

# Photoluminescence Properties of MoS<sub>2</sub> on h-BN/Au Heterostructure

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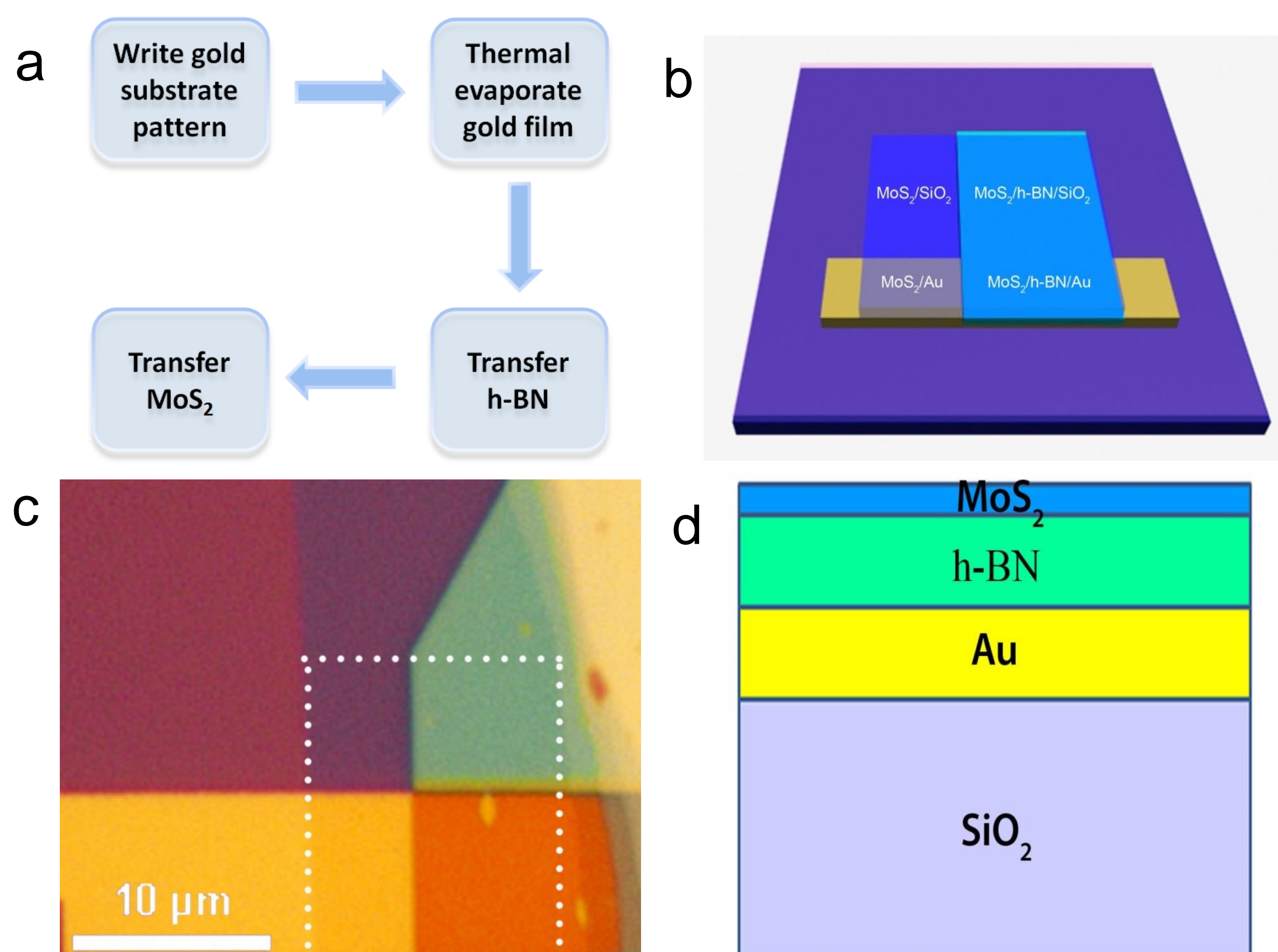
## Abstract

