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Photochemical Reactivity of Ba1-xSrxTiO3 Solid Solutions AN-DREW GAMALSKI, Arizona State University, NINA V. BURBURE, GREGORY S. ROHRER, Carnegie Mellon University — The photochemical reactivities of SrxBa1-xTiO3 solids solutions were investigated as a function of composition. Samples with x = 0 to 1 were used to reduce Ag+ in aqueous solution to Ag0 on the sample surface and the relative heights and densities of the silver deposits, as measured by atomic force microscopy, were used to quantify the relative reactivities. A local maximum in the reactivity is observed at x = 0.26, which is near the cubictetragonal phase boundary. Based on these observations, it is concluded that the increase in the dielectric constant at this composition increases the width of the depletion layer and this promotes electron transfer to the surface and increases the reactivity.

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