

Review

Effect of Nanoparticle Interaction on Structural, Conducting and Sensing Properties of Mixed Metal Oxides

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Abstract: This review analyzes the studies published, mainly in the last 10–15 years, on the synthesis, structure, and sensor properties of semiconductor nanocomposites. Particular attention is paid to the interaction between nanoparticles of the sensitive layer, and its effect on the structure, sensitivity, and selectivity of semiconductor sensor systems. Various mechanisms of interaction between nanoparticles in metal oxide composites are considered, including the incorporation of metal ions of one component into the structure of another, heterocontacts between different nanoparticles, and core-shell systems, as well as their influence on the characteristics of gas sensors. The experimental data and studies on the modeling of charge distribution in semiconductor nanoparticles, which determine the conductivity and sensor effect in one- and two-component systems, are also discussed. It is shown that the model which considers the interactions of nanoparticles best describes the experimental results. Some mechanisms of detection selectivity are considered in the conclusion.

Keywords: mixed metal oxide sensor; nanoparticle interaction; morphology; nanoparticle structure; sensitivity; selectivity



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1. Introduction

Sensors based on semiconductor metal oxides are currently widely used to detect various gases in the environment. The principles for production of highly efficient sensors, based on metal oxides, for the detection of various substances have been investigated for more than four decades (see, for example, reviews [1–5]). In this review, we consider studies elucidating the influence of the composition, structure, and chemical properties of semiconductor composites, on the sensor characteristics, undertaken mainly in the last 10–15 years. Particular attention is paid to the interaction between the components of nanostructured composites and its effect on the chemical processes that determine the operational properties of metal oxide sensors. All the studies cited below have used composites prepared by different methods, including the mixing of ready-made nanopowders, various impregnation methods, hydro- and solvothermal treatment, the sol-gel technique, the electrospinning method, and others. The techniques used for the preparation of different composites in the literature and associated references are summarized in two tables in a subsequent section.

The operation of metal oxide sensors is based on the change in their conductivity, under the influence of the analyzed gases. Depending on the oxygen content in the environment, this process is described by two different models. The ion-sorption model describes the sensor process at a high oxygen concentration. The chemisorbed oxygen molecules present in the sensitive layer of the particle capture electrons from the conduction band of the metal oxide, creating oxygen anion centers, such as O^- , O_2^- , and O^{2-} on the surface. As a result, a negatively charged layer is formed on the surface of the nanoparticles, and the

near-surface region is depleted of conduction electrons (the depletion region), due to their escape to the surface from the near-surface layer [6].

The resistance of the sensor depends on the parameters of the barrier that the electrons overcome during the transition between particles, and the concentration of electrons in the layers near the surface of the particles. The distribution of electrons in a nanoparticle is determined by the equilibrium conditions (see Section 3). Correspondingly to the change in the concentration of electrons in the layers near the surface of the particles, the sensor resistance increases.

The interaction of the molecules of the analyzed gas, adsorbed on the surface of metal oxide nanoparticles with the active oxygen centers, results in the release of trapped electrons, which subsequently return to the conduction band of the semiconductor. There occurs an increase in the conductivity of electronic (*n*) semiconductors and a decrease for hole (*p*) semiconductors.

At low oxygen concentrations, for example, to detect impurities of reducing gases in an inert gas, the sensor process proceeds in accordance with the vacancy model [7,8]. In this case, the reducing gas reacts with the oxygen ions of the O^{2-} lattice of the metal oxide on the surface of the sensitive layer nanoparticles, to form oxygen vacancies containing weakly bound electrons, which also increases the conductivity of the sensor [8]. This review considers the processes occurring in air ambience and therefore, the most common ion-sorption model is used.

The type of oxygen ions adsorbed on the surface of a semiconductor metal oxide depends on temperature (see Figure 1). At temperatures in the range of 200–400 °C, at which metal oxide sensors mainly “work” for the detection of reducing gases, oxygen anions (O^-) are the active centers [8–10].

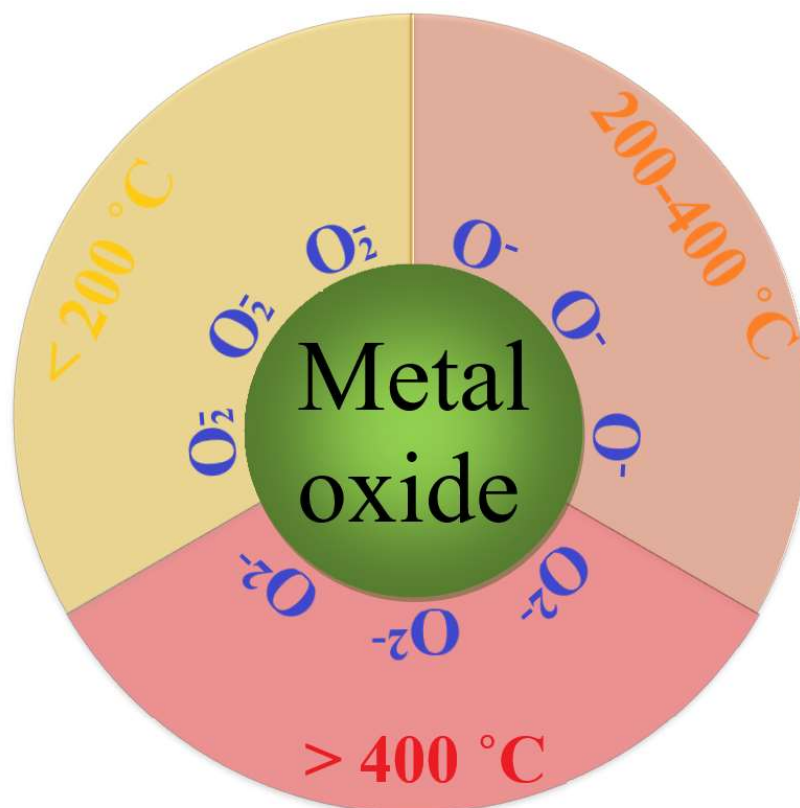


Figure 1. Schematic representation of the priority formation of active oxygen species on the surface of metal oxide nanoparticles at different temperatures.

As for the O^{2-} ions, there are no convincing experimental data on the state and behavior of these particles on the metal oxide surface, and the calculations in some previous

studies [11,12] give contradictory results. In addition, these calculations do not consider surface oxygen vacancies, which play an important role in the sensor process. All of the above support the sensor response model involving the radical ion O^- .

A generally accepted approach to improving the sensitivity, rate, and selectivity of the sensors is to use composite systems that combine metal oxides with different electronic characteristics and chemical properties (see, for example, [13–20]). Until recently, only systems containing small additives (no more than 5–10%) of modifying oxides were considered in the study of sensor characteristics of metal oxide composites. A detailed study of the behavior of composites containing measurable amounts of different oxides began with the studies in Refs [20–22]. In the last 5–6 years, studies by other authors have appeared, that considered sensor systems consisting of mixtures of oxides over a wide range of compositions (see, for example, [23,24]).

The sensor properties of metal oxide composites are determined by the electronic structure of the nanoparticles, which, in turn, is affected by the interaction between the components of the composite. It is this interaction that determines the sensor activity of metal oxide nanocomposite sensors.

Depending on the relative content of metal oxides, it is reasonable to classify the two-component systems into single-phase and two-phase systems. A single-phase composite is a solid solution in which an ion of the doping oxide enters the lattice of nanocrystals of another oxide. Such solutions are formed only in the range of doping oxide concentrations not exceeding 10%. The difference in the sizes and charges of the ions forming the lattice produces additional deformations and defects in the lattice. This stimulates the adsorption of oxygen and detected gases on the surface of nanocrystals, which increases the sensor effect.

Binary two-phase systems, formed at comparable concentrations of oxides, consist of a solution of a small addition of one oxide (up to 10%) in the lattice of another oxide, without nanocrystal of the first component in the system. The influence of the interaction of components on the sensitivity of the sensor system is due to the contact between nanocrystals of different phases.

An analysis of the available literature shows that the sensor effect of nanostructured metal oxide systems is determined mainly by two factors [25,26]. The first is the chemisorption of the detected gas and oxygen on the surface of sensor particles, and the subsequent interaction of the chemisorbed gas with active oxygen centers. The second factor is the change in the electronic structure of metal oxides during the sensor reaction, which affects the conductivity of the sensitive layer. It is important to note that in a system consisting of metal oxides with different electronic and physicochemical properties, both of the above factors depend on the interaction between these components. This interaction largely determines the behavior of metal oxide nanocomposite sensors and makes it possible to purposefully change their performance characteristics.

Since the metal oxides used in sensors are ionic compounds, their contact in mixed oxide composites can lead to the transition of metal ions of one component into the nanoparticles of another (doping of nanoparticles). This causes the transfer of electrons between particles, their mutual charging, and changes in the structure of the nanoparticles [27,28]. The introduction of small concentrations (up to 5–7%) of metal ions with a higher charge into the metal oxide results in the appearance of an additional positive charge.

Compensation of this charge occurs due to a decrease in the concentration of positively charged oxygen vacancies and an increase in the concentration of conduction electrons. Thus, when In_2O_3 is doped with tin oxide, In^{+3} ions are replaced by Sn^{+4} ions, which are accompanied by an increase in conductivity and a decrease in the sensor response to hydrogen [19]. Replacing lattice ions with lower valence ions creates localized negative charges on metal ions embedded in the lattice. In this case, the concentration of positively charged oxygen vacancies in the lattice increases [29], leading to a decrease in the concentration of conduction electrons in the metal oxide.

An important role in the sensor process is also played by the transfer (spillover) of chemisorbed molecules and products of their reactions with metal oxides between the contacting nanoparticles. Thus, a noticeable sensor effect in a system of SnO₂ nanofibers, coated with a layer of catalytically active Pd nanoparticles, is explained by the spillover of oxygen atoms formed on the surface of Pd nanoparticles, upon dissociation of O₂ [30]. Oxygen atoms transfer to SnO₂ nanofibers, capturing conduction electrons from them, and, as a result, are converted into O[−] radical anions.

The sensor process proceeds according to a similar scheme in a system containing comparable concentrations of ZnO and In₂O₃ [22]. An increase in the sensor effect upon the addition of ZnO occurs because of the spillover of O[−] radical anions formed during the dissociation of oxygen, on catalytically active ZnO, into tin oxide nanoparticles. These O[−] radical anions determine the conductivity of the composite.

The theory of the sensor effect in single-component systems, allowing for the distribution of conductive electrons, was developed in previous studies [14,31–35]. A comparison is made in these studies with experimental literature data, and the approaches of other authors to the description of the sensor process are discussed. The theoretical description of the sensor effect is largely based on the distribution of electrons in semiconductor nanoparticles [31,36,37]. In the case of nanoparticles with high electron content in the conduction band, the important role of the negatively charged layer is considered. Various mechanisms of interaction between nanoparticles are also considered, and the distribution of electrons and the sensor effect in two-component systems is described [37].

2. Mechanisms of Sensitivity of Semiconductor Metal Oxide Sensors in the Detection of Reducing Gases

The sensor effect, which characterizes the sensitivity of the sensor, is mainly determined by three factors: the specific surface area of the sensitive nanostructured layer; the adsorption of oxygen and analyzed compounds on the surface of nanoparticles; and the reactivity of adsorbed compounds with respect to active oxygen centers [2].

