

# Screening excitonic insulator by liquid metal in proximity

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Excitons in semiconductors are extensively utilized in optoelectronics and photovoltaics and the control over their binding energy is important for various applications. On the other hand, coherent quantum states of excitons can be formed in artificial electron-hole bilayers and also as a ground state in rare and exotic materials called an excitonic insulator. Here we demonstrate the systematic control over the excitonic order parameter in a unique excitonic insulator  $\text{Ta}_2\text{NiSe}_5$ . We investigate using angle-resolved photoemission spectroscopy (ARPES) the band structure of  $\text{Ta}_2\text{NiSe}_5$  during the surface doping by alkali adsorbates. Our polarization-dependent ARPES measurements identify that alkali adsorbates form a liquid metallic layer with its 4s band partially-filled without substantial doping into  $\text{Ta}_2\text{NiSe}_5$ . However, the excitonic band gap decreases systematically upon the increase of the alkali coverage (and the 4s band occupation). This result indicates that excitons are well screened by the metallic surface layer in proximity. We call this effect proximity screening, which may be utilized widely for the control over the exciton binding energy in various different materials.

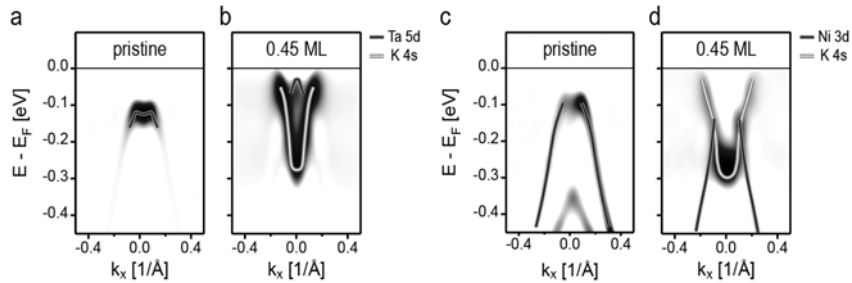


Fig 1. The second derivatives of ARPES data. a-b, vertical polarization data. Ni 3d is invisible. c-d, horizontal polarization data. Ta 5d is invisible. The order parameter is decreased by the liquid K induced proximity screening.