

Electronic Supplementary Material (ESI) for Journal of Materials Chemistry A. This journal is © The Royal Society of Chemistry 2021

Supporting Information for

**Pt-on-Pd bimetallic nanodendrites stereoassembled on MXene  
nanosheets for use as high-efficiency electrocatalysts toward the  
methanol oxidation reaction**

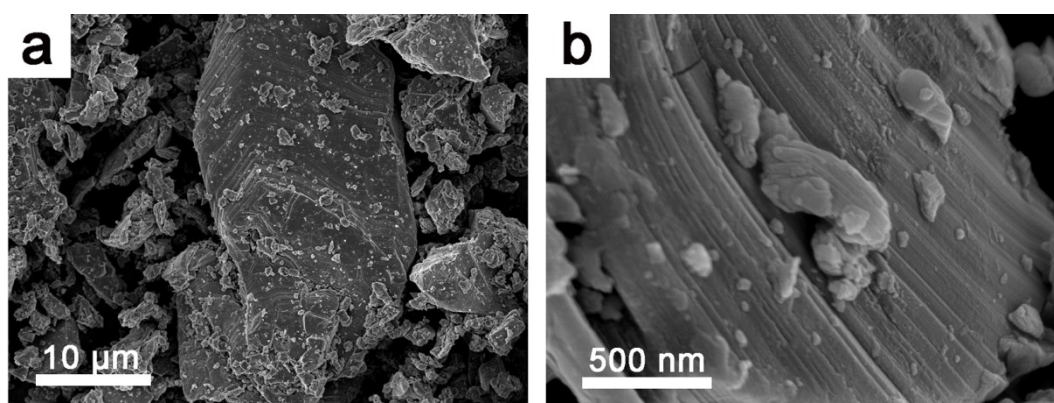
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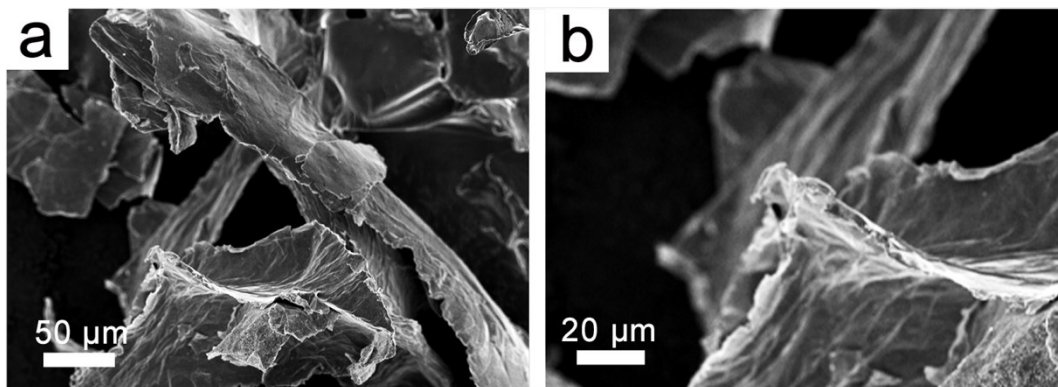
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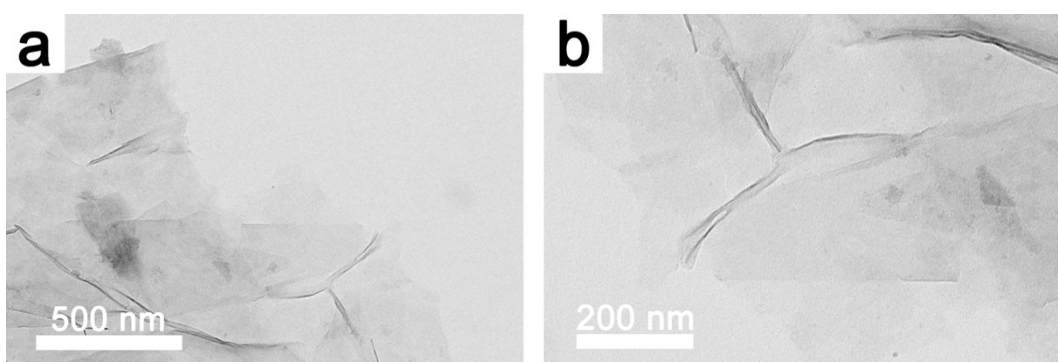
**Supplementary Results**