The sensitivity of semiconductor metal oxide sensors can be significantly increased by using nanostructured sensitive layers based on two-component materials, consisting of metal oxides with different electronic characteristics and chemical properties (see, for example, [13–16]). The mechanisms for the increase in sensitivity depend on the interaction between the metal oxide components, which is affected by their phase state. With this in mind, we will classify sensor systems into *single-phase* and *two-phase* types.

As noted above, a single-phase, two-component system is a solid solution in which metal ions of one component enter the lattice of nanocrystals of another component. The presence in the lattice, of contacting metal ions of different sizes and charges, contributes to the formation of additional deformations and defects in the crystal (in particular, oxygen vacancies), which stimulate the adsorption of oxygen and analyzed gases on the surface of nanocrystals and increase the sensor effect.

However, metastable solid solutions in a two-component metal oxide system are formed only in the region of low doping oxide concentrations, which depend on the size and valence of metal ions, and do not exceed 10%. At higher concentrations, segregation of solid solutions occurs when the particles of the segregated material coexist with the particles of the solid solution [14,19,38].

In binary two-phase systems, the influence of the interaction of components on the sensitivity of the sensor system is due to contacts between nanocrystals of different phases. The transfer of electrons between nanocrystals with different work functions, during the contact of oxides, leads to mutual charging of the nanocrystals [27], which, in turn, affects the adsorption of molecules, especially dipole types.

Of particular interest are contacts between catalytically active metal oxides and electron-rich nanoparticles that form the conduction paths of a sensor system (see, for example, [14,19,39]). In this case, the increase in sensitivity is due to the transition into

conducting nanocrystals of atomic and molecular particles, arising from the dissociation of oxygen molecules and the adsorption of the analyzed gas.

2.1. Mechanisms of Increasing the Sensitivity of Two-Component, Single-Phase Metal Oxides

Various processes occur in the sensitive layer of two-component, single-phase metal oxides that affect their properties. In the following sections, we consider the effects associated with the doping of nanoparticles of one component with the molecules or atoms of another. These effects include the change in the particle lattice during doping, the role of doping in the adsorption of oxygen and detected gases, and the influence of the nature of doping ions.

2.1.1. Change in the Lattice of Particles during Doping—Dependence of the Interaction of Doping Ions with Ions of Main Lattice on Their Valence

The formation of binary metal oxide systems, by coprecipitation of components from solutions containing a mixture of salts or organometallic precursors or by impregnation of crystals of one metal oxide with another metal, can be accompanied by the incorporation of one of the components into the crystal structure of the second oxide, i.e., doping [13,16,40]. In this case, the metal ions of the base oxide are replaced by doping ions. There are two types of doping, namely isovalent doping, in which the valences of the main ions that make up the crystal lattice and the doping ions are the same, and heterovalent doping, when the valences of the metal ions are different.

The change in the structure and properties during isovalent doping is due to the deformation of bonds in the main crystal when doping ions are introduced into its lattice, the size of which differs from the size of the main ions. The energy of ionic bonds (E_{MO}) in the metal oxide crystal depends on the surface charge density of the cation in the lattice and increases with decreasing cation size.

The influence of lattice deformations on the concentration of oxygen vacancies depends on the method of formation of a two-component system. A typical example is In_2O_3 nanocrystals doped with aluminum oxide. During the formation of $Al_2O_3-In_2O_3$ nanocrystals from a mixed solution of indium and aluminum salts by electrospinning, the In^{+3} ions (radius 0.80 Å) are partially replaced by the smaller Al^{+3} ions (radius 0.54 Å). This leads to the contraction of the lattice of In_2O_3 nanocrystals and their rearrangement into a stronger rhombohedral phase, which is accompanied by an increase in E_{MO} and a decrease in the concentration of oxygen vacancies [41].

A similar result was obtained for SnO_2 nanocrystals synthesized by electrospinning (Sn^{+4} radius 0.71 Å) and doped with Ti^{+4} ions (radius 0.64 Å) [42]. At the same time, $Al_2O_3-In_2O_3$ nanocrystals formed by the hydrothermal method retain the weaker cubic structure of In_2O_3 crystals. Lattice compression upon the doping of crystals with aluminum ions leads to the formation of defects and an increase in the concentration of oxygen vacancies [43]. These vacancies form centers of oxygen chemisorption, which increase the sensor effect in the detection of reducing compounds [43,44].

An increase in the concentration of oxygen vacancies and the associated increase in the sensor effect are more pronounced because of lattice expansion upon the isovalent doping of metal oxide nanocrystals with metal ions of larger size than the main lattice ions. This is clearly seen in the example of In_2O_3 nanocrystals with In^{3+} ions (radius 0.80 Å) doped with La^{3+} ions (radius 1.03 Å) [45,46], which is explained by the weakening of bonds in the In_2O_3 lattice under the influence of La^{3+} ions.

The most significant effect on the sensitivity of two-component, single-phase metal oxides is provided by heterovalent doping. Due to the difference in the valence of metal ions, not only local deformations but also new local charges arise in the lattice of a nanocrystal. The change in the sensor properties of the metal oxide during doping depends on the ratio of the charges of the doping ions to the main ions of the crystal lattice, which determines the conductivity.

The introduction of up to 5–7% of metal ions with a larger charge into the basic oxide lattice produces an additional positive charge in the system. The concentration of positively charged oxygen vacancies decreases and the concentration of conduction electrons increases. As a result, when In_2O_3 is doped with SnO_2 and In^{3+} ions are replaced by Sn^{4+} ions, an increase in conductivity and a decrease in the sensor response to hydrogen are observed [19,40].

When lattice ions are replaced by ions with a lower valence, localized negative charges appear on metal ions built into the lattice, the concentration of positively charged oxygen vacancies increases [29], and the concentration of conduction electrons decreases. Such a process is considered in the example of the formation of the SnO_2 - In_2O_3 composite [29,40]. Since the size of In^{+3} is larger than that of Sn^{+4} , the substitution of In^{+3} for Sn^{+4} results in an increase in interplanar distances in the SnO_2 lattice and an increase in the response to hydrogen [40,47].

When lattice ions are replaced by doping ions, in contrast to their incorporation into the intercrystallite space, the emerging vacancies can form complexes with these ions. Such complexes were characterized for In_2O_3 nanocrystals doped with Mn^{+2} and Co^{+2} ions [48]. The vacancies formed when these ions are included in the In_2O_3 lattice, and enter the first coordination sphere of oxygen anions surrounding the doping ion [49].

Many metal ions form specific bonds with organic molecules. Such complexes, in which the centers of oxygen chemisorption (oxygen vacancies) are associated with the centers of adsorption of the detected gas on embedded metal ions, create especially favorable conditions for the sensor process. Thus, the impregnation of nanocrystalline In_2O_3 by divalent cobalt, which is accompanied by the incorporation of Co^{+2} ions into the In_2O_3 lattice, produces a material with an exceptionally high response to CO (Figure 2) [50]. Impregnation of In_2O_3 nanoparticles by other metals also increases the sensor effect, but much less so than in the case of Co^{+2} ion incorporation.

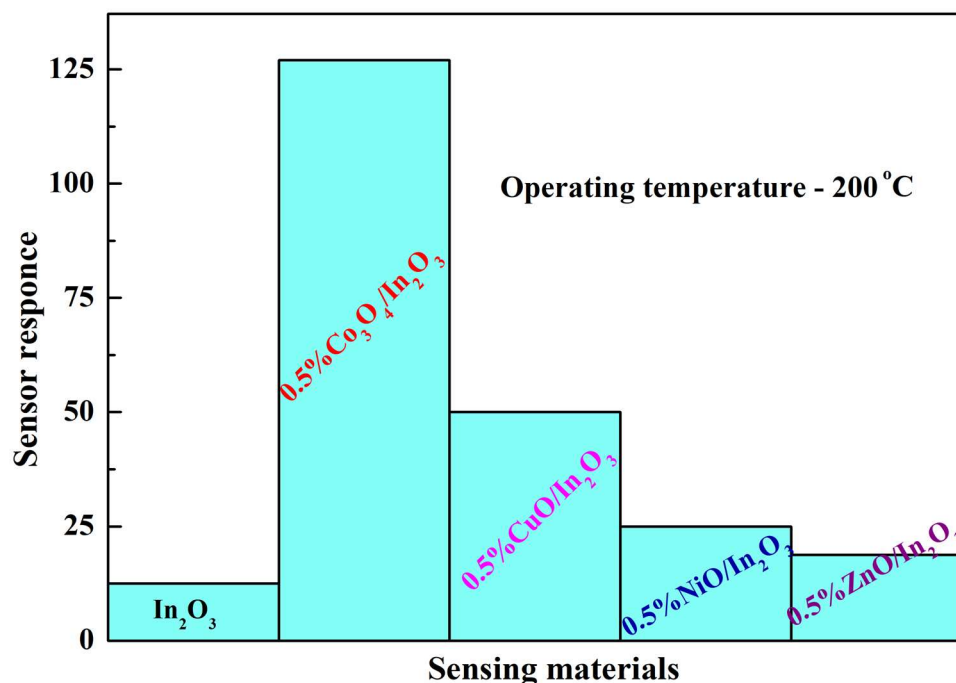


Figure 2. Sensor response of In_2O_3 -based nanocomposites impregnated by various metal oxides in the detection of 1000 ppm CO.

The activity of In_2O_3 doped with Co^{+2} ions is attributed to the fact that complexes of cobalt ions with oxygen vacancies are active centers of the sensor reaction [48]. The bonds of these ions with CO molecules [51] increase the adsorption of CO on such a complex.

2.1.2. Effect of Doping on Adsorption of Detected Gases

A change in the structure of metal oxide nanocrystals, due to their doping with metal oxides, leads to the formation of new adsorption centers. Two types of adsorption bonds between adsorbed organic molecules and metal oxide nanocrystals of the sensitive layer can be distinguished: acid–base bonds [52] and bonds between π -electrons of adsorbed molecules and d-electrons of metal ions in metal oxides [53,54].

Acid–base bonds are characteristic of adsorption on metal oxides with polar functional groups containing nitrogen or oxygen atoms. Doping of metal oxide nanocrystals with basic metal ions, including alkaline earth metal ions, creates adsorption centers for acidic compounds in their lattice. Such compounds are alcohols, aldehydes, ketones, etc., related to Brønsted acids [55,56]. Doping not only increases the number of adsorbed molecules but also can change the strength of their bond with nanocrystals, depending on the acidity of the molecules.

The influence of doping of In_2O_3 nanocrystals with Ca, Sr, and Ba ions on sensor effects in the detection of formaldehyde, ethanol, and acetone by sensors based on these nanocrystals has been studied [57]. The sensor effect in the detection of formaldehyde decreases from Ca to Ba [55]. This is due to a drop in the electronegativity of metal ions and a decrease in their ability to attract electrons from the surrounding oxygen ions of an oxide. Their electron donation and the energy of bonds between the metal ions and adsorbed acid-type molecules also increase [58].

The sensor effect for In_2O_3 nanocrystals doped with alkaline earth metal ions increases sharply [55] with an increase in the acidity of the analyzed compounds in the series acetone ($\text{pK}_a = 20$), ethanol ($\text{pK}_a = 15.9$), and formaldehyde ($\text{pK}_a = 13.27$). Thus, there is a correlation between the acid–base bonds of the detected molecules with the sensitive layer and the magnitude of the sensor effect.

