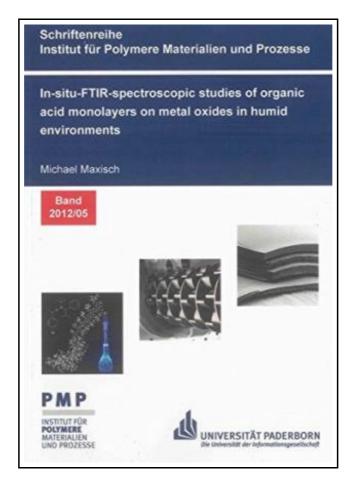
In-situ-FTIR-spectroscopic studies of organic acid monolayers on metal oxides in humid environments



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Reviews

It in one of my favorite book. Sure, it is actually engage in, nonetheless an interesting and amazing literature. I am happy to let you know that this is basically the finest book i have got study inside my very own existence and might be he finest publication for ever. (Randal Reinger)

IN-SITU-FTIR-SPECTROSCOPIC STUDIES OF ORGANIC ACID MONOLAYERS ON METAL OXIDES IN HUMID ENVIRONMENTS



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Shaker Verlag Jun 2012, 2012. Buch. Book Condition: Neu. Neuware - A new in-situ setup combining ultra-high vacuum (UHV), plasma application, modern photoelastic-modulated-infrared-reflection-absorptionspectroscopy (PM-IRRAS) and quartz crystal microbalance (QCM) was developed to analyze the formation and stability of selfassembled monolayers (SAMs) on engineering metals in detail. At first the adsorption behaviour of different organophosphonic acids on nickel-titanium (NiTi) shape memory alloy (SMA) were analysed. PM-IRRAS prove that the binding mechanism is based on a mono- or bidental bond. In addition it was shown that an alkyl chain length of 17 CH2 groups is required for the formation of a SAM, which is stable in aqueous environment. Furthermore in-situ measurements were undertaken to determine the formation, the barrier properties and the stability under high water activities of octadecylphosphonic acid (ODPA) and nonadecanoic acid (NDA) SAMs. PM-IRRAS reveal the self-assembly of both molecules on oxyhydroxide covered aluminium surfaces and prove their stability during exposure to humid air. For ODPA and NDA SAMs a strongly reduced amount of water - based on the inability of water to form hydrogen bonds to the low energy aliphatic surfaces was observed. However, the ODPA monolayer covered surfaces did not show a significant inhibition of the H2O/D2O isotope exchange reaction between a D2O gas phase and the hydroxyl groups of the aluminium oxyhydroxide film. The high stability of the interfacial phosphonate group as well as the constant ordering of the SAM are referred to the strong acid-base interaction of the phosphonate head group with the Al ions as well as the molecular interactions of the aliphatic chains. For NDA monolayer covered Al substrates the kinetic of surface hydroxylation by dissociative adsorption of H2O molecules could be significantly reduced while the interfacial carboxylate group as well as the orientation of the NDA monolayer is not affected by...

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