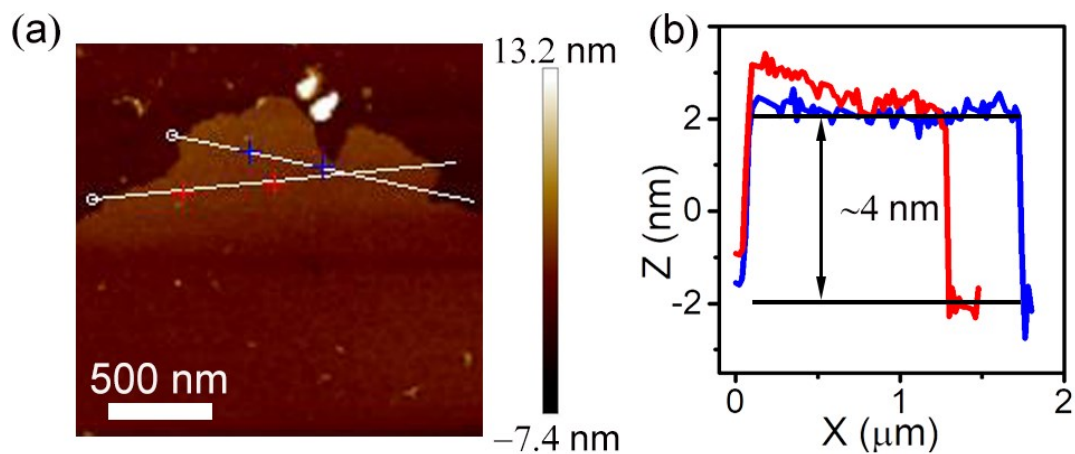
**Fig. S1** Representative SEM images of the bulk Ti<sub>3</sub>AlC<sub>2</sub> powder.



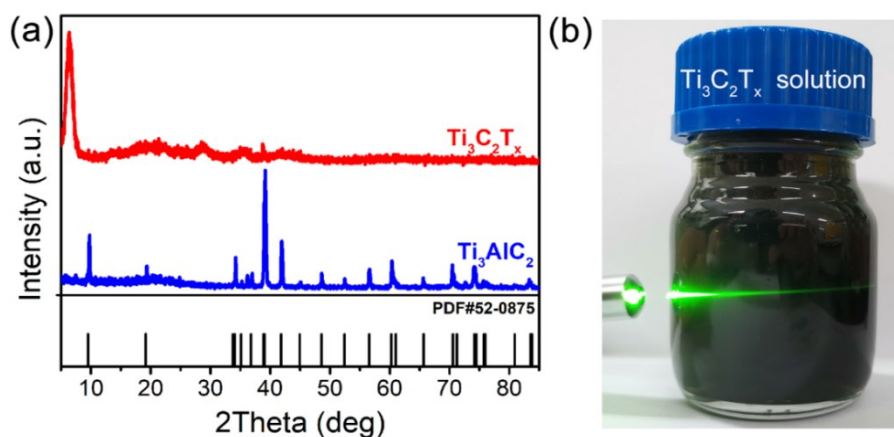
**Fig. S2** Representative SEM images of the exfoliated Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> nanosheets.