The nature of the adsorption bonds of π -electron molecules with a sensitive metal oxide layer depends on the electronic structure of the cations that make up this layer. Such bonds have been the most studied in relation to the CO molecule [59]. For effective detection, CO bonds with metal ions must be strong enough to ensure reliable CO adsorption on the surface of the sensor layer, at temperatures of 200–300 °C.

These bonds should not interfere with the sensor reaction of adsorbed molecules with oxygen active sites, mainly O^- . In this respect, the doping of oxides with divalent metal ions, in particular, Mg^{2+} , Mn^{2+} , Fe^{2+} , Co^{2+} , Ni^{2+} , and Zn^{2+} , which are characterized by reversible binding to CO molecules, is of particular interest [51]. Simultaneously, upon doping with these ions, oxygen vacancies appear in the system, which are the active centers of the sensor reaction. Ions in which d-electron shells are either absent (Mg^{2+}) or completely filled (Zn^{2+}) have the lowest binding energy with CO molecules. This bond is due to the electrostatic attraction between the metal ion and the lone pair of electrons of the 5σ orbital of the C atom in the CO molecule; the energy of such a bond is about 8 kcal/mol [53,60].

When CO interacts with metal cations with unfilled d-electron shells (Co^{2+} , Ni^{2+}), the electron pair of the 5σ -orbital of carbon is connected with the d-electrons of the metal. Concurrently, due to the overlapping of the unoccupied d-orbitals of the metal ion by the antibonding π^* -orbitals of CO, the π^* -d interaction appears [54]. This leads to an increase in the binding energy of the ion with CO, which, for Co^{2+} and Ni^{2+} , reaches about 13 kcal/mol [51].

In a series of CO sensors obtained by the hydrothermal method from indium oxide doped with Cu^{2+} , Pd^{2+} , and Ni^{2+} ions, sensors containing Ni^{2+} have the highest sensor effect, which is six times the effect in the absence of additives [61]. Doping of In_2O_3 with Zn^{2+} ions also increases the sensor effect approximately 3.5 times [62].

A similar effect of the interaction energy of CO with doping ions on the sensor effect was observed in sensors based on SnO_2 doped with Zn^{2+} and Ni^{2+} ions [63]. Thus, as in systems with acid–base bonds between the analyzed gas and the sensitive layer, an increase in the energy of CO adsorption bonds, due to the π^* -d interaction of CO with doping ions, causes an increase in the sensor effect.

2.1.3. Influence of the Nature of Doping Ions on the Reactivity of Oxygen Anionic Centers

In the reactions of anions localized on the surface of a metal oxide sensitive layer, the activation energy that determines the sensor effect depends on the binding energy of the anion with the metal oxide. The effect of these bonds was demonstrated in the detection of formaldehyde by a sensor based on In_2O_3 , doped with ions of various metals [64–66]. The Fermi level (see Figure 3) of the sensitive layer increased or decreased depending on the nature of the doping ions.

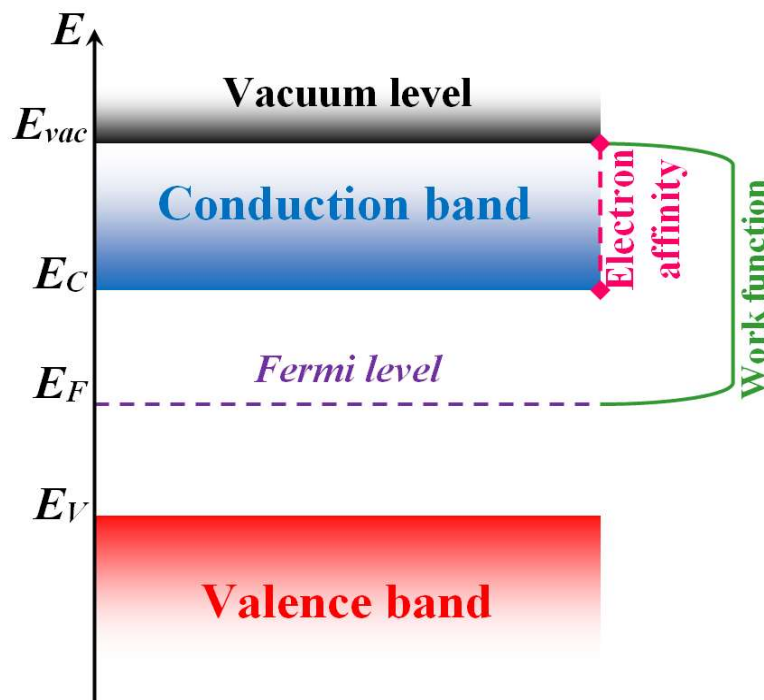


Figure 3. Semiconductor band structure. Here, E_{vac} is minimal vacuum energy of electron, E_C is bottom of the conduction band, E_F is Fermi level and E_V is valence band ceiling.

An increase in the Fermi level leads to an increase in the sensor effect in the detection of formaldehyde and other reducing compounds [59]. This trend can be attributed to the increase in the reactivity of active oxygen anions, due to a reduction in their binding energy to the surface. The reduced binding energy is caused by a decrease in the electron affinity χ_e of the metal oxide sensitive layer from the increase in the Fermi level.

An increase in the Fermi level of the sensitive layer leads to the increase in the concentration of adsorbed oxygen anions. The change in concentration is due to the difference between the Fermi level of the metal oxide and the energy of electrons captured by adsorbed oxygen atoms or molecules, as well as the increase in the number of electrons passing from the semiconductor to oxygen, to achieve equilibrium. An increase in χ_e , on the contrary, causes a decrease in the sensor effect [64]. A similar dependence of the sensor effect on the Fermi level (and χ_e) of the sensitive layer was observed in [55,57].

The influence of the metal oxide layer on the reactivity of the active oxygen particles associated with this layer was considered, as well as the dependence on the average characteristics, such as the electron affinity related to the entire layer [64]. In reality, oxygen molecules are adsorbed on certain lattice defects, where the coordination between metal ions and oxygen ions of the metal oxide lattice is disrupted. In this case, bonds of coordinating-unsaturated metal ions with environmental molecules can be formed. Essentially, such defects are oxygen vacancies [44], whose properties depend on the structure of the crystal lattice and the location of the vacancy. Accordingly, the chemical activity of a vacancy in the sensor process is changed [67].

The properties of vacancies depend on the conditions of metal oxide annealing during the formation of the sensitive layer. Thus, nanocrystalline SnO_2 , obtained by the hydrothermal method of tin hydroxide annealing, contains both isolated oxygen vacancies and clusters of oxygen vacancies, in combination with vacancies of tin ions [68]. The number of vacancies and the ratio between vacancies of different types depends on the medium in which the annealing is carried out—in air, in oxygen, or in helium. In accordance with this, the sensor response to acetone, ethanol, and formaldehyde is changed [68].

As noted in the previous section, the chemical activity of the metal oxide, and the sensor effect in the detection of reducing agents, increase with a decrease in the bond energy between oxygen ions and metal ions. To clarify this mechanism, the sensitivity of MnO_2 nanofibers doped with niobium ions to CO was studied [69]. The Mn–O–Nb sequences formed in doped nanofibers contain oxygen ions weakly bound with the ions of these metals. Therefore, the reaction, $\text{CO} + \text{Mn–O–Nb} \rightarrow \text{CO}_2 + \text{Mn–V}_\text{O}\text{–Nb}$, with the formation of oxygen vacancies (V_O) proceeds efficiently, even at room temperature. This reaction leads to a sharp drop in the resistance of nanofibers, but the reasons for this are not discussed in [69].

It can be assumed that, since the electron levels of the formed vacancies lie near the conduction band of the nanofibers [44,65], electrons from oxygen vacancies easily pass into them. As a result, the conductivity of the composite increases, leading to a sensor effect. After replacing the air mixture, with CO, with pure air, the vacancies are filled with oxygen, and the resistance of the sensor system returns to its original value.

This system has exceptionally high sensitivity to CO, which makes it possible to detect the gas, even at its concentration in air of 2 ppm [69]. A high selectivity of CO detection is thus achieved in the presence of organic compounds, which can be explained by the selective adsorption of CO on Mn–O–Nb structures [70]. These structures are very stable. Therefore, the formation of oxygen vacancies under the influence of CO and their death upon interaction with oxygen are completely reversible processes.

A similar increase in the reactivity of oxygen ions was found for cerium oxide doped with samarium ions [71]. The Ce–O–Sm structures formed in this material also contain labile oxygen ions weakly bonded to the lattice. As a result, the rate of CO oxidation with the formation of oxygen vacancies increases, and the maximum of oxidative activity shifts to lower temperatures. All this testifies to the key role of oxygen vacancies and complexes of vacancies with metal ions in the sensor process.

2.2. Mechanisms for Increasing Sensitivity of Two-Component, Two-Phase Metal Oxide Composites

In contrast to single-phase systems, different conduction paths are formed in binary two-phase systems, by aggregates of component particles contacting with each other. Below, we consider the influence of the nature and structure of conduction pathways, as well as the interaction between components, on the conductivity and sensor properties of nanocomposites. The mechanisms of chemical and electronic sensitization of sensor processes are also discussed.

2.2.1. Effect of Conduction Pathways on Sensor Properties of Binary Two-Phase Nanocomposites

The conductivity of the sensor layer is determined by the aggregates of metal oxide particles that are in contact with each other and form current flow paths. In a two-component system, in which components A and B form different crystalline phases, there are three types of intercrystalline contacts, namely homophase contacts A–A and B–B, and heterophase contacts A–B. As a rule, nanoparticles A and B have different work functions (W), such that, upon their contact, electron transfer occurs at $W_A > W_B$ from B to A, and at $W_B > W_A$, in the opposite direction. Such an electron transfer between nanoparticles corresponds to their mutual charging [27].

An internal electric field develops at heterogeneous contacts. In this case, the potential barrier, that prevents the passage of current through a heterogeneous contact between

particles, depends on the mutual direction of the internal and external fields [72]. In a disordered system of nanoparticles, which includes heterogeneous contacts, when the direction of the external field is opposite to the direction of the field in the contact, high barriers inevitably arise on the current path. The current flow path “bypasses” such contacts, choosing the homogeneous contacts with lower barriers for electron transfer [72].

Thus, in two-phase nanocomposites consisting of particles of components A and B, two parallel homophase conduction paths can be realized through the contacting nanoparticles A-A or B-B. The current flows through chains of contacting nanoparticles, which form conducting clusters that connect the electrodes of the measuring system (“endless” clusters [72,73]). The nanoparticles of the second component, that did not enter the conduction pathways, are modifiers, the interaction with which changes the chemical and electronic characteristics of the conducting nanocrystals, and, consequently, the conductivity and sensor effects.

In a disordered two-phase conglomerate from particles of two components, the conduction paths are determined by percolation theory [73], and depend on the relative content of the components and the ratio between the sizes of the corresponding nanocrystals [72].

In nanostructured composites ranging from *n*- and *p*-modifications of TiO₂, three conduction regions are possible [72], based on the percolation model, depending on the relative content of the components. These are regions I and II, with conducting clusters of nanoparticles of only one type, and region III, in which conducting clusters of both types coexist. In region III, the conductivity of the composite depends on the ratio between the concentrations of components, the size of the particles, and the concentration of current carriers (electrons or holes) in the components [72].

