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and SrTiO₃(110) surfaces in r.t. and Ar atmosphere were studied by the in situ electron diffraction and Raman spectroscopy. The (101) surface was adsorbed by 4-methylpyridine. The formation of CO was observed by in situ electron spectroscopy for reaction of 4-methylpyridine with the (101) surface. The cross-sectional views by electron microscopy were obtained by consecutive steps of reaction with and without argon. The (101) surface covered by 4-methylpyridine was subsequently irradiated, and the cross-sectional views were again obtained by

electron microscopy. The adsorbed 4-methylpyridine covered by a thin film of CO and the 4-methylpyridine derived from the 4-methylpyridine were generated on the (101) surface. For the reaction of 4-methylpyridine with SrTiO₃ (110), (110) surface was covered with the intermediate. The intermediate reacted with H₂O to form OH on the surface and then reacted with 4-methylpyridine. The resultant 4-methylpyridine was adsorbed on the substrate. The hydrogen bonding of 4-methylpyridine to OH and to surface was confirmed by

adsorption on

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