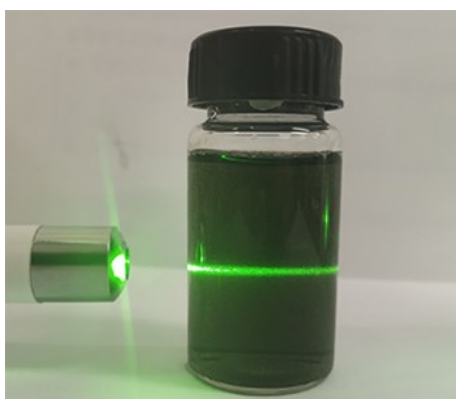
**Fig. S3** Representative TEM images of the exfoliated Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> nanosheets.



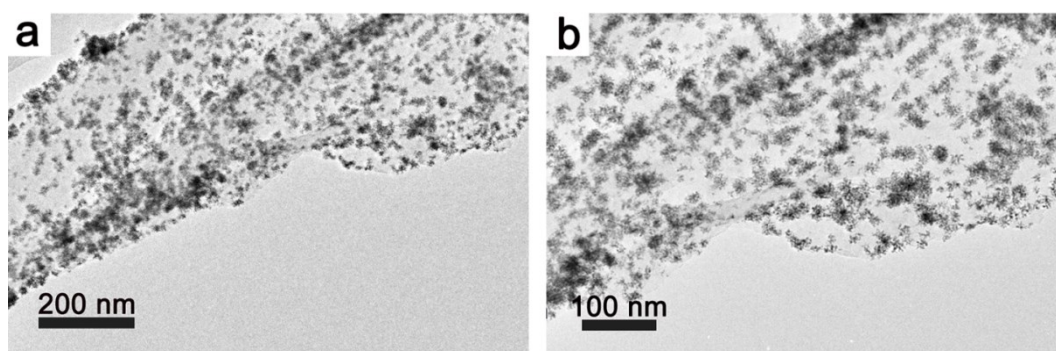
**Fig. S4** (a) AFM image of the exfoliated Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> nanosheets and (b) the corresponding thickness analysis along the white lines in (a) discloses that the uniform thickness of the Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> nanosheets is about 4.0 nm.



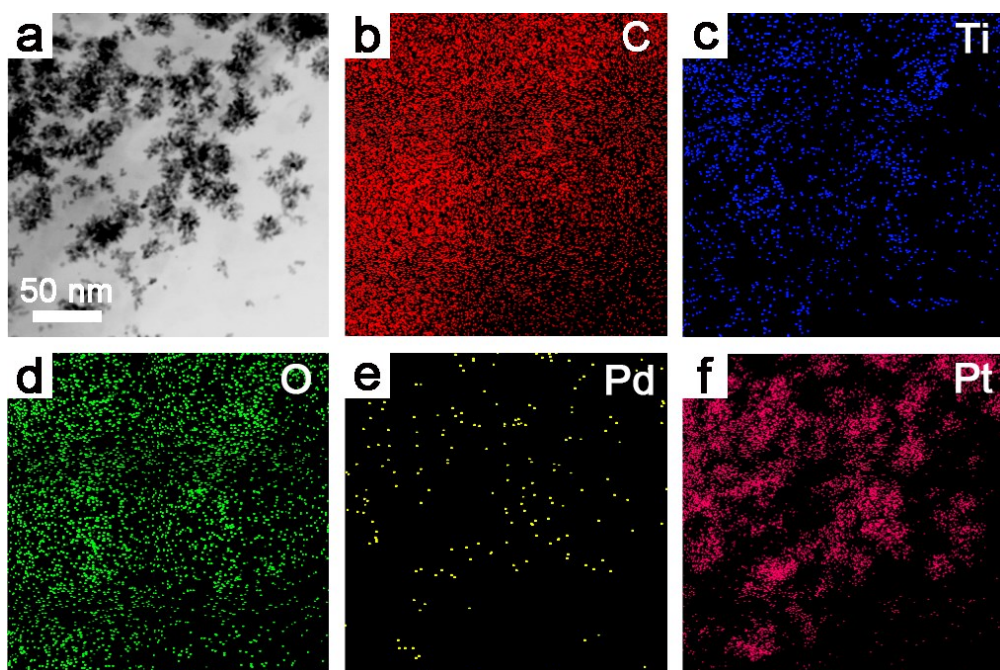
**Fig. S5** (a) The XRD patterns of the  $\text{Ti}_3\text{AlC}_2$  powder and  $\text{Ti}_3\text{C}_2\text{T}_x$  nanosheets; (b) The digital photographs of  $\text{Ti}_3\text{C}_2\text{T}_x$  solution showing the Tyndall scattering effect.