In a nanostructured composite of metal oxides with different work functions, the magnitude and direction of the sensor effect are determined by the electron transfer between the conducting clusters and the modifier. A characteristic example is the result obtained from the study of the conductivity and sensor effect in a composite of SnO₂ nanofibers with nanoclusters of TiO₂ or WO₃ modifiers located on the surface of the nanofibers [74]. In SnO₂ nanofibers and modifier nanoclusters, the work function is different; therefore, the effect of modifiers on conductivity and sensor properties depends on the difference between the W_{SnO_2} and W of the modifier. Modification of nanofibers with TiO₂ nanoclusters ($W_{\text{TiO}_2} < W_{\text{SnO}_2}$) leads to an increase in the conductivity of the composite, due to electron transfer from TiO₂ to SnO₂, while modification with WO₃ nanoclusters ($W_{\text{WO}_3} > W_{\text{SnO}_2}$) reduces the conductivity, due to electron transfer from SnO₂ to WO₃ [74].

The modification of SnO₂ nanofibers also changes sensor effects, in addition to the conductivity. An increase in conductivity under the influence of TiO₂ causes an increase in the sensor effect for oxidizing gases but has almost no effect on the sensor effect for reducing agents. The decrease in conductivity caused by WO₃ nanoclusters also leads to an increase in the sensor effect in the detection of reducing gases [74].

A similar increase in the sensor response to reducing gases, accompanied by a decrease in the conductivity of the composite, is observed for the deposition of WO₃ nanoclusters on the surface of In₂O₃ nanofibers. In contrast, the deposition of In₂O₃ on WO₃ nanofibers increases their conductivity and sensor response to oxidizing gases but reduces the sensitivity of the composite to reducing agents [75].

In numerous reviews on two-phase nanocomposites (see, for example, [76,77]), as in the studies considered above, only systems with conduction paths from particles of one component are mostly discussed. Particles of the second component, interacting with conductive particles, change their properties and affect sensor effects.

Binary nanocomposites with different conduction paths have different sensitivity to detected gases. The authors of [78,79] studied systems consisting of metal or metal oxide nanoparticles in combination with thin nanolayers of reduced graphene oxide (rGO). With minimal coverage of rGO layers by Pd particles, these layers are current flow paths, and the conductivity decreases in the presence of hydrogen. This is due to the transformation of Pd

nanoparticles into PdH_x , followed by electron transfer from PdH_x to rGO and a decrease in the concentration of charge carriers—holes—in the *p*-semiconductor rGO [78].

With an increase in the concentration of Pd particles, the response to H_2 decreases, due to a change in the conduction paths—from rGO layers to clusters of Pd nanoparticles. This occurs during the formation of a continuous Pd film on the surface of the rGO layers. Due to the change in the structure of the Pd film, under the influence of hydrogen dissolved in it, the response increases [78]. In the absence of Pd nanoparticles, the conductivity of rGO layers largely does not react to the presence of nonpolar molecules in the air, for example, H_2 . The effect of current flow paths in nanocomposites combining rGO layers with metal oxide nanoparticles on conductivity and sensor properties was studied in detail using the response to ammonia as an example [80].

As rGO was added to nanostructured SnO_2 , the type of conductivity of the composite changed from electronic conductivity, characteristic of pure SnO_2 , to hole conductivity, characteristic of rGO. Due to the difference in the shape of the components (thin, extended rGO nanolayers and quasi-spherical SnO_2 nanoparticles), the percolation transition from *n*- to *p*-composites occurs already at small (about 0.5 wt.%) additions of rGO to SnO_2 .

These experimental results agree with the calculations [80]. The response of the sensor to ammonia is due to its reaction with oxygen anions on the surface of SnO_2 particles. In *n*-type nanocomposites, where the current flows through clusters of SnO_2 nanoparticles, the addition of ~0.5 wt.% rGO leads to an increase in the electronic conductivity of the composite [80]. Since W_{rGO} (5.2 eV) of a completely reduced graphene oxide film [81] is larger than W_{SnO_2} (4.9 eV), electrons pass from SnO_2 to rGO. As a result, the electron concentration in conducting clusters of SnO_2 nanoparticles decreases, which increases the response to reducing gases [74,82].

In connection with the mutual charging of contacting SnO_2 nanoparticles and rGO nanolayers, during the transition of electrons from SnO_2 to rGO, favorable conditions are created for the adsorption of polar molecules in the contact region in the nanocomposite. This also applies to NH_3 molecules and additionally increases the sensitivity of the composite sensor to ammonia. An increase in rGO concentration to 1 wt.% leads to a percolation transition from conduction through SnO_2 particles to conduction paths through rGO layers; that is, to hole conduction [80].

The paths of hole and electron conduction are different in nanostructured metal oxides [3,77,83]. The electron current flows through contacting nanoparticles, crossing their surface layers where potential barriers are formed between the particles, due to oxygen ionosorption accompanied by the capture of electrons from the conduction band. The resistance of the surface layers is significantly increased, compared to the resistance in the bulk of the particles. The height of such barriers on the current path is determined by the number of electrons captured by oxygen, which decreases when O^- interacts with a reducing gas. This leads to an increase in conductivity.

In the case of hole conduction, the current flows mainly in the near-surface layers of *p*-nanoparticles, wherein, the negative charge of O^- on the surface is compensated by the positive charge of holes in the near-surface layer. The concentration of holes in this layer is naturally higher than in the volume of the particles [83,84]. The total resistance to the hole current in the volume of the particles and surface layers is determined by the morphology of the nanostructured metal oxide [85].

At the hole conduction, the total resistance (R_{sen}) of the sensor layer includes a volume component which is largely independent of the processes on the particle surface. Therefore, the sensitivity to reducing gases in such systems is significantly less than in the case of electronic conduction, when the change in resistance is determined by the reactions of the analyzed gases with O^- on the surface of the particles. SnO_2 -rGO sensors to ammonia, with hole conduction along the paths formed by rGO layers, are characterized by greater stability and a shorter response time to ammonia, and relaxation time after its removal [80].

The conductivity and sensor properties of nanocomposite fibers containing *n*- SnO_2 and *p*- Co_3O_4 nanoparticles were studied in [23]. The resistance of the $x\text{SnO}_2 - (1 - x)\text{Co}_3\text{O}_4$

system and its sensor response to CO, C₆H₆, and C₃H₆O reach their maximum values at $x = 0.5$. The nanofiber of this composition has electronic conductivity. Since the arrangement of nanoparticles in a fiber is chaotic, the conductivity of a nanocomposite fiber is determined by the percolation model [72,73].

According to this model, the electron current in such a system flows along *n*-type homophase paths, excluding heterophase *n-p* contacts, the inclusion of which in the current path would lead to the occurrence of large potential barriers. Heterophase contacts between *n*- and *p*-nanoparticles of composite lead to the loss of electrons, due to electron-hole recombination. As a result, the electron concentration in the conducting channels decreases. In some cases, this leads to an increase in the response to reducing gases [83].

Similar effects are also characteristic of other *n-p* nanocomposites [86,87], in which the maximum response is achieved in the systems with electronic conductivity. The sensitization of the sensor response by *p*-nanoparticles contacting with *n*-nanoparticles of conducting pathways is due to a decrease in the electron concentration in conducting pathways, resulting from electron-hole recombination. At *n-n*-contacts, the concentration of conduction electrons does not decrease, but the electrons pass into a lower-energy conductivity band, forming an “accumulation layer” of electrons in acceptor centers [83,88]. Such a transition also reduces the concentration of conduction electrons in the conducting paths of the composite, but to a lesser extent since the transition of electrons is hindered by the resulting electric field. Therefore, *n-p* contacts are more effective for sensitizing the composites undergoing the electronic conductivity.

2.2.2. Effect of Interaction between Conducting and Modifying Nanocrystals on Sensor Properties of a Composite

In binary systems, the interaction of nanoparticles, constituting current flow paths with modifier nanoparticles surrounding these paths, is a source of sensor sensitization. There are two types of sensitizations: chemical and electronic [13,89].

Chemical sensitization is caused by catalytically active particles of the modifier, which chemisorb oxygen and molecules of detected gases with the formation of reactive particles, in particular, atoms or radicals. The most common variant of chemical sensitization is the adsorption and dissociation of oxygen molecules on modifier particles with a subsequent spillover of oxygen atoms into conducting clusters. After that, oxygen atoms capture electrons from the conduction band of nanoparticles constituting conducting clusters, forming O[−] radical anions.

Since the reaction of reducing gases with O[−] on the surface of conducting clusters “releases” trapped electrons and increases the conductivity of the composite, the concentration of adsorbed oxygen should be increased to raise the sensitivity. This is clearly facilitated by the catalytic activity of modifier nanoparticles [14,90].

The detection of several π -electron compounds, by sensors based on SnO₂ nanofibers modified with Au, Pd, and Pt nanocrystalline particles, was studied to determine the optimal conditions for achieving the maximum sensitivity [91]. The study indicated that the adsorption bonds, of π -electron molecules with metal particles, are determined by the location of the metal d-electron band, relative to the π -electron orbitals of the molecules of the analyzed gases. Therefore, by varying the electronic structure of the contact between a gas molecule and a metal particle on the surface of SnO₂ nanofibers, chemisorption can be increased. The spillover of adsorbed molecules from metal particles onto SnO₂ fibers increases the gas concentration on nanofibers and the sensitivity of the composite. The conditions were determined for the most efficient chemical sensitization of the sensor effect by Au, Pd, and Pt particles in the detection of CO, benzene, and toluene [91].

Chemical sensitization depends both on the structure of the detected molecule and the properties of the catalytically active particle. A typical example is the sensitization of CO and H₂ detection by a sensor based on nanocrystalline In₂O₃, containing nanoparticles of the well-known catalyst, ZrO₂. The particles of ZrO₂ increase the sensor effects in the detection of H₂, but practically, do not affect the detection of CO [92].

The effect of chemical sensitization of the sensor response to CO on this system could be expected, due to the dissociation of O₂ on the surface of ZrO₂ nanoparticles with the subsequent formation of O[−] active centers. However, the dissociation of O₂ requires a temperature above 800 °C [93]. The efficiency of H₂ detection is probably ensured by the dissociation of H₂ molecules under the influence of ZrO₂. As a result of the spillover of the formed hydrogen atoms onto In₂O₃ particles, H atoms react with O[−] radical anions localized on the In₂O₃. In this case, the electrons captured by oxygen return to In₂O₃.

The conductivity of a nanofiber network is determined by potential barriers to electron transfer between contacting nanofibers, and depends on the electron concentration in their surface layers (n_s) [83]. The decoration of SnO₂ fibers, with electron-donating TiO₂ ($W_{\text{TiO}_2} < W_{\text{SnO}_2}$) or electron-withdrawing WO₃ ($W_{\text{WO}_3} > W_{\text{SnO}_2}$) nanoclusters, leads to a redistribution of conduction electrons between conducting channels and decorating clusters [74]. The transfer of electrons from SnO₂ nanofibers to WO₃ nanoclusters reduces n_s and increases the barriers for electron transfer between contacting nanofibers [83]. All this affects the sensor effect [68], and can be considered as electronic sensitization.

