## **Supporting Information to**

## Fabrication of Luminescent Monolayered Tungsten Dichalcogenides Quantum Dots with Giant Spin-Valley Coupling

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**Figure S1.** (a) XRD patterns of WS<sub>2</sub> falkes raw material, residual WS<sub>2</sub> sediment after preparation and dried suspension which contains WS<sub>2</sub> QDs. (b) Local enlarged XRD patterns of residual WS<sub>2</sub> sediment and dried suspension which contains WS<sub>2</sub> QDs. For better presentation, the zero intensities of the patterns were shifted. The diffraction peaks were indexed base on JCPDS: 84-1389. Disappearence of the (002) diffraction peak in dried suspension which contains WS<sub>2</sub> QDs suggest thin structure of the QDs. Due to destroy of layered structure (exfoliation and disintegration), the (002) diffraction of residual WS<sub>2</sub> sediment are much weakened from that of WS<sub>2</sub> raw materials. In the XRD pattern of dried suspension, the other diffraction peaks should from the diffractions of K salt.



**Figure S2.** AFM image of  $WS_2$  QDs. The line was plotted to across the dots' centers (three marked) to minimize measurement errors. This thickness was also proved by three-dimensional AFM image (Figure S3).



**Figure S3.** Three-dimensional view of the AFM image in Figure S2. The majority dots under AFM observation have same thickness (monolayered).



**Figure S4.** Plots of absorbance ( $\lambda_{ex}$  = 393 nm) normalized to cell length, versus the concentration of dialyzed WS<sub>2</sub> QDs product in water solution. The errors arise from uncertainty in the weighing and diluting processes.

Calculation of the Quantum Yield of  $WS_2$  QDs: anthracene in ethonal (QY=0.3) was

chosen as standard. The quantum yield of BN QDs (in water) was calculated according to:

 $\Phi_x = \Phi_{st}(I_x / I_{st})(\eta_x^2 / \eta_{st}^2)(A_{st} / A_x)$ 

Where  $\Phi$ , *I*,  $\eta$  and *A* is the quantum yield, measured integrated emission intensity, refractive index of the solvent and the optical density of the standard (*st*) anthracene and WS<sub>2</sub> QDs (*x*).

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Sample	Integrated	Absorptance at	Refractive	Quantum
	Emission	360 nm ( <i>A</i> )	Index of	Yield $(\Phi)$
	Intensity (I)		Solvent $(\eta)$	
Anthracene	286101	0.0875	1.36	30%
				$(Known)^{1-3}$
WS <sub>2</sub> QDs	75033	0.1642	1.33	~4.0%
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Table S1. Quantum yield of WS<sub>2</sub> QDs suspension (in H<sub>2</sub>O) using anthracene as a reference.



**Figure S5.** (a) The instrument response function. (b) The single exponential function of the instrument response (before afterpulse) showing a 0.55 ns decay time, which is different from the decay time in Figure 4b and confirmed that all the decay time in Figure 4b is not the instrument response.

## Reference

1. Demas, J. N.; Grosby, G. A. The Measurement of Photoluminescence Quantum Yield. A Review. *J. Phys. Chem.* **1971**, *75*, 991-1024.

2. Dawson, W. R.; Windsor, M. W. Fluorescence Yield of Aromatic Compounds. J. Phys. Chem. 1968, 72, 3251-3260.

3. Weber, G.; Teale, F. W. J. Determination of The Absolute Quantum Yield of Fluorescent Solutions. *Trans. Faraday. Soc.* **1957**, *53*, 646-655.