

# Control of active species in the afterglows of N<sub>2</sub> and N<sub>2</sub>-O<sub>2</sub> microwave plasmas and the role of those active species in the selective surface nitridation of TiO<sub>2</sub> nanocrystals

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TiO<sub>2</sub> is widely used as a catalyst in various applications including photocatalysis, photovoltaics and sensors. However, the performance of TiO<sub>2</sub> is strongly determined by the detailed chemical structure of surfaces. So, the selective control of surface structure can be a good choice for the control of activities associated with it. As a one way to achieve such a goal, plasma treatments are widely used as in the case of enhancing the photoresponse of TiO<sub>2</sub> materials. Plasma treatments of TiO<sub>2</sub> have been used to enhance the visible light absorption for enhanced photocatalytic activity [1]. To understand the origin of enhancement, it is necessary to study the chemical states and bonding structure of the surface-modified TiO<sub>2</sub> materials.

In this study, anatase TiO<sub>2</sub> nanocrystals was exposed to the post-discharge region of N<sub>2</sub> microwave plasma and the chemical bonding states of surface nitrogen species on the surface-modified TiO<sub>2</sub> were carefully evaluated using X-ray photoemission spectroscopy (XPS) [2]. We observed that the surface treatments in the afterglows can induce stable nitrogen species which are formed at or near the surface of TiO<sub>2</sub>. However, the detailed bonding configuration of N species can vary strongly depending on the type of active species present in the afterglow. Fig. 1 shows an example of N 1s core level spectra taken from the TiO<sub>2</sub> sample treated in the afterglows with N<sub>2</sub>-O<sub>2</sub> microwave plasma. Here, it shows two distinct N species induced by the interaction of TiO<sub>2</sub> surface with the active species in the afterglow region. The high concentrations of O or NO in the afterglows of N<sub>2</sub>-O<sub>2</sub> plasmas can induce the formation of nitrate species in addition to the interstitial N species. In addition, a prolonged exposure in the early afterglow was found to induce additional N species at lower binding energies which is attributed to substitutional N species.

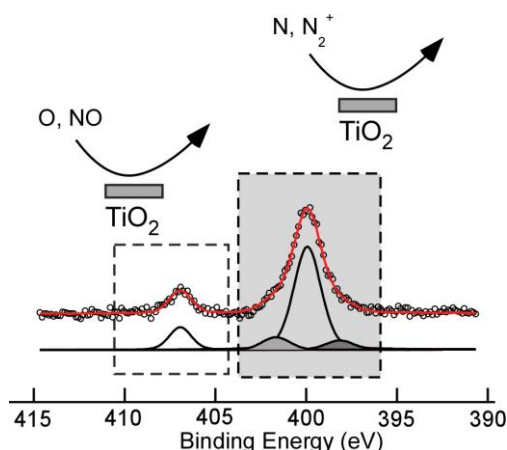


Fig. 1. N 1s core level spectrum of TiO<sub>2</sub> nanocrystals treated in the afterglow of N<sub>2</sub>-O<sub>2</sub> microwave plasma showing different N species induced by different active species in the afterglow.

## References:

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