The sensor properties of nanocomposites depend on the method of their formation, which determines their morphology. In this regard, the indicators are the results of studies of composites obtained by physical mixing of oxide nanopowders or by laser sputtering of one oxide on the surface of crystals of another [21,22,94–96]. In such composites based on electron acceptor SnO₂ or ZnO oxides, electron-donating additives of In₂O₃ or TiO₂ nanoparticles do not decrease the sensor effect for reducing gases, as in [74,97,98], but increase it.

It should be noted that the sensor characteristics of binary composites obtained by mixing nanocrystalline powders, in contrast to the systems considered in [74,97,98], are mainly determined by the contact of particles with different electron affinities and clearly marked interphase boundaries [94], where the charges, formed as a result of mutual charging of such particles, are localized. Charges in the interfacial regions of the nanocomposite, as mentioned above [80], can increase the chemisorption of oxygen and detected gases and, thus, increase the sensor effect.

The effect of interfacial contacts on the sensor properties of nanocomposites was not considered in [74,75]. This is probably due to the small size of electron-donor and electron-acceptor clusters located on the surface of conducting nanofibers, because of which, the distribution of electrons in the region of these contacts is largely blurred.

It can be assumed that the differences in the sensor properties of nanocomposites obtained by different methods are due to the structure of the interfacial regions in such composites, which determines the adsorption of gases and their activity in sensor reactions. The largest increase in the sensor effect is achieved due to the joint effect of chemical and electronic sensitization by chemically active modifiers with a high work function [90].

Such sensitization of the sensor effect occurs, in particular, as a result of the modification of the surface of conducting metal oxide particles by nanoclusters of catalytically active metals, Au, Pd, and Pt [91]. Nanoclusters of these metals, as shown above, catalyze the formation of highly reactive atoms and radicals on the surface of metal oxide nanocrystals, which corresponds to chemical sensitization. These nanoclusters, which have a higher work function than conductive metal oxide nanocrystals, capture the electrons of the nanocrystals, lowering n_s . Thus, there is also an electronic sensitization of the sensor effect.

A characteristic example of the synergism of electronic and chemical sensitization is the detection of ethanol vapor by sensors based on In₂O₃ nanotubes, decorated by nanoclusters, Co₃O₄, Fe₂O₃ or a combination of these nanoclusters [99]. The deposition of nanoclusters, of even a single metal oxide on the surface of nanotubes, increases the sensor effect due to electron sensitization, since the work function of these metal oxides is higher than that of In₂O₃. In this case, Co₃O₄ clusters have a greater influence, although W for Fe₂O₃ (5.3 eV [100]) is higher than for Co₃O₄ (4.8 eV [101]).

Co_3O_4 nanoclusters are both electronic and chemical sensitizers, which is due to the catalytic activity of Co_3O_4 in the dissociation reactions of adsorbed oxygen molecules with the formation of O^- . The largest increase in the sensor effect occurs with the joint application of Co_3O_4 and Fe_2O_3 , when high chemical sensitization of the sensor effect under the action of Co_3O_4 is combined with high electronic sensitization under the influence of Fe_2O_3 [99].

The synergism of the sensitizing action of different agents is also manifested in sensors based on porous In_2O_3 nanospheres, modified with NiO and Au nanoclusters [102]. The interaction of NiO clusters with In_2O_3 leads to the formation of oxygen vacancies on the surface of In_2O_3 in such systems, due to the difference in the charges of Ni^{+2} and In^{+3} ions. Concurrently, Au nanoclusters catalyze the dissociation of these molecules, with the capture of electrons and the formation of O^- active centers.

The combined action of NiO and Au clusters on In_2O_3 nanoparticles produces high sensitivity in the detection of reducing compounds, in particular, toluene [102]. Using such a sensor, toluene can be detected, even at concentrations of the order of 100 ppb, since Au clusters adsorb toluene and thus increase the sensor effect. A similar synergistic effect of electronic and chemical sensitization of the sensor effect was also found for In_2O_3 nanotubes containing NiO and PdO nanoclusters on the surface [39].

The results presented in this section have shown that the sensor response for two-component, two-phase metal oxide composites can differ markedly for various gases. Table 1 summarizes the results of previous investigations and makes it possible to compare the response magnitudes to various gases of sensors synthesized with different methods under diverse conditions.

Table 1. Sensor response to various gases of two-component two-phase metal oxide composites.

Material	Synthesis Method	Sensor Response (Gas Concentration)	Temperature	References
$0.5\text{SnO}_2\text{-}0.5\text{Co}_3\text{O}_4$	electrospinning method	18.7 (1 ppm C_6H_6)	350 °C	[23]
$\text{WO}_3\text{-SnO}_2$	sputtering high-purity Ti or W targets on SnO_2	140 (1 ppm H_2)	300 °C	[74]
TiO-SnO_2		232 (1 ppm O_2)		
$\text{WO}_3\text{-In}_2\text{O}_3$	sol-gel method	27 (200 ppm NO_2)	300 °C	[75]
graphene-Pd/ SnO_2 composites	grapheme by CVD method, SnO_2 gas phase synthesis method	14.8% (1% $\text{C}_2\text{H}_5\text{OH}$)	200 °C	[79]
rGO- SnO_2	rGO by hydrothermal treatment of aqueous dispersion of GO, rGO- SnO_2 composite by mixing	1.3 (50 ppm NH_3)	22 °C	[80]
$0.5\text{SnO}_2\text{-}0.5\text{NiO}$	electrospinning process	36 (10 ppm NO_2)	300 °C	[86]
$\text{SnO}_2\text{-}2.78\text{CuO}$	sol-gel route	200% (400 ppm CO)	350 °C	[87]
$\text{In}_2\text{O}_3\text{-Co}_3\text{O}_4$	Electrospinning method	39 (200 ppm HCHO)	260 °C	[88]
Pt- SnO_2	photolithographic process and γ -ray radiolysis method	40 (1 ppm C_7H_8)	300 °C	[91]
$10\%\text{Co}_3\text{O}_4\text{-}90\%\text{In}_2\text{O}_3$	Mixing metal oxides	1300 (1100 ppm H_2)	250 °C	[92]
$20\%\text{ZrO-}80\%\text{In}_2\text{O}_3$		280 (1100 ppm H_2)	315 °C	
$\text{TiO}_2/\text{SnO}_2$	hydrothermal process	52.3 (100 ppm triethylamine)	260 °C	[96]
$\text{ZnO@In}_2\text{O}_3$	hydrothermal method	28.6 (100 ppm $\text{C}_2\text{H}_5\text{OH}$)	160 °C	[97]
$\text{Au-NiO/In}_2\text{O}_3$	solvothermal method	80.6 (10 ppm toluene)	250 °C	[102]

The consideration of the processes of chemical and electronic sensitization of sensor reactions has shown that such a division is rather arbitrary. For example, oxygen molecules dissociate on the surface of nanoparticles with the formation of chemically active

centers, which corresponds to chemical sensitization, wherein, the formed oxygen atoms actively capture electrons from the near-surface layer, thereby depleting it; this is already characteristic of electron sensitization. Thus, in various sensor processes, both types of sensitization mechanisms can occur simultaneously, but the chemical or electronic nature will be manifested to a greater extent.

2.3. Interaction of Nanocomponents in Core–Shell Sensor Systems

Two-phase structures of the core–shell type have been intensively studied in the last decade. Such structures consist of a core of metal oxide covered by a layer of another semiconductor oxide (see, for example, reviews [14,103–106]).

With the right choice of core and shell materials, such structures can achieve exceptionally high sensitivity and selectivity in the detection of various substances. This also makes it possible to significantly lower their operating temperature [14,104,105,107–109]. These materials have generally different morphologies, but the highest sensitivity is exhibited by structures formed by composite nanofibers (see Figure 4). The sensor characteristics of such a material depend on the path of current flow. According to the percolation model of conductivity in a disordered system of semiconductor particles with different electron work functions [72,73], the current in the sensitive layer passes through the fiber shell, overcoming the barriers of the homocontacts, but bypassing the potential barriers of heterocontacts in the core–shell system [110].

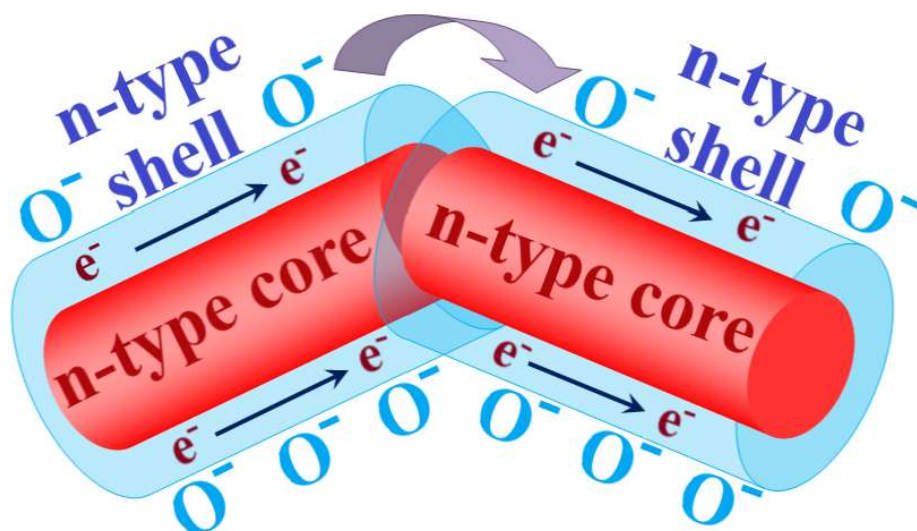


Figure 4. Scheme of the sensitive layer of composite nanofibers of the core–shell type. « $e^- \rightarrow e^-$ » indicates the current flow in the shell of nanofibers.

Such a scheme has been confirmed by the results obtained from the detection of a reducing agent (benzene) and an oxidizing agent (NO_2) by composites of $n\text{-ZnO-p-CuO}$ nanofibers [111] (here and hereinafter, the first oxide is the core, and the second is the shell). The conductometric sensor response to these compounds in the sensitive layer of $n\text{-ZnO-p-CuO}$ nanofibers indicates hole conductivity through the nanofiber shells without crossing the fibers. In this case, the value of the sensor response, due to the features of hole conductivity [112], is much lower than in the sensitive layer from pure zinc oxide nanofibers with electronic conductivity [111]. An increase in the sensor response, when $p\text{-CuO}$ nanoparticles are included in such a layer, is due to a change in the electronic conductivity as a result of contacts between $n\text{-ZnO}$ and $p\text{-CuO}$ nanoparticles and electron transfer from ZnO to CuO.

The resistance of the sensitive layer, as in other nanostructured metal oxide semiconductors, is determined by the resistance of contacts between nanoparticles (in this case, nanofibers), which in systems with electronic conductivity increases with a decrease in the electron concentration in the shells.

Two factors affect the concentration of electrons in the shell nanofiber. The first factor is the capture and localization of conduction electrons by electron-withdrawing centers on the outer and inner surfaces of the shell, which leads to the depletion of the shell in electrons. On the outer surface, such centers are adsorbed forms of oxygen, which, at the “operating” temperatures of the sensors above 200 °C, are mainly oxygen atoms. On the inner surface separating the nanofiber shell from the core, electrons are trapped by lattice defects that arise during the formation of the composite, due to the difference in the structures of the core and shell [113].