Two-dimensional transition metal dichalcogenides (TMDs) are an emerging group of materials suitable for optoelectronic applications because of their large bandgap related to graphene. When the layer number decreases from bulk to monolayer, their photoluminescence (PL) is greatly enhanced due to an indirect to direct bandgap transition. There are many methods for enhancing the PL of TMDs, such as surface plasma of noble metal nanoarray, reducing the number of trions and defect sites by chemical doping, and choosing suitable substrates. The PL of MoS<sub>2</sub> is low compared to other TMDs due to n doping. Here we report a significant PL enhancement of MoS<sub>2</sub> on h-BN/Au heterostructure. The PL intensity saturation as the laser power is also observed. Such large enhancement is attributed to reduced n doping and field enhancement related mechanisms in this heterostructure.

## Introduction

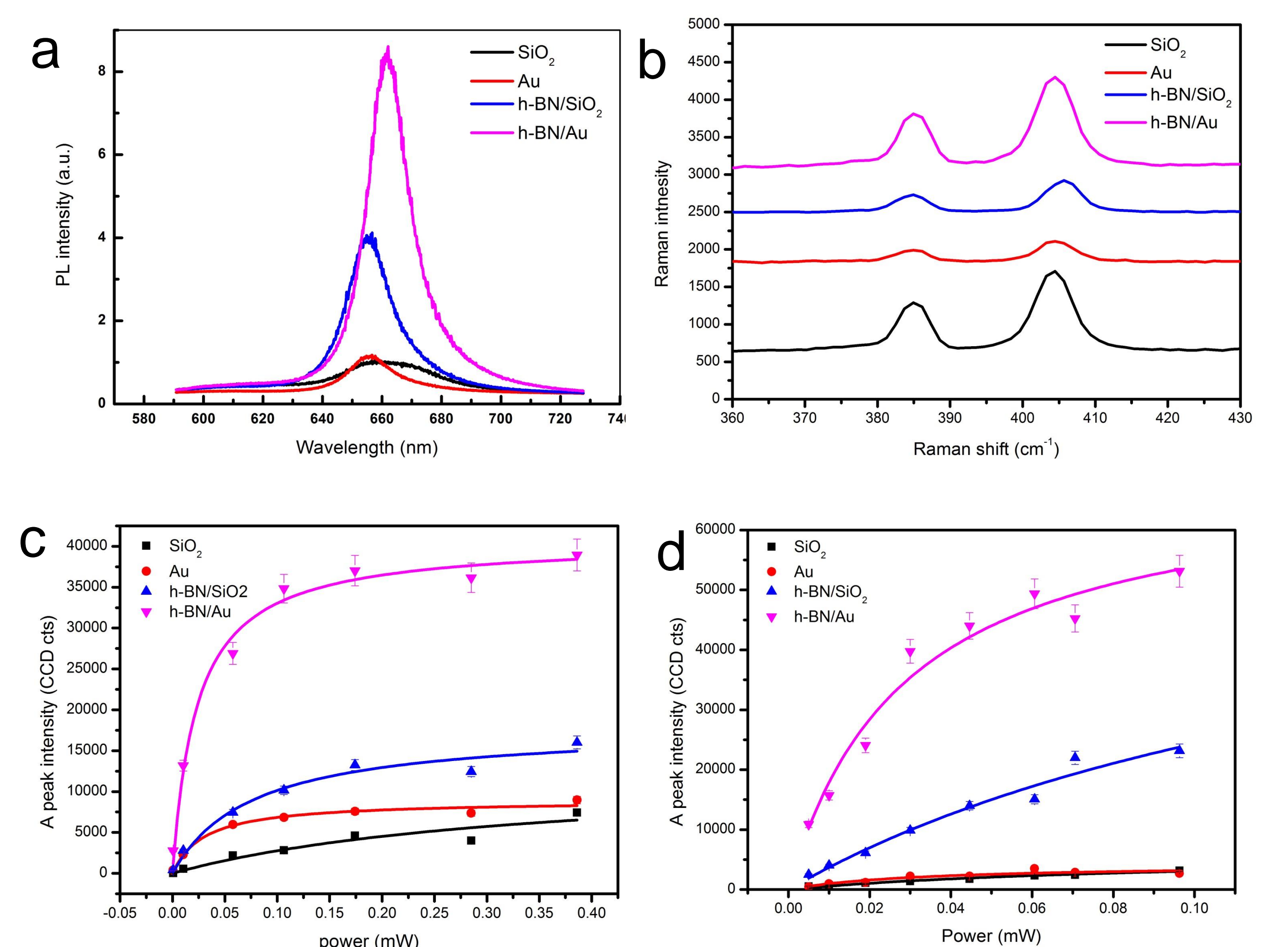
Monolayer two-dimensional (2D) transition metal dichalcogenides (TMDs) are an emerging group of materials suitable for electronic and optoelectronic applications because of their large direct bandgap compared to graphene<sup>[1]</sup>. The large choice of TMD bandgaps and the predicted type II alignment nature<sup>[2]</sup> offer myriad possibilities of fabricating vertical and lateral hybrid structures for photovoltaic<sup>[3]</sup>, photodetector<sup>[4]</sup> and light emission<sup>[5]</sup> applications. The optical properties of monolayer TMDs are dominated by excitons with much stronger binding energies than conventional semiconductors<sup>[6]</sup>. The PL of MoS<sub>2</sub> is low compared to other TMDs. So the PL enhancement of MoS<sub>2</sub> is studied here.

