In this work, organic–inorganic hybrids based on nanocrystalline indium oxide and ruthenium (II) heteroleptic complexes were used as sensitive materials for room temperature light-activated NO\textsubscript{2} detection. In\textsubscript{2}O\textsubscript{3} was obtained by chemical precipitation method and then annealed at three different temperatures (T = 300, 500, 700 \degree C) in order to investigate the influence of the microstructure of indium oxide on sensor characteristics of hybrid materials and on kinetics of the rise and fall of photoconductivity. The results of the X-ray phase analysis demonstrated that the obtained materials are single-phase with a cubic bixbyite structure. The Ru (II) heteroleptic complex, which was used as a photosensitizer, made it possible to shift the optical sensitivity range of the hybrids to the low energy region of the spectrum and to use a low-power LED (\(\lambda_{\text{max}} = 470\) nm) source for the photoactivation process. The sensor properties were investigated toward NO\textsubscript{2} at sub-ppm range at room temperature. It was found that for pure oxides, the sensor signal correlates with a specific surface area, while for hybrid materials, both the sensor signal and photoresponse increase with increasing the matrix crystallinity. In this case, the main role is played by traps of nonequilibrium charge carriers, which are structural defects in the matrix.

Keywords: nanocrystalline indium oxide; Ru(II) heteroleptic complex; organic–inorganic hybrid materials; room temperature gas sensor; light activation; nitrogen dioxide; microstructure effect

1. Introduction

Since the early 2000s, organic–inorganic hybrid materials have become a rapidly developing investigation objects and attract special attention within the framework of fundamental research and industrial application in the field of gas sensors [1–4]. This is due to the fact that such systems can have new or improved physicochemical functional properties through a synergistic interaction between the constituent components in the interface. In this regard, organic dyes and QD (quantum dots) as photosensitizers are of particular interest as the receptor part of the system, which allow shifting the optical sensitivity of wide-gap semiconductor oxides towards lower energies due to their high extinction coefficient in the visible region of the spectrum [5–10]. The location of HOMO and LUMO levels allows one to control the generation of photoexcited charge carriers and their transfer to the semiconductor conduction band [11–16], while the composition (central cation and ligands) and structure (chirality and steric effects) of the complex itself can contribute to additional and specific interaction with gas phase molecules [17–19].
However, there are pitfalls in the form of physical and chemical parameters of materials, such as composition and microstructure that can either limit or facilitate their use as gas sensors [20–24].

Previously, the authors of the works [25,26] already carried out a study of the influence of the In$_2$O$_3$ microstructure on the NO$_2$ detection. It was concluded that the dependence of the sensor signal on the size of nanocrystals is nonmonotonical; it is connected with the concentration and mobility of the electrons and was explained on the basis of the potential barriers on the boundary of nanocrystals. However, the influence of the matrix microstructure in hybrid systems, where the photosensitizer provides the transfer of charge carriers, still remains unexplored.

The possibility of using the organic–inorganic hybrid materials based on nanocrystalline SnO$_2$ or In$_2$O$_3$ and Ru(II) heteroleptic complexes for the efficient photoactivated detection of nitrogen oxides (NO$_2$ and NO) at room temperature was published in our earlier works [8,9,27]. A recent study of a series of Ru(II) complexes based on 1H-imidazo[4,5-f][1,10] phenanthroline derivatives (ImPh) demonstrated that the complex with the bithiophene ligand demonstrates the most effective photosensitization effect [16]. At the same time, the NO and NO$_2$ gas sensor properties of hybrids based on In$_2$O$_3$ turned out to be two orders of magnitude higher than that of SnO$_2$, which may be due to a smaller band gap and higher number of nonequilibrium charge carriers of the first one [8,9].

Based on the background experience, we have focused our attention in this work on the study of the influence of the In$_2$O$_3$ microstructure on the NO$_2$ detection in sub-ppm range at room temperature by hybrid materials with the Ru(II) heteroleptic complex as the photosensitizer. Particular consideration is paid to the photoelectrical properties of the obtained materials.