Another factor is associated with the presence of electron heterojunctions between the core and shell. Such a transition is determined by the difference in the values of the electron work function W for the core W_C and the shell W_{Sh} of the fiber. If $W_{Sh} < W_C$, conduction electrons pass from the shell to the core. In this case, the shell of the composite fiber is even more depleted of electrons.

According to existing concepts, the most promising are sensors based on core-shell nanofibers, which contain weakly conductive catalytically active layers of SnO_2 or ZnO . Modification of core-shell metal oxide systems with noble metal nanoparticles can have a significant effect on their sensor response and selectivity. Sensors based on such nanofibers make it possible to detect even traces of various organic compounds in the air (see, for example, [114–117]).

The response to 0.1 ppm CO of core-shell sensors, based on SnO_2 - ZnO nanofibers with Au nanoclusters located on the surface of ZnO shells, is 25, while for pure SnO_2 and ZnO nanofibers, it is only 3 and 2, respectively [114,118]. The maximum sensor response in the detection of 100 ppb benzene, by SnO_2 - ZnO nanofibers with Pd nanoclusters on the surface of ZnO shell, is 71 [116]. This is much higher than the sensor effect for SnO_2 (1.9) and ZnO (2.1) nanofibers, as well as for SnO_2 - ZnO sensors that do not contain Pd clusters (11.2). It is also interesting that the deposition of silver nanoparticles on the surface of a nanosystem consisting of tungsten oxide ($\text{W}_{18}\text{O}_{49}$), included in a polypyrrole shell, leads to an increase in the sensor response to ammonia and an improvement in the selectivity of this system [115].

To determine the mechanism of sensor effects in composite nanofibers, it is important to find out how the electrical and sensor characteristics of fibers change when the material of the core and shell are swapped. Sensor effects in the detection of CO and aromatic hydrocarbons were determined for core-shell ZnO - SnO_2 nanofibers [118]. A comparison with data for similar layers of SnO_2 - ZnO nanofibers [109] makes it possible to elucidate important features of the resistance and sensor effects in these systems.

In both cases, shells were formed on monocrystal nanorods by deposition of atomic layers. According to [108,109], the interaction between the core and shell in SnO_2 - ZnO nanofibers leads to an increase in the resistance of the sensor system, due to the contacts between the layers. The sensor effect is also increased.

The values of resistance and the sensor effect of core-shell composites largely depend on the thickness of the shell. For the SnO_2 - ZnO nanocomposite, the maximum values of these parameters are achieved at a shell thickness of about 30 nm; that is, close to the Debye charge screening length in ZnO (λ_{ZnO}), which is 22–35 nm [116]. A similar dependence on the shell thickness was also observed in another sensor system of the same type.

The maximum sensor response of the Nb_2O_5 - ZnO system of the core-shell type to 1% H_2 at 300 °C is observed at a ZnO shell thickness of about 50 nm, which is close to $2\lambda_{\text{ZnO}}$ [119]. These data indicate the presence of electron-withdrawing defects in the Nb_2O_5 - ZnO system at the interface core-shell, which reduce the electron concentration in the shell. Therefore, the length of the screening of charges on oxygen centers, formed on the surface of the shell during oxygen chemisorption, increases. Meanwhile, the reaction of these centers with the analyzed reducing agent and the “release” of trapped electrons, respectively, causes a sensor response, as evidenced by an increase in conductivity.

Judging by the data on the resistance of SnO_2 - ZnO and ZnO - SnO_2 nanofiber layers [114,117], the concentration of conduction electrons in the shells of these nanofibers

decreases mainly due to defects at the boundary between SnO_2 and ZnO. The transfer of electrons between SnO_2 and ZnO increases or decreases the electron concentration in the shells, depending on which of these materials forms the core or shell of the nanofibers. The value of the sensor effect, in the detection of reducing compounds (CO , C_6H_6 , C_7H_8) by SnO_2 -ZnO and ZnO- SnO_2 sensors, changes according to the change in the resistance of the sensitive layer [114,117].

The attribution of the large value of the sensor effect in core-shell nanofibers, when both the core and the shell have catalytic activity, to only the influence of the system resistance, is not sufficiently substantiated. Indeed, in [116–118], a significant increase in the sensor effect was observed when nanoclusters of noble metals were located on the surface of shells, although such additives only reduce the resistance of the film. In addition, a very noticeable sensitivity and selectivity was found in the detection of ethanol by In_2O_3 -ZnO nanofibers [120] containing In_2O_3 core with a very high electron concentration, which can pass into the ZnO shell, since the work function of the electron for In_2O_3 is lower than for ZnO. The sensor effect for composite nanofibers is 10 times that for In_2O_3 nanofibers. In this case, In_2O_3 , as in mixed systems of nanoparticles, is an electron supplier, while ZnO is a catalytically active agent.

The study of the sensor layer based on core-shell nanofibers was also carried out for the ZnO- SnO_2 system, which has a slightly different crystal structure. These fibers consist of a monocrystal ZnO core and a continuous SnO_2 shell, formed during the thermal reaction of $(\text{CH}_3)_4\text{Sn}$ with oxygen on the ZnO surface [121]. The sensitivity of such a layer to reducing gases (CO , CH_4 , H_2 , $\text{C}_2\text{H}_5\text{OH}$) at 200 and 300 °C is very low and almost does not differ from the sensitivity of a layer of ZnO nanofibers. An increase in the sensor effect, under the influence of the SnO_2 shell located on the surface of ZnO nanofibers, manifests itself only when the temperature rises to 400 °C. Therefore, in ZnO- SnO_2 nanofibers [121], conduction electrons, trapped by defects at the interface between the core and shell, are held more strongly in defects than in the ZnO- SnO_2 and SnO_2 -ZnO nanofibers considered above, with shells formed by the deposition of atomic layers [114,117].

Thus, the sensor properties of composite metal oxide nanofibers of the core-shell type strongly depend on the composition and method of syntheses of the nanofibers (Table 2) and, accordingly, on their morphology. A detailed study of such systems will elucidate the role of the interaction between the core and shell in the sensor process, and will contribute to the development of new, highly efficient sensor materials.

Table 2. Sensor responses of core-shell materials to various gases.

Material	Synthesis Method	Sensor Response (Gas Concentration)	Temperature	References
Ag- α - Fe_2O_3 core-shell composites	two-step reduction-sol gel approach, including Ag nanoparticles	9 (500 ppm)	250 °C	[105]
Au-ZnO core-shell nanoparticles	facile low-temperature solution route	103.9 (100 ppm H_2)	300 °C	[107]
SnO_2 -ZnO core-shell nanowires	two-step process	25 (10 ppm NO_2) 75 (10 ppm C_7H_8) 83 (10 ppm C_6H_6) 77 (10 ppm CO)	300 °C	[108]
SnO_2 -ZnO core-shell nanofibers	two-step process	6.5 (1 ppm CO) 48 (1 ppm NO_2)	300 °C	[110]
ZnO-CuO core-shell nanowires	facile three-step process	29 (10 ppm NO_2)	350 °C	[111]
In_2O_3 /ZnO core-shell nanowires	thermal evaporation of indium powder in an oxidizing atmosphere, followed by the atomic layer deposition of ZnO	196 (1000 ppm $\text{C}_2\text{H}_5\text{OH}$)	300 °C	[113]

Table 2. Cont.

Material	Synthesis Method	Sensor Response (Gas Concentration)	Temperature	References
SnO ₂ -ZnO core-shell nanowires functionalized by Au nanoparticles	Vapor-liquid-solid growth method	26.6 (100 ppb CO)	300 °C	[114]
Ag functionalized W ₁₈ O ₄₉ @PPy core-shell nanorods	polymerizing the uniform PPy shell film on surface of W ₃ nanorods with AgNO ₃ as oxidant and DBSA as modifier.	2.5 (20 ppm NH ₃)	40 °C	[115]
Pd-functionalized SnO ₂ -ZnO core-shell nanowires	two-step growth technique. Pd functionalized SnO ₂ -ZnO by using the ray radiolysis technique	71 (100 ppb C ₆ H ₆)	300 °C	[116]
SnO ₂ -ZnO core-shell nanowires functionalized Pt nanoparticles	two-step growth technique. Pt functionalized SnO ₂ -ZnO by using the ray radiolysis technique	279 (100 ppb C ₇ H ₈)	300 °C	[117]
ZnO-SnO ₂ core-shell nanowires	two-step growth technique	41.13 (10 ppm CO) 39.48 (10 ppm C ₇ H ₈) 40.34 (10 ppm C ₆ H ₆)	300 °C	[118]
Nb ₂ O ₅ /ZnO core-shell nanorod	two-step growth process	156 (100 ppm H ₂)	300 °C	[119]
In ₂ O ₃ -ZnO core-shell nanowires	two-step growth process	265 (400 ppm C ₂ H ₅ OH) 7 (2000 ppm H ₂)	350 °C	[120]
ZnO-SnO ₂ core-shell nanowires	two-step vapor growth method	66.3 (10 ppm NO ₂)	200 °C	[121]

3. Modeling of the Electronic Subsystem of Semiconductor Nanoparticles and Properties of Associated Sensitive Layers

The operation mechanism of semiconductor sensors has been described in many studies [10,14,82]. Due to the presence of oxygen vacancies in the oxide nanoparticles that make up the sensitive layer, a noticeable electron density appears in the conduction band. Oxygen molecules from the air are adsorbed on the surface of nanoparticles and dissociate. The resulting oxygen atoms, which are much deeper electron traps than oxygen vacancies, capture electrons from the bulk of the nanoparticles, and the conductivity of the sensitive layer decreases sharply. After the addition of the analyzed gas, its molecules are adsorbed on the surface of the nanoparticles and react with oxygen ions. The captured electrons are subsequently released and returned to the conduction band of the nanoparticle. The conductivity of the sensitive layer increases, constituting the sensor effect.

Many articles have been published on the modeling of the charge distribution in semiconductor nanoparticles and sensor processes. A discussion of these studies, with reference to the numerous assumptions made within them, including erroneous ones, is presented in a review [14]. The most notable of the erroneous assumptions are the use of a flat instead of a real boundary between nanoparticles, the absence of a relationship between the number of oxygen ions on the surface and the electron concentration in the bulk, and the assumption that the density of positively charged donors is constant along the nanoparticle radius, which is true only in the case of fully ionized donors.

In early studies aimed at determining the distribution of electrons in a nanoparticle, it was assumed that positive charges are immobile and, most importantly, that the concentration of electrons in the negatively charged layer on the surface of nanoparticles was chosen arbitrarily. The role of this layer was clearly overestimated.

The nanoparticles that make up the sensitive layers can be conditionally classified into two types, depending on the concentration of conducting electrons. The first type consists of the nanoparticles with a low electron concentration in the conductive zone. Thus, in sensors

based, for example, on SnO_2 , the concentration of conduction electrons is $n_c < 10^{16} \text{ cm}^{-3}$. At a diameter of 100 nm, one such nanoparticle contains no more than one electron, and a negatively charged layer cannot be created. Thus, there is no interference, as a result of shielding electron transfer between nanoparticles.

