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Superfluid p-H₂ Monolayer in Carbon Nanostructures MILEN KOSTOV, Florida State University — A fluid of para-hydrogen $(p-H_2)$ molecules is a prime candidate for potential superfluid, due to the light mass (half the mass of helium) and the existence of a compound boson ground state. In bulk $p-H_2$ superfluidity is not observed because, unlike helium, molecular hydrogen solidifies at a temperature (triple point T=13.8 K) significantly higher than that (T~2K) at which such phenomena as Bose Condensation and, possibly, superfluidity (SF) might occur. This is due to the fact that H_2 - H_2 interaction is significantly stronger than the He-He one (more than a factor of three in the well depth). One way to attain a liquid ground state at low T is to reduce the effective attraction between the H_2 molecules. Here a novel solution to the problem is proposed, which implies that a SF monolayer p-H₂ can be achieved in a carbon slit-pore with height $H \sim 5.8$ Å, where the alignment of the graphitic planes corresponds exactly to the AB stacking sequence in a pristine hexagonal graphite crystal. Our approach is based on the idea to attain a liquid ground state of p-H₂ monolayer at low T (T \sim 2K), through a substantial renormalization of the pair interaction of $p-H_2$ molecules due to their interaction with the surface electrons of the carbon slit pore. In this environment, the resulting de Boer quantum parameter η for the adsorbed p-H₂ film lies in the vicinity of the threshold value for zero-temperature Bose liquid.

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