2. Materials and Methods

2.1. Materials Synthesis

2.1.1. Synthesis of Nanocrystalline In$_2$O$_3$

The synthesis of nanocrystalline In$_2$O$_3$ was carried out by deposition of an aqueous solution of indium (III) chloride with ammonia solution followed by heat treatment. A solution of NH$_3$·H$_2$O (1 M) was added dropwise into a water solution of InCl$_3$·4H$_2$O (3.5 M) with vigorous stirring until pH ~ 6 was reached. The synthesis was carried out at room temperature. The following reaction took place:

$$2\text{InCl}_3 + 6\text{NH}_3\cdot\text{H}_2\text{O} + (x - 3)\text{H}_2\text{O} \rightarrow \text{In}_2\text{O}_3\cdot x\text{H}_2\text{O} \downarrow + 6\text{NH}_4\text{Cl}$$  (1)

The formed dense gel-like precipitate was separated by centrifugation, then it was washed with deionized water, and with 0.01 M NH$_4$NO$_3$ solution after the beginning of peptization until the chloride ions were removed. An opalescence absence was verified by adding a AgNO$_3$ solution (0.01 M). Next, the In$_2$O$_3$·xH$_2$O gel was dried at 50 °C for 24 h. The resulting glassy mass was grounded to a powder state in an agate mortar and annealed in air at three different temperatures (T = 300, 500 and 700 °C) for 24 h.

2.1.2. The Heteroleptic Ru(II) Complex as Photosensitizer

A detailed description of the synthesis method of heteroleptic Ru(II) complex is given in Supplementary Materials. Table 1 shows the structural formula, the energies of HOMO and LUMO and the label of the complex in this work.

2.1.3. Synthesis of Hybrid Materials

Organic–inorganic hybrid materials were obtained both in the form of powders for the composition and optical characterization and in the form of thick films to study the photoelectric and sensor properties. The sensitization of indium oxide in the form of a powder was carried out by the impregnation method; thick films of pure oxides were formed on microelectronic hotplates and then the solution of the heteroleptic Ru(II) complex
was directly deposited dropwise on their surface. The concentration of the solution was chosen so that the total content of [Ru] relative to [In] was 0.5 wt%. The microelectronic hotplate consists of a metal base with leads, in the center of which a dielectric substrate made of Al₂O₃ (0.9 × 0.9 × 0.25 mm) is fixed on with platinum wires (Figure 1). The substrate is covered with Pt electrodes (0.18 mm in width) on the top side for resistance measurements and a Pt heater (0.18 mm in width) on the back side.

Table 1. Structural formula, the energies of HOMO and LUMO and the label of the Ru(II) heteroleptic complex.

| Structure | E₉HOMO, eV | E₉LUMO, eV | Label | Name |
|-----------|------------|------------|-------|------|
| ![Structural formula](image) | −6.15 | −3.46 | RuTT | bis(2,2′-bipyridine-k²N¹,N¹’)[2-(2,2′-bithiophen-5-yl)-1H-imidazo[4,5-f][1,10]phenanthroline-k²N⁷,N⁸) ruthenium (II) dichloride |

Figure 1. Microelectronic hotplate (a) before and (b) after deposition of a thick semiconductor film on the substrate and (c) after sensitization with a Ru(II) heteroleptic complex; SEM images of the sensitive layer deposited on the substrate, (d) side view and (e) top view.

Figure 1d,e shows the side view and top view SEM images of the sensitive layer deposited on the substrate, respectively. As the sensitive layer is deposited from a suspension with α-terpineol, the surface dries unevenly; this explains why the surface has a wavy shape with an average thickness of 110 ± 10 μm. The surface of the thick film after annealing in the flow of dry air at T = 300 °C during 3 h is porous and consists of agglomerated and sintered grains. The size of agglomerates is about 50–80 nm.
2.2. Materials Characterization