In another type of nanoparticles at operating temperatures, the sensitive layer contains a sufficiently large number of conduction electrons. In a nanoparticle of the same size (for example, In_2O_3 , in which $n_c \approx 10^{19}–10^{20} \text{ cm}^{-3}$ [122,123]), the number of electrons can reach $\sim 10^4–10^5$. Since a significant part of these electrons is captured on the surface of nanoparticles by oxygen atoms, this leads to the appearance, in this case, of a negatively charged layer. Assuming that the nanoparticles are spherical and that the surface is charged uniformly, we use the well-known Gauss theorem, according to which, an electrical charge uniformly distributed on a spherical surface does not create a field inside the sphere. If, on the whole, the nanoparticle is electrically neutral, then the field outside it is also equal to zero.

The electric field inside the nanoparticle is created, due to the uneven distribution of negative and positive charges inside the nanoparticle [31]. The current in the system is determined by the state of electrons in the near-surface layer. To determine the efficiency of the sensor, it is necessary to calculate the distribution of charges over the radius of the nanoparticle, in both the presence and absence of the studied gas. This will make it possible to compare theoretical and experimental results on the sensitivity of sensors based on both single and mixed oxides.

3.1. Distribution of Electrons in Oxide Nanoparticles

In sensors based on nanoparticles with a low concentration of conduction electrons (for example, SnO_2), there is no negatively charged layer, the distribution of electrons over the volume is uniform, and there is no problem of finding the charge distribution in the nanoparticle.

For sensors with a high concentration of conduction electrons, the situation is much more complicated. To find the distribution of electrons along the radius of a nanoparticle, one can use the free energy of the system, which consists of the free energies of conduction electrons and electrons on adsorbed oxygen atoms, as well as the potential energy of positively charged ionized donors and the interaction of all negative and positive charges of the nanoparticle [31]. The total free energy is minimized with respect to the chemical potential. Further, the resulting equations are solved, together with a system of stationary equations for the quantity of oxygen atoms, ions, and molecules on the surface of nanoparticles [31,35].

As a result, for nanoparticles of different radii, one can find the distribution of conduction electrons along the nanoparticle radius as a function of temperature. The distribution is found to be inhomogeneous, and the inhomogeneity increases with increasing temperature. This is explained by an increase in the number of electrons on the surface at increasing temperature [37].

3.2. Sensor Effect in One-Component Systems

A theory has been constructed that describes the sensor properties of semiconductor oxides containing both small and large quantities of conduction electrons. In the case of one-component sensors, terms and equations for the detected gas are added to the system of stationary kinetic equations for the concentrations of oxygen atoms, ions, and molecules. The solution of this system, together with the equations for the concentration of conduction electrons, makes it possible to find the sensitivity of the sensor as a function of the pressure of the detected gas, the temperature, and the average radius of the nanoparticles [32–35].

The theory developed was compared with the experimental data. For example, in [33] a comparison was made of the theoretical and experimental results obtained in a study on the dependence of sensitivity to different hydrogen concentrations on temperature for sensors with large [124] and small [33] average radii of SnO_2 nanoparticles. In addition,

the dependence of the sensitivity of a sensor based on tin oxide nanoparticles, on their size, was experimentally studied [125]. It has been shown that the maximum sensitivity of the sensor decreases with an increase in the average size of the nanoparticles. Calculations of the dependences of the sensitivity of the SnO₂ sensor, on temperature and nanoparticle size, showed good agreement with experiment [32,33].

Sensors with a high concentration of conduction electrons have also been studied [34,35]. The technique of determining the electron distribution along the nanoparticle radius has been used for comparing theory with experiment [31]. The electron concentration in the near-surface layer was calculated in the presence and absence of the studied gas. The ratio of these concentrations corresponds to the sensor effect. The study showed a fairly good agreement between the experimental and theoretical results, at reasonable values of the fitting parameters [34,35]. Some discrepancy between theory and experiment is explained by the fact that the calculations were carried out for spherical nanoparticles, which, in fact, differ markedly from reality [35]. In addition, the study assumed average particle radius, without consideration for the dispersion of nanoparticles.

3.3. Electron Distribution and Sensor Effect in Two-Component Systems

The distribution of particles and the sensor effect are determined by different types of interaction between nanoparticles in two-component semiconductor systems [19,37,39,126]. There are many experimental studies devoted to the sensor properties of mixed oxides (see, for example, reviews [13,14,103]). The influence of one nanoparticle on the electronic structure of another largely determines these properties. A theoretical consideration of the distribution of electrons in a mixed system was carried out for the first time in [37]. The distribution of electrons in In₂O₃ nanoparticles surrounded by CeO₂ nanoclusters was studied. The interaction of nanoobjects leads to the effective dissociation of O₂ molecules on the surface of CeO₂ nanoclusters [127], followed by the transfer of oxygen atoms to neighboring In₂O₃ nanoparticles and the capture of electrons from their bulk [126]. It should be noted that in experiments where the CeO₂-In₂O₃ sensor system was studied [14,20,103,126,128–130], the result did not give a simple summation of sensor effects on CeO₂ and In₂O₃ nanoobjects but showed their synergistic interaction.

The experimental and theoretical results on the CeO₂-In₂O₃ sensor system were subsequently compared [131]. A nanostructured system, corresponding to the insert in Figure 5, was considered. The distribution of electrons in the mixed nanocomposites [37] can be used to describe the experimental results [126].

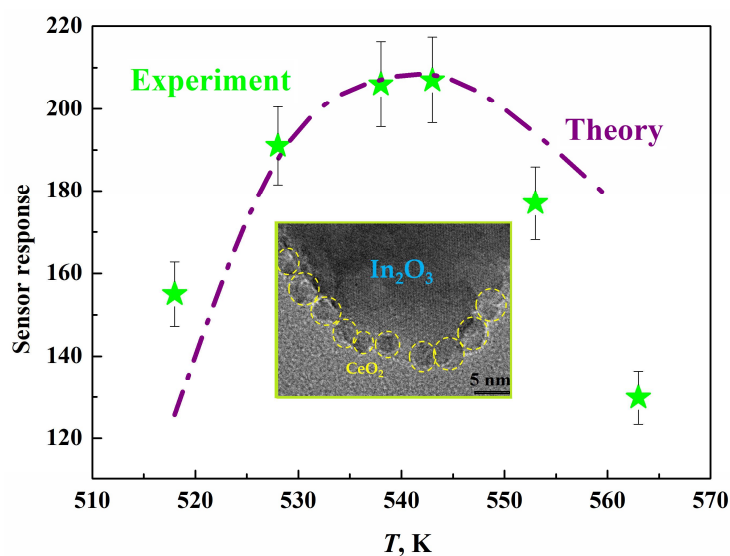


Figure 5. Temperature dependence of sensor response of impregnated 3% CeO₂-97% In₂O₃ system to hydrogen. Green stars are experimental data [127] and dot-dash line is the theory; the insert is the TEM structure of composite.

Using the mechanism of nanoparticle interaction considered above, it was possible for the first time to describe the temperature dependence of the sensor effect (Figure 5) in a mixed system [132].

4. Mechanisms of Sensor Selectivity

One of the main characteristics of a sensor is the selectivity of its response, S_A , to the detected chemical compound. In the case of reducing compounds, which are considered in this review, the selectivity is determined mainly by two factors: (i) the selectivity of the adsorption of oxygen molecules and analyzed molecules from air in the near-surface sensor layer, and (ii) the different activity of the adsorbed molecules in the reaction with oxygen anionic centers (mainly O^-), capturing the conduction electrons of this semiconductor layer.

At present, there is considerable interest in the detection of harmful anthropogenic substances, the accumulation of which in air is associated with the rapid development of industry [132–134]. The selectivity of the detection of such substances is extremely important for the development of methods for protecting the body from the harmful effects of the external environment.

4.1. Selectivity of Molecule Adsorption on Surface of Sensor Layer

Harmful anthropogenic substances include reducing organic compounds. The reaction of reducing agents with oxygen anion centers on the surface of a semiconductor (mainly metal oxide), which is used for the selective detection of these compounds by the conductometric method. The oxygen anion centers are formed when oxygen molecules capture electrons from the conduction band of the semiconductor.

This process can be conditionally divided into two stages: the adsorption of reagent molecules, and the spillover of the adsorbed molecules from the adsorption center to the oxygen center, accompanied by the reaction of this molecule with the “release” of the trapped electron and its transition to the conduction band of the semiconductor. Such a distinction is valid if the formation of adsorption bonds, between molecules and semiconductor nanoparticles of the sensitive layer of the sensor, does not significantly affect the chemical properties of the molecules, including the reactivity of molecules with respect to the oxygen centers of the sensitive layer.

It is assumed that this feature of adsorption bonds is typical for the interaction of alcohols, aldehydes, and ketones with a metal oxide semiconductor sensor layer. For the selective detection of these compounds, the Brønsted acids are used as nanostructured layers, modified with inclusions of metal oxides with highly basic ions, especially alkali and alkaline earth metal ions. These ions are introduced into the lattice of oxide nanocrystalline particles of the sensitive layer, and form adsorption centers for acid-type compounds. The energy of adsorption bonds, in this case, increases with a decrease in the electronegativity and, consequently, with an increase in the basicity of the introduced ions [135].

The study of nanocrystalline metal oxide layers containing these ions showed that, for such layers, the sensor effect varies, depending on the acidity of adsorbed oxygen-containing compounds, which is determined by their structure. This dependence is the main selectivity factor for the conductometric detection of these compounds by metal oxide sensors [55,57,136].

The use of acid–base bonds, between the analyzed molecules and the nanostructured metal oxide layer, also underlies the selectivity in the detection of compounds related to bases, including amines of various structures [137]. Centers for the selective adsorption of compounds with acidic and basic properties can appear in composites as a result of the mutual charging of molecules during the transfer of electrons between metal ions with different work functions [138,139]. The data obtained indicate a correlation between the acid–base bonds of the detected molecules and the sensitive layer, as well as the magnitude of the sensor effect.

The composite sensitive layers of sensors are considered to contain particles that provide preferential adsorption of molecules of a certain type, with semiconductor particles that form conduction paths. The increase in conductivity in this case, as noted above, is due to the spillover of adsorbed molecules onto the conducting semiconductor layer, where they react with oxygen anionic centers, which gives a selective sensor response.

Further application of this approach to the formation of selective conductometric sensors, by combining adsorbing and conductive components in the sensitive layer, was carried out in [140], where modifiers adsorbing analyzed compounds were grafted onto conductive SnO₂ nanoparticles. Long-chain acid molecules with functional groups were chosen as such modifiers [140]. This method was used to obtain sensors that make it possible to distinguish hydrocarbons, in particular, methane, benzene, and toluene, by the magnitude of the sensor response.

Many harmful anthropogenic substances are present in the atmosphere that belong to various π -electronic compounds, in particular, benzene derivatives. The greatest interest, in the high selectivity detection of these compounds, is on nanostructured metal oxide layers containing transition metal inclusions with partially filled d-electron orbitals.

Adsorption bonds between π -electron molecules and nanoparticles of noble transition metals, in particular, Au, Pd, and Pt, are mainly due to the overlap of the π -electron orbitals of the molecules with the orbitals of the d-electron band of the metal [90]. These bonds depend on the position of the π -electron levels, relative to the center of the d-electron band. It is precisely the formation of these bonds with the subsequent spillover of adsorbed molecules, onto conducting metal oxide nanofibers, that ensures the sensitization of the sensor effect in the composite of nanofibers with noble metal nanoparticles [91].