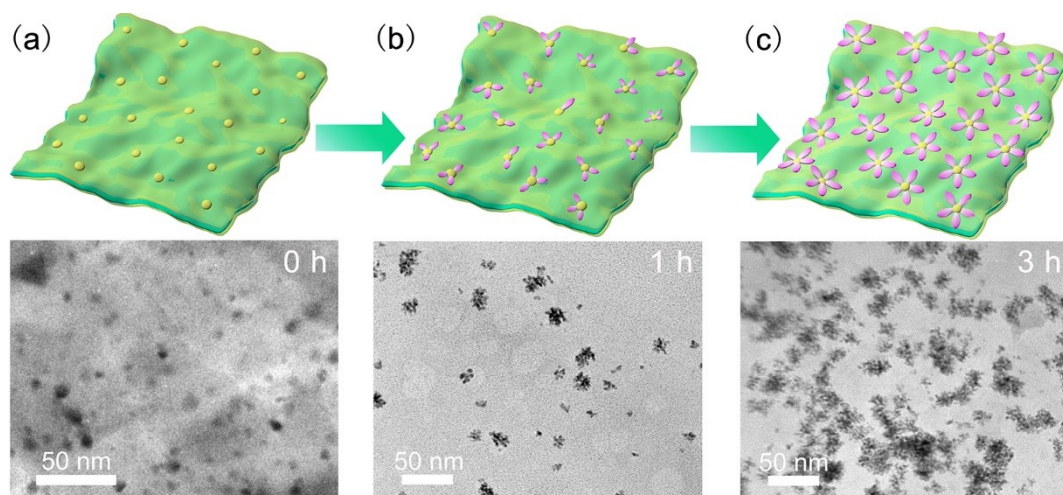
**Fig. S6** The Tyndall effect confirming the colloidal nature of the PVP-functionalized  $\text{Ti}_3\text{C}_2\text{T}_x$  solution.



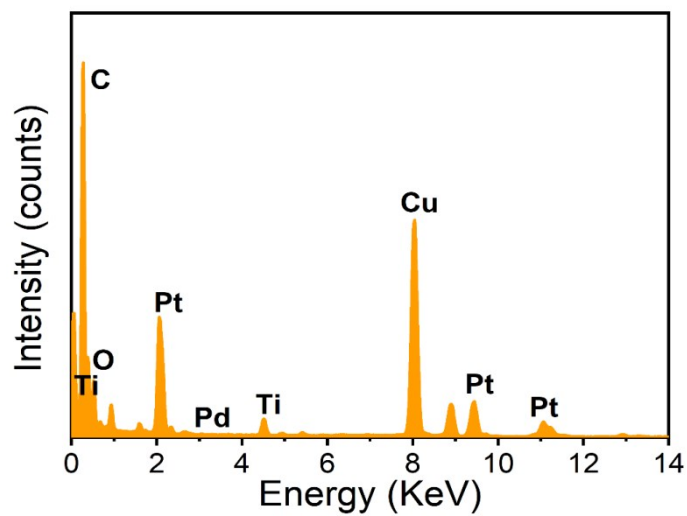
**Fig. S7** TEM images of the Pt-on-Pd/ $\text{Ti}_3\text{C}_2\text{T}_x$  hybrid with different magnifications, implying a uniform distribution of Pt-on-Pd nanoflowers on the  $\text{Ti}_3\text{C}_2\text{T}_x$  nanosheets.