The phase composition of the obtained In$_2$O$_3$ was investigated by X-ray diffraction on a DRON-4-07 diffractometer (radiation source—CuKα with λ = 1.5418 Å). The phase composition was determined using the STOE WinXPOW software. The ICDD PDF database was used to identify the peaks. The crystallite size ($d_{XRD}$) was calculated using the Scherrer equation. The crystal structure of the synthesized samples was also studied by Raman spectroscopy. The studies were carried out on an i-Raman Plus spectrometer (BW Tek, Newark, Delaware, USA) equipped with a BAC 151C microscope. Raman spectra were recorded in the range of 90–2000 cm$^{-1}$ with 4 cm$^{-1}$ resolution; a laser with a wavelength of 532 nm was used as a radiation source. The thickness and morphology of the films were studied by scanning electron microscope (SEM) using a Carl Zeiss SUPRA 40 FE-SEM instrument (Carl Zeiss AG, Oberkochen, Germany) with Inlens SE detector. The specific surface area ($S_{surf}$) of nanocrystalline oxides was measured by low-temperature nitrogen adsorption using a Chemisorb 2750 instrument (Micromeritics). The surface area available for adsorption was calculated using the BET model (Brunauer, Emmett, Teller). Oxidation centers on the surface of the synthesized materials were investigated by the thermo-programmed hydrogen reduction (TPR-H$_2$) method, and also on a Chemisorb 2750 instrument (Micromeritics, Norcross, GA, USA) in a quartz reactor using a 10% H$_2$/Ar gas mixture (50 mL/min) under heating up to 900 °C with 10 K/min rate.

Fourier Transform Infrared (FTIR) spectroscopy measurements were performed on a Spectrum One (Perkin Elmer Inc., Beaconsfield, UK) spectrometer in the transmission mode in the wavenumber range of 400–4000 cm$^{-1}$ with a step of 1 cm$^{-1}$. Samples (about 5 mg) were ground with 50 mg of potassium bromide (Aldrich, for FTIR analysis) and pressed into tablets 0.5 mm thick and 6 mm in diameter.

X-ray fluorescence analysis (XRF) was used to determine the elemental composition of hybrid materials. The study was performed on a M1 Mistral spectrometer (Bruker) with a tube voltage of 50 kV. The diameter of the analyzed region was 1.5 mm; the signal accumulation time was 1 min. The absorption spectra of semiconductor oxides and hybrid materials were obtained on a Perkin-Elmer Lambda-950 spectrophotometer in the diffuse reflection mode in the range of 200–800 nm with a step of 1 nm. The measurements of spectral dependence of the photoconductivity were studied in the range of 350–700 nm and were performed by using a halogen lamp as a white light source and MDR-206 monochromator to obtain irradiation with narrow spectral band. The illumination time for each wavelength was 1 min, and then followed by a stage of the material’s conductivity recovery to the initial stationary value at dark conditions. The conductivity of the samples was measured by a Keithley 6517A electrometer. The following equation was used to calculate the photoconductivity:

$$\frac{\Delta \sigma}{\sigma_0} = \frac{\sigma(\lambda) - \sigma_0}{\sigma_0}$$

where $\sigma(\lambda)$—is the conductivity of the thick film after 1 min of illumination with the corresponding wavelength; $\sigma_0$—is the conductivity of the thick film in stationary state in dark conditions.

The gas sensor properties of hybrid materials were investigated by measuring the electrical conductivity of thick films in situ in a flow cell under a controlled gas flow of 100 ± 0.1 mL/min. The gas mixture for measurements was prepared by dilution with dry synthetic air using RRG-12 electronic mass-flow controllers. Sensor measurements were carried out at room temperature under conditions of periodic illumination with a blue light-emitting diode (LED, 8 mW/cm$^2$, $\lambda_{max}$ = 470 nm, OTdiode, Shenzhen, China) located at a distance of 4 cm above the sensors.
3. Results and Discussion

3.1. Characteristics of Nanocrystalline In$_2$O$_3$

Figure 2a shows X-ray diffraction patterns of nanocrystalline In$_2$O$_3$ samples annealed at temperatures of 300, 500 and 700 °C. According to the results of X-ray phase analysis and ICDD PDF-2 database, the obtained indium oxide powders are single-phase corresponding to a cubic bixbyite structure. It was found that increasing the annealing temperature leads to an increase in crystallite size ($d_{XRD}$) and a decrease in specific surface area ($S_{surf}$) (Table 2).

