. Shi, W. Li, W. C Li, R. Si, F. Schüthc, A. H. Lu, *Catal. Sci. Technol.*, **2018**, 8, 472-479.
- [2] Y. Nakamura, T. Murayama, W. Ueda, *ChemCatChem* , **2014**, 6, 741-744.
- [3] Q. N. Wang, L. Shi, A. H. Lu, *ChemCatChem*, **2015**, 7, 2846-2852.
- [4] D. Sannino, V. Vaiano, P. Ciambelli, G. Carotenuto, M. DiSerio, E. Santacesaria, *Catal. Today*, **2013**, 209, 159-163.
- [5] J. J. Murcia, M. C. Hidalgo, J. A. Navío, V. Vaiano, P. Ciambelli, D. Sannino, *Int. J. Photoenergy*, **2012**, 2012, 1-9.