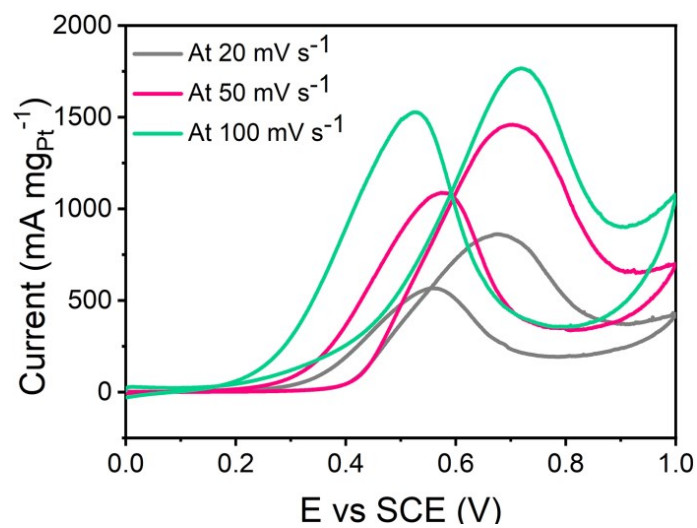
**Fig. S8** (a) TEM image of the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> hybrid and the corresponding elemental mapping images of (b) C, (c) Ti, (d) O, (e) Pd and (f) Pt elements.



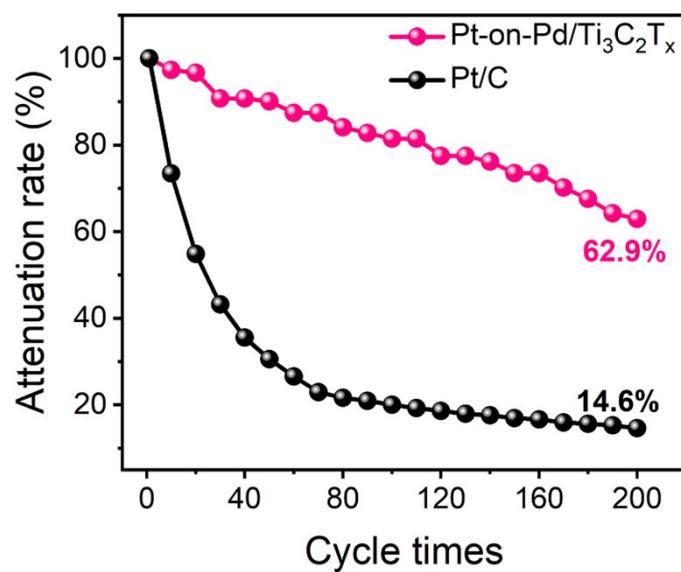
**Fig. S9** TEM images of intermediate products extracted from the reaction mixture of Pt-on-Pd bimetallic nanodendrite/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> hybrids at different reaction times : (a) 0 h; (b) 1 h; (c) 3 h.



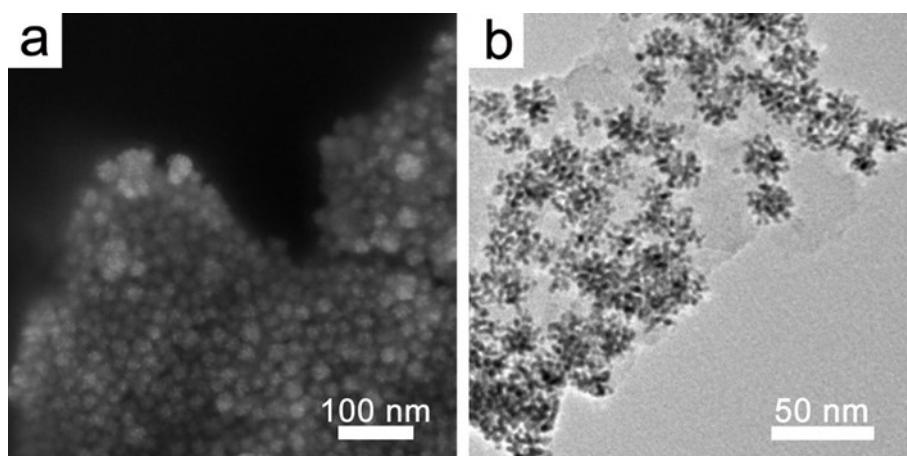
**Fig. S10** EDX spectrum of the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> samples confirms the co-existence of C, O, Ti, Pd and Pt components in the material. The Cu peaks were also observable because the sample was held on a Cu grid.