![Figure 2a: X-ray diffraction patterns of nanocrystalline In$_2$O$_3$ samples.](image)

**Table 2.** Phase composition, crystallite size ($d_{XRD}$) and specific surface area ($S_{surf}$) of nanocrystalline In$_2$O$_3$ samples annealed at temperatures of 300, 500 and 700 °C.

| Sample       | Phase Composition  | $d_{XRD}$, nm | $S_{surf}$, m$^2$/g |
|--------------|--------------------|---------------|---------------------|
| In$_2$O$_3$(300) |                    | 13 ± 1        | 72 ± 3              |
| In$_2$O$_3$(500) | In$_2$O$_3$ bixbyite | 24 ± 2        | 6 ± 1               |
| In$_2$O$_3$(700) |                    | 28 ± 2        | 3 ± 1               |

Figure 2b represents the Raman spectra of nanocrystalline In$_2$O$_3$ samples annealed at temperatures of 300, 500 and 700 °C. The main characteristic Raman modes for the body-centered cubic lattice of In$_2$O$_3$ were observed. The peak at 122.5 cm$^{-1}$ is associated with the vibrations of the In-O bond of the [InO$_6$]$_i$ and the peaks at 298.5 cm$^{-1}$, 487.6 and 621.5 cm$^{-1}$ are due to the deformation and stretching vibration of the [InO$_6$] octahedrons, respectively. The stretching vibrations of In-O-In bonds were corresponded at 357.2 cm$^{-1}$ [28–30]. However, the In$_2$O$_3$(300) sample contains a broad band at 441.5 cm$^{-1}$, which corresponds to a superposition of surface modes. These vibration modes are characteristic of nanosized powders and indicate a highly defective surface layer [31]. It should also be noted that with an increase in the annealing temperature, the background of the Raman spectra descends and becomes horizontal. The defect structure leads to luminescence of the sample under the influence of a powerful radiation source, which leads to the rise in the baseline at higher values of the Raman shift. This statement confirms the defect structure of the surface of the samples annealed at lower temperatures.

The curves of the TPR-H$_2$ method are represented in Figure 3a. Two regions of hydrogen consumption can be distinguished—low-temperature (up to 400 °C) and high-temperature (above 400 °C). The absorption of hydrogen at low temperatures is due to the reduction of surface OH groups and various forms of chemisorbed oxygen:

$$\text{OH}_{\text{ads}} + \frac{1}{2} \text{H}_2 = \text{H}_2\text{O}$$  \hspace{1cm} (3)
In the high-temperature region, the \( \text{H}_2 \) consumption is due to the reduction of \( \text{In}_2\text{O}_3 \) to metallic indium:

\[
\text{In}_2\text{O}_3 + 3\text{H}_2 \rightarrow 2\text{In} + 3\text{H}_2\text{O}
\]  

(5)

The amount of consumed hydrogen was calculated from the peak area, and the calibration was performed using a standard \( \text{Ag}_2\text{O} \) sample. The calculation results of the \( \text{H}_2 \) consumption for two temperature ranges are presented in Table 3. In the low-temperature region, the amount of absorbed hydrogen is 2 times higher in the case of the most dispersed sample (\( \text{In}_2\text{O}_3(300) \)), which indicates a greater number of surface OH groups and chemisorbed \( \text{O}_2 \). A shift in the maximum \( \text{H}_2 \) consumption peak and an increase in its amount with an increase in the annealing temperature of the sample can be observed in the high-temperature region. This is associated with an increase in the particle size and crystallinity degree of \( \text{In}_2\text{O}_3 \), which increases the crystal lattice energy, and therefore its destruction during bulk reduction requires large amounts of thermal energy. The total amount of hydrogen consumed during the reduction of \( \text{In}_2\text{O}_3 \) ranges from \( n = 3.0 \) to \( 3.3 \) mol of \( \text{H}_2 \) per mol of \( \text{In}_2\text{O}_3 \), which is in good agreement with the theoretical value (\( n = 3 \) mol) of the reduction of indium oxide to metal (reaction (5)).

