

FROM MONOLAYER TO MULTILAYER SILICENE

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Silicene has attracted a huge interest since publication of the archetype 3×3 reconstructed monolayer phase on a silver (111) substrate in 2012 [1]. It is a synthetic emergent two-dimensional Si-based material that might rival graphene for logic applications in electronics [2] : indeed, the first monolayer silicene device, a Field Effect Transistor with ambipolar characteristics, has been fabricated in 2015 [3].

Ordered hydrogenation of 3×3 silicene has been achieved recently in Beijing [4], next in our Lab in Marseille, with further High Resolution Electron Energy Loss spectroscopy results [5].

Multilayer silicene has been also synthesized ; it possesses an intrinsic $\sqrt{3} \times \sqrt{3}$ surface reconstruction, hosts Dirac fermions, and is self-protected in ambient air by its native oxide [6,7]. However, despite such evidences, its mere existence has been severely questioned.

Here, we will show thanks to novel *in situ* Photoelectron Spectroscopy data as well as *ex situ* Surface X-Ray Diffraction and Raman spectroscopy results that multilayer silicene is effectively grown in a rather low temperature regime ($\sim 200^\circ\text{C}$), while, instead, diamond-type silicon terminated by the well-known $\text{Si}(111)\sqrt{3} \times \sqrt{3}\text{-Ag}$ superstructure is formed at higher temperatures.

Keywords: Silicene; 2D Phases; Structural and Electronic Characterizations

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