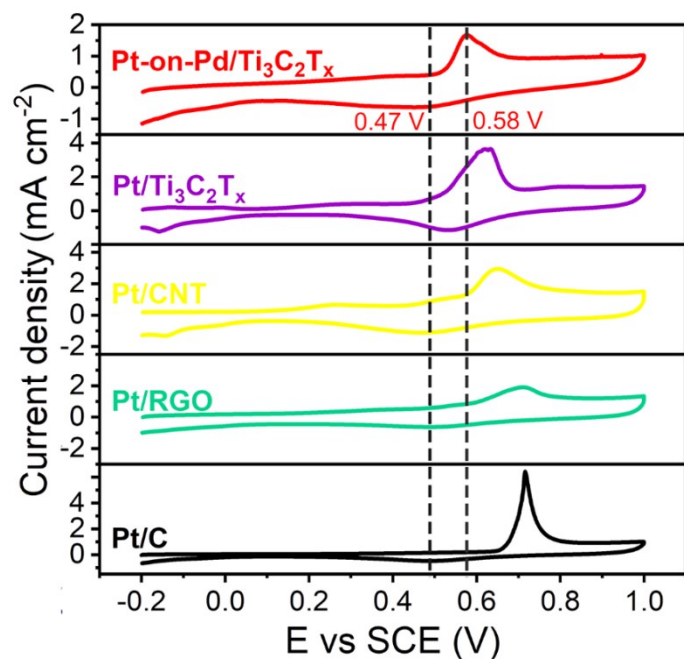
**Fig. S11** The CV curves of the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> catalyst with different scan rates.



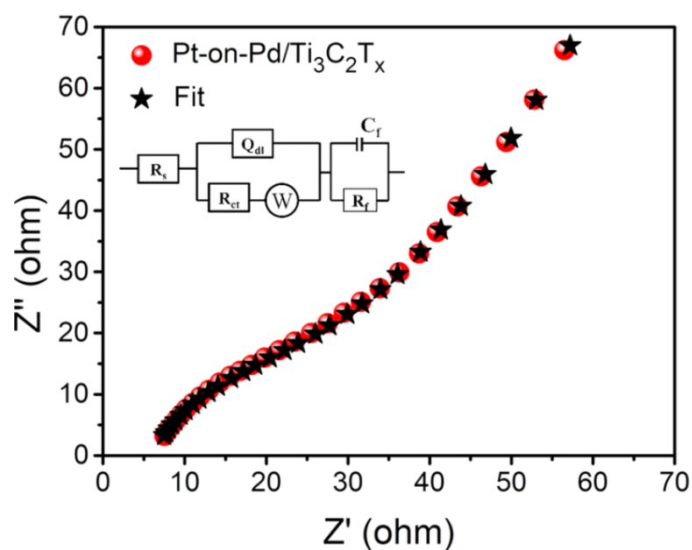
**Fig. S12** The stability behaviors of Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> and Pt/C catalysts toward methanol oxidation, revealing the better cycling stability of Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> catalysts.



**Fig. S13** Typical (A) SEM and (B) TEM images of the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> nanoarchitecture after the long-term chronoamperometric test.