## Methods



(a) The schematic diagram of the device fabrication process. (b) Schematic diagram and (c) the real optical picture of the hybrid structure. The white dashed region is the area where PL and Raman data are collected. (d) The cross section diagram of the MoS<sub>2</sub>/h-BN/Au/SiO<sub>2</sub> structure.

## Results



(a) PL spectra of MoS<sub>2</sub> on SiO<sub>2</sub> (black), Au (red), h-BN/SiO<sub>2</sub> (blue), h-BN/Au (magenta) substrates. The PL spectra on four different substrates were normalized to the PL peak intensity of MoS<sub>2</sub> on SiO<sub>2</sub>. The PL spectra were collected with 532 nm laser, 0.01mw, spot diameter 0.5 μm. (b) Raman spectra of MoS<sub>2</sub> on SiO<sub>2</sub>, Au, h-BN/SiO<sub>2</sub> and h-BN/Au structures. The Raman spectra were collected with 532 nm laser, 0.657mw, spot diameter 0.5 μm. (c) MoS<sub>2</sub> PL power dependence of 532 nm laser. (d) MoS<sub>2</sub> PL power dependence of 633 nm laser.

## Conclusions

The PL intensity of MoS<sub>2</sub> on h-BN/Au heterostructure is greatly enhanced compared to that on SiO<sub>2</sub>. Such large enhancement is attributed to reduced n doping and field enhancement related mechanisms on this heterostructure.

## References

1. Wang, Q. H.; Kalantar-Zadeh, K.; Kis, A.; Coleman, J. N.; Strano, M. S. Nature nanotechnology 2012, 7, (11), 699-712.
2. Kang, J.; Tongay, S.; Zhou, J.; Li, J.; Wu, J. Applied Physics Letters 2013, 102, (1).
3. Furchi, M. M.; Pospischil, A.; Libisch, F.; Burgdörfer, J.; Mueller, T. Nano letters 2014, 14, (8), 4785-4791.
4. Koppens, F. H.; Mueller, T.; Avouris, P.; Ferrari, A. C.; Vitiello, M. S.; Polini, M. Nature nanotechnology 2014, 9, (10), 780-93.
5. Withers, F.; Del Pozo-Zamudio, O.; Mishchenko, A.; Rooney, A.; Gholinia, A.; Watanabe, K.; Taniguchi, T.; Haigh, S.; Geim, A.; Tartakovskii, A. Nature materials 2015, 14, (3), 301-306.
6. Tongay, S.; Suh, J.; Ataca, C.; Fan, W.; Luce, A.; Kang, J. S.; Liu, J.; Ko, C.; Raghunathan, R.; Zhou, J.; Oglethorpe, F.; Li, J.; Grossman, J. C.; Wu, J. Scientific reports 2013, 3.