**Table 3.** The results of the TPR-\( \text{H}_2 \) experiments.

| Sample          | \( \text{H}_2 \) Consumption, mol \( \text{H}_2 \) per 1 mol \( \text{In}_2\text{O}_3 \) |
|-----------------|--------------------------------------------------------------------------------------------------|
|                 | Total | 25–400 °C | 400–900 °C | \( T_{\text{max}}, ^\circ \text{C} \) |
| \( \text{In}_2\text{O}_3(300) \) | 3.0 ± 0.1 | 0.12 ± 0.01 | 2.88 ± 0.1 | 582 |
| \( \text{In}_2\text{O}_3(500) \) | 3.2 ± 0.1 | 0.05 ± 0.01 | 3.15 ± 0.1 | 589 |
| \( \text{In}_2\text{O}_3(700) \) | 3.3 ± 0.1 | 0.06 ± 0.01 | 3.24 ± 0.1 | 589 |

FTIR spectra of the samples are shown in Figure 3b. The spectra contain peaks corresponding to the vibrations of bonds in the crystal lattice of In-O (400–600 cm\(^{-1}\)), adsorbed water (1628 cm\(^{-1}\)) and hydroxyl groups (3000–3670 cm\(^{-1}\)). With an increase in the annealing temperature, a decrease in OH groups on the surface is observed, while the...
In$_2$O$_3$(300) sample contains the largest number of OH groups, which is in good agreement with the TPR-H$_2$ results.

The electrophysical properties of the In$_2$O$_3$(300), In$_2$O$_3$(500) and In$_2$O$_3$(700) samples were investigated in the temperature range of 300–25 °C in a dry air flow. The temperature dependence of the conductivity has a complicated form for all samples, which may be associated with the chemisorption of oxygen from air. Therefore, in order to study the electrophysical properties, the temperature dependence of the conductivity was plotted in the Mott coordinates [32] (6):

$$G = \frac{G_M}{T^{0.5}} \exp \left[ -\left( \frac{T_M}{T} \right)^{0.25} \right]$$

where $G_M$ and $T_M$ are the Mott characteristic parameters.

This dependence is well linearized in the temperature range of 140–25 °C. The value of $T_M$ was calculated from the slope of the straight line ln(GT$^{0.5}$) = f(T$^{-0.25}$) (Figure 4). Then, using equations (7–9), the following characteristic parameters were calculated: $N(E_F)$, $R_{hop}$ and $W_{hop}$, where $N(E_F)$ is the density of localized states near the Fermi level, $R_{hop}$ is the electron “jump” length, $W_{hop}$ is the “jump” energy electron, $\alpha$ is a quantity that describes the spatial degree of localization of the wave function, and is equal to 1.24 nm$^{-1}$ and $k_B$ is the Boltzmann constant.

$$T_M = \frac{16\alpha^3}{k_B N(E_F)}$$

$$R_{hop} = \left( \frac{9}{8\pi\alpha k_B TN(E_F)} \right)^{0.25}$$

$$W_{hop} = \frac{3}{4\pi R_{hop}^3 N(E_F)}$$

![Figure 4. Temperature dependence of conductivity of the In$_2$O$_3$(300), In$_2$O$_3$(500) and In$_2$O$_3$(700) samples in Mott coordinates in the temperature range of 140–25 °C.](image-url)

Table 4 shows the parameters characterizing the conductivity of the studied samples according to the Mott model [32]. The Mott model assumes the activation character of conductivity, which indicates the presence of barriers, and the height of the barrier is directly proportional to the resistance of the material. In this case, the resistance value correlates with the grain size in the range of In$_2$O$_3$(300) > In$_2$O$_3$(500) > In$_2$O$_3$(700). The calculated values of the electron “jump” energy or activation energy ($W_{hop}$) for the temperature range of 25–140 °C also changes monotonically, and may be associated with an increase in the
crystallite size, which leads both to a decrease in the height of intergrain barriers and to a shift of the Fermi level to the edge of the conduction band.