The dependence of the sensor effect on the energy level of the d-electron band of a metal makes it possible to achieve a high selectivity of the sensor response, by using metal particles with different levels and widths of this band, and also considering the levels of π -electron orbitals of the detected molecules [91,141]. Indeed, doping SnO₂ with Pd, Pt, or Au nanoparticles makes it possible to selectively determine the concentration in air of even sub-ppm amounts of C₆H₆, C₇H₈, or CO, respectively.

The selectivity observed in these systems is due to the chemical sensitization of the sensor process due to different interactions between the catalytically active metals and the analyzed gas. Thus, the sensor response of Pd-SnO₂ to 1 ppm C₆H₆ at 300 °C is 25.5, which is almost an order of magnitude higher than the response to C₇H₈, C₂H₅OH, CO, and H₂S. On the other hand, SnO₂ composites with Pt or Au have the maximum response to 1 ppm CO (21.1) and C₇H₈ (40), which also significantly exceeds the response to other gases investigated.

It is important to note that the electronic structure of the surface layer of metal nanoparticles, and the parameters of the d-electron band of this layer, can change due to ligands bound to nanoparticles and deformations of nanoparticles [142]. This opens new possibilities for obtaining selective sensors using noble metal nanoparticles.

4.2. Selective Activity of Adsorbed Molecules in Reaction with Oxygen Anions

In composite metal oxide layers containing clusters of noble metals, the adsorption of detected molecules, especially molecules with π -electron bonds, occurs mainly on these clusters, which have an increased affinity to such molecules. The next stages are the spillover of adsorbed molecules onto the conducting layer of semiconductor metal oxide particles, and the reaction of these molecules with chemisorbed oxygen atoms or molecules that “captured” electrons from the conduction band of the semiconductor. This reaction is accompanied by the return of the “liberated” electrons to the conduction band and, consequently, an increase in the conductivity of the sensor.

However, such a scheme is doubtful at a high bonding energy, between an adsorbed molecule and a metal cluster. For example, the adsorption energy of benzene on a platinum cluster is over 30 kcal/mol and, at the same time, this system has a particularly high sensor response and an extremely short response time [91].

It should be noted that complexes of noble metal clusters, with metal oxide semiconductor nanoparticles in air, already contain chemisorbed oxygen, and this can drastically change the pattern of the sensor process. Most likely, the detected molecule is adsorbed on the complex of the catalytically active metal cluster, with the metal oxide and the negatively charged oxygen center of the sensor reaction, for example, O^- .

Using the example of a CO molecule adsorbed on such a complex, it has been shown that the adsorption of molecules “triggers” the reaction of CO oxidation and the rearrangement of the initial complex [143]. The reaction, $CO + O^- \rightarrow CO_2 + e^-$, produces a sensor response, i.e., leads to an increase in conductivity. Thus, in contrast to the two-stage process considered above, in this case, both adsorption and reaction merge into one reaction.

A sensor process of this type is observed at the interaction of reducing compounds with oxygen centers of nanocomposites consisting of *n*- and *p*-nanoparticles. Sensor characteristics of the *n-p*-nanocomposite depend on the interaction between the particles of its components. In *p-n*-composite semiconductor systems, in which potential barriers at the boundaries of *n*- and *p*-composite nanoparticles prevent the recombination of electrons and holes between particles, the sensor effects depend on the electronic and hole conductivity of the composite. The resulting current flowing through the composite is a combination of the electron current in the conduction band of the composite particles (electron current), and the current, due to electron hopping between positively charged holes in the valence band, in the direction of the electric field (hole current). This situation arises when conducting paths consisting of *n*- and *p*-nanoparticles [23] are simultaneously present in the composite.

In the *p-n*-nanocomposite, when reducing compounds react with oxygen centers on the surface of nanoparticles that have captured conduction electrons, the concentration of these electrons in the particles increases. This leads to an increase in the electronic conductivity (ΔI_e) in the paths consisting of the *n*-type nanoparticles. In *p*-type particles, due to the recombination of positively charged holes with electrons released during the reaction on the surface of these particles, the hole concentration decreases and the hole conductivity (ΔI_h) decreases. As a result, the increase in overall conductivity of a nanostructured *n-p*-composite sensor, upon reaction with a reducing gas (i.e., sensor response), should be less than in a nanostructured sensor consisting only of *n*-nanoparticles. This is confirmed by the data of some studies [24,144].

The sensor response of a nanocomposite *p-n* sensor to reducing compounds depends, firstly, on the adsorption of the detected molecules on the nanoparticles of the sensor's sensitive layer. The difference in the adsorption of detected gases on *n*- and *p*-nanoparticles of this layer significantly increases the selectivity of gas detection.

In particular, the use of a nanostructured *n-SnO₂-p-Co₃O₄* composite sensor made it possible to achieve exceptionally high selectivity in the detection of hydrogen in a mixture with carbon monoxide [24]. This became possible due to different adsorption of H_2 and CO on the *n*- and *p*-nanoparticles of the composite. H_2 is adsorbed and reacts with adsorbed oxygen mainly on the *n-SnO₂* nanoparticles, while CO is equally adsorbed on the *n-SnO₂* and *p-Co₃O₄* nanoparticles [24]. This means that the decrease in the conductivity considered above, and the decrease in the sensor response of the *n-SnO₂-p-Co₃O₄* composite with an increase in the content of the *p*-hole component (*p-Co₃O₄*) in it, will be greater for the detection of CO than the detection of H_2 . Accordingly, the difference between sensor responses to H_2 and CO will increase.

At a certain concentration of Co_3O_4 , near the percolation transition from the electronic conductivity of the composite to the hole conductivity (in the system [24], at a Co_3O_4 content of about 15%), the fraction of the conductivity of the “hole” component of Co_3O_4 , in the total conductivity of the composite, becomes so small that the sensor response of the composite almost does not differ from the response to hydrogen for SnO_2 [24].

High selectivity in the detection of H_2 and CO, in a mixture of these gases, was also found for nanocomposite sensors, *n-SnO₂-p-Mn₃O₄* [144] and *n-ZnO-p-NiO* [145], where the selectivity has the same characteristics as in composites considered in [24]. It was also shown that the annealing temperature of the intermediate mixture of oxides, during the

formation of composites, affects the temperature of electron-hole transitions and sensor responses to CO and H₂ [145]. The observed influence of the annealing temperature on the response opens up new possibilities for controlling the sensor process.

The above results have shown that the doping of sensors based on n-type metal oxides with catalytically active additives leads to the production of sensor materials with high selectivity in the detection of various gases (see Table 3). The influence of the sensor composition on selectivity is due to the electronic and chemical sensitization of the sensor process that occurs during the interaction of the sensitive layer with the analyzed gas.

Table 3. Selectivity of various sensor systems in the detection of various chemical compounds.

Material	Synthesis	Gas Selectivity	Temperature	References
ZnO/NiO	ZnO nanorods by the one-pot chemical method, NiO by hydrothermal method	H ₂ > CO, NO ₂ , CO ₂ , CH ₄	237 °C	[145]
Pd-ZnO/NiO	Pd by chemical reduction method, ZnO nanorods by the one-pot chemical method, NiO by hydrothermal method	H ₂ > CO, NO ₂ , CO ₂ , CH ₄	225 °C	[145]
SnO ₂ -Mn ₃ O ₄	sol-gel technique	H ₂ > CO	350 °C	[144]
SnO ₂ /Co ₃ O ₄	sol-gel technique	H ₂ > CO	350 °C	[24]
SnO ₂ -ZnO	precipitation method	C ₂ H ₅ OH > CO	300 °C	[139]
SnO ₂	precipitation method	CO > C ₂ H ₅ OH	250–300 °C	[139]
SnO ₂ -ZnO	hydrothermal method	Trimethylamine > other gases	330 °C	[138]
Nb-doped TiO ₂ nanotubes	anodic oxidation method	Dimethylamine > NH ₃ , C ₂ H ₅ OH, CO	300 °C	[137]
5% Ca-In ₂ O ₃	electrospun method	Formaldehyde > other volatile organic compounds	130 °C	[136]
5Y-In ₂ O ₃	electrospun method	formaldehyde > other gases	100–120 °C	[135]
NiO/ZnO	one-step hydrothermal method	formaldehyde > other volatile organic compounds	200 °C	[134]

5. Conclusions

The influence of the interaction of nanoparticles on the sensitivity and selectivity of semiconductor sensors in the detection of reducing gases is considered and analyzed. The primary focus is on various detection mechanisms in two-component metal oxide systems. The mechanisms discussed include the influence of doping on the lattice structure, the adsorption of detected gases, the reactivity of oxygen centers, and various possibilities for sensitization of the sensor process. The role of conduction pathways is also considered.

Particular attention is paid to a direction that has been intensively developing recently—sensors based on core-shell nanofibers. The reasons for the high sensitivity and selectivity of such materials are discussed in sufficient detail. In addition, the possibilities of a theoretical description of the electronic structure of nanoparticles and the sensor process in one-component and two-component systems are briefly considered.

One of the important issues considered in this review is the problem of selectivity. Among the various ways to achieve greater selectivity, this review is limited to the consideration of such mechanisms as the selectivity of the process of adsorption of molecules on the sensitive layer surface, and the selectivity of the reaction of oxygen anions with adsorbed molecules.

It is interesting that different authors note both an increase and a decrease in the sensor effect during doping. The sensitivity of sensors is known to be determined by the ratio of the density of conduction electrons in the surface layers of nanoparticles,

in the presence and absence of the detected gas ($n_s(P)/n_s(0)$, where P is the pressure of the detected gas). Sensitivity can be increased either by increasing the numerator or decreasing the denominator. Obviously, at high concentrations of the detected gas, it is better to use nanoparticles with large n_s to enhance the numerator, while at low concentrations, n_s should be small, in order to affect the denominator. In other words, for sensors designed to determine the concentrations of toxic and explosive substances close to the MPC (maximum permissible concentrations), it is preferable to use semiconductors with a high concentration of conduction electrons.

The efficiency of the sensor is determined by the following characteristics: sensitivity, sensor response time and relaxation, optimal operating temperature and selectivity. The values achieved for the first three characteristics can be considered quite satisfactory, especially considering the results of operando mode. The situation is more complicated for temperature and selectivity.

Almost all conductometric gas sensors operate at high temperatures (300–500 °C).

Such high temperatures limit the scope of the application of gas sensors, including the impossibility of their integration into mobile devices, as well as the significant power consumption requirement and risk of explosion. Some reduction in operating temperature has been achieved by using mixed nanocomposites, such as the sensitive layer. In the last decade, studies on the photoactivation of detection at room temperatures have been reported, but the efficiency and response/recovery time of such sensors leave much to be desired. The same can be said about the metal-containing sensitive layers synthesized by low-temperature, solid-phase polymerization.

Although the situation with the selectivity of sensors remains unsatisfactory, new opportunities have been opened for solving this problem. The use of composites based on noble metal nanoparticles, for example, is promising in this regard. Moreover, the sensor effect of nanocrystalline metal oxides containing alkali and alkaline-earth metal ions varies depending on the acidity of adsorbed oxygen-containing compounds. This is the main factor for the selective detection of such compounds. Nevertheless, further research is needed to better address this issue.

Another area that needs to be developed is the theoretical study of the structure of semiconductor nanoparticles and the mechanisms of the sensor effect. This is especially important for two-component sensitive layers, because single-component systems have already been widely explored.

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