Photosensitive composites for light activated gas sensors

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This work is devoted to the functionalization of semiconductor oxides to create gas sensors operating under photoactivation instead of thermal heating in order to reduce energy consumption. This goal is achieved by modifying the sensitive layer of semiconductor oxides with photosensitizers. The objects of the study can be divided into 2 groups depending on the type of photosensitizer used. In both cases, wide-band metal oxide semiconductors SnO₂ and In₂O₃ were chosen as the matrix. These semiconductor oxides, on the one h and, are characterized by different concentration of bulk defects that determine the concentration and mobility of free charge carriers, and on the other hand, they differ in their chemical properties that affect the type and concentration of surface active centers.

In the first group, the photosensitizers are Ru (II) heterocyclic complexes. Their photoexcitation with blue LED ($\lambda_{max} = 470$ nm) leads to electron-hole pairs generation. The electrons are transferred from the LUMO level to the conduction band of the oxide matrix, which is lower in energy. At the same time, photoexcited holes remaining in the organic complex can also be transferred to an oxide matrix, where they recombine with electrons localized in chemisorbed molecules. Our obtained and published results showed the promise of using organic-inorganic hybrid materials based on semiconductor oxides (SnO₂, In₂O₃) and Ru(II) heterocyclic complexes for NO₂ and NO detection [1, 2].

In the second group, nanocrystalline titanium dioxide was chosen as a photosensitizer, because of its high photocatalytic activity in oxidation of volatile organic compounds (VOCs) at room temperature under UV light. This part of the work is devoted to the detection of VOCs using TiO₂@SnO₂ nanocomposites under UV photoactivation ($\lambda_{max} = 365$ nm) [3]. The obtained results showed that SnO₂/TiO₂ nanocomposites can be perspective in formaldehyde detection in sub-ppmrange at T = 150 °C. Further modification of SnO₂/TiO₂ nanocomposites with Au NPs increases the sensor signal at T = 100°C, while modification with Pt NPs gives rise to the appearance of sensor response at T = 25°C and 50°C. The reactions of VOCs oxidation are activated by Au and Pt nanoparticles that lead to increased selective sensitivity toward HCHO, especially in 0.1 – 1 ppm concentration range [4].

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