**Fig. S14** CO stripping voltammograms for the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> samples Pt/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub>, Pt/CNT, Pt/RGO, and Pt/C catalysts tested in 0.5 M H<sub>2</sub>SO<sub>4</sub> solution.



**Fig. S15** AC impedance spectrum of the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> electrode and the corresponding fitting curve. The inset is the equivalent circuit: R<sub>s</sub> and R<sub>ct</sub> represent the resistances for electrolyte and catalyst, respectively, Q<sub>dl</sub> is a constant phase element, W represents semiinfinite diffusion at the electrolyte/electrode interface, R<sub>f</sub> and C<sub>f</sub> are the resistance and capacitance for the Nafion-carbon film, respectively.

**Table S1.** Compiled study comparing ECSA and CV results for different catalysts.

Electrode	ECSA (m <sup>2</sup> g <sup>-1</sup> )	Mass activity (mA mg <sup>-1</sup> )	Specific activity (mA cm <sup>-2</sup> )
Pt-on-Pd/Ti <sub>3</sub> C <sub>2</sub> T <sub>x</sub>	157.3	1461.7	0.93
Pt-on-Pd/C	67.9	325.7	0.48
Pt/Ti <sub>3</sub> C <sub>2</sub> T <sub>x</sub>	31.9	215.0	0.67
Pt/RGO	27.6	313.3	0.72
Pt/CNT	27.1	173.0	0.64
Pt/C	23.1	106.0	0.46

**Table S2.** Comparison of methanol oxidation behavior on the Pt-on-Pd/Ti<sub>3</sub>C<sub>2</sub>T<sub>x</sub> hybrids and various Pt-based electrocatalysts.

Catalyst	ECSA (m <sup>2</sup> g <sup>-1</sup> )	Mass activity (mA mg <sup>-1</sup> )	Scan rate (mV s <sup>-1</sup> )	Electrolyte	Ref.
Pt-on-Pd/Ti <sub>3</sub> C <sub>2</sub> T <sub>x</sub>	157.3	1461.7	50	0.5 M H <sub>2</sub> SO <sub>4</sub> + 1 M CH <sub>3</sub> OH	This work
AuPtCu nanowires	N.A.	~500.0	50	0.1 M HClO <sub>4</sub> + 1 M CH <sub>3</sub> OH	S1
FePtPd nanowires	N.A.	488.7	50	0.1 M HClO <sub>4</sub> + 0.2 M CH <sub>3</sub> OH	S2
PtPd/graphene	81.6	647.2	50	0.5 M H <sub>2</sub> SO <sub>4</sub> + 1 M CH <sub>3</sub> OH	S3
Pt/N-doped G	N.A.	~400.0	200	0.5 M H <sub>2</sub> SO <sub>4</sub> +1MCH <sub>3</sub> OH	S4
Pt/N-doped G nanoribbon	64.6	~390.0	20	1 M H <sub>2</sub> SO <sub>4</sub> + 2 M CH <sub>3</sub> OH	S5
Pt/imidazolium-salt ionic liquid/CNT	67.6	~410.0	50	0.5 M H <sub>2</sub> SO <sub>4</sub> + 0.5 M H <sub>3</sub> OH	S6
Pt/[BMIM]BF <sub>4</sub> /CNT	N.A.	155.0	50	0.5 M H <sub>2</sub> SO <sub>4</sub> + 1 M CH <sub>3</sub> OH	S7
Pt/mesoporous carbon	N.A.	~450	20	0.5 M H <sub>2</sub> SO <sub>4</sub> + 1 M CH <sub>3</sub> OH	S8
Pt/macroporous carbon	N.A.	81.6	50	0.5 M H <sub>2</sub> SO <sub>4</sub> + 0.5 M CH <sub>3</sub> OH	S9
Pt/3D MoS <sub>2</sub> -G	62.3	~91.8	10	1 M H <sub>2</sub> SO <sub>4</sub> + 2 M CH <sub>3</sub> OH	S10



## Reference

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