Table 4. Parameters characterizing the conductivity of the studied samples according to the Mott model for In$_2$O$_3$(300), In$_2$O$_3$(500) and In$_2$O$_3$(700) samples.

| Sample          | $T_M \times 10^9$, K | $N(E_F) \times 10^{17}$, eV$^{-1}$ cm$^{-3}$ | $R_{hop}$, nm | $W_{hop}$, eV |
|-----------------|----------------------|-------------------------------------------|---------------|--------------|
| In$_2$O$_3$(300)| 14.5                 | 0.24                                      | 26.05         | 0.55         |
| In$_2$O$_3$(500)| 3.5                  | 1.01                                      | 18.26         | 0.38         |
| In$_2$O$_3$(700)| 1.16                 | 3.05                                      | 13.85         | 0.29         |

3.2. Characteristics of Hybrid Materials

The results of a quantitative analysis of the composition of hybrid materials obtained by X-ray fluorescence analysis (XRF) are presented in Table 5. It can be observed from the table that the ruthenium content in the hybrids is close to the theoretically specified value $[\text{Ru}]/([\text{Ru}] + [\text{In}]) = 0.5$ wt.%. These values are averaged and taken from 4 different areas of the sample.

Table 5. Elemental composition of hybrid materials.

| Sample              | $[\text{Ru}]/([\text{Ru}] + [\text{In}])$, wt.% | $[\text{In}]/([\text{Ru}] + [\text{In}])$, wt.% |
|---------------------|-----------------------------------------------|-----------------------------------------------|
| In$_2$O$_3$(300)+RuTT | 0.53 ± 0.01                                    | 99.47 ± 0.13                                  |
| In$_2$O$_3$(500)+RuTT | 0.58 ± 0.01                                    | 99.42 ± 0.11                                  |
| In$_2$O$_3$(700)+RuTT | 0.54 ± 0.01                                    | 99.46 ± 0.12                                  |

The optical absorption spectra (a,b) and normalized spectral dependences of the photoconductivity (c,d) of the sensitized composites and pure In$_2$O$_3$ samples are shown in Figure 5.

![Figure 5](attachment:image.png)

Figure 5. Optical absorption spectra (a,b) and normalized spectral dependences of the photoconductivity (c,d) of pure semiconductors and hybrid materials.

As can be observed from the figures, unmodified In$_2$O$_3$-based samples are transparent in the visible spectral range. All spectra exhibit an absorption band in the UV region, which corresponds to a direct transition from the valence band to the conduction band,
as the energy of this transition exceeds the semiconductor band gap. The curves of the absorption spectra coincide and are overlapped on the curves of the spectral dependence of the photoconductivity for all samples. However, some features can be noticed. In particular, in Figure 5c, a shoulder is observed in the blue region of the spectrum, while its intensity increases with a decrease in the size of indium oxide crystallites. This may be related to excitation of the impurities within the bandgap, because the In$_2$O$_3$(300) sample has a more defective structure.

An absorption edge is observed in the visible region of the spectrum in hybrid materials due to the presence of organic complexes in their composition. Moreover, the maximum of the absorption edge for In$_2$O$_3$(500)+RuTT and In$_2$O$_3$(700)+RuTT samples is shifted to longer wavelengths by about 10 nm; for these composites, a bathochromic shift is observed. This shift may be associated with the stabilization effect of the excited states of Ru(II) heteroleptic complexes by the solid phase of the semiconductor oxide. The increase in photoconductivity for hybrid materials in the visible range of the spectrum indicates the transfer of photoexcited electrons from the Ru(II) heteroleptic complex to the conduction band of the semiconductor oxide.

The photoconductivity of nanocrystalline In$_2$O$_3$ and hybrid materials was studied under constant illumination with blue LED ($\lambda_{\text{max}} = 470$ nm). The curves of changes in the conductivity of the studied samples under illumination are shown in Figure 6. When the light is turned on, the electrical conductivity of materials increases; turning off the LED leads to relaxation of the conductivity. The effect of the annealing temperature of In$_2$O$_3$ on the behavior of photoconductivity under the influence of blue LED radiation is nonmonotonic. The maximum value of the photoresponse, calculated as the ratio of the conductivity under illumination to the conductivity under dark conditions, was achieved by the In$_2$O$_3$(500) sample. In this case, the change in conductivity can be associated with the presence of acceptor levels in the band gap of the semiconductor, as the value of the quantum energy corresponding to the emission of a blue LED is less than the band gap of indium oxide.

As can be observed, the difference in photoresponse between the hybrid material and the corresponding matrix increases with the increasing annealing temperature (Figure 7). The In$_2$O$_3$ annealed at 300 °C has the largest number of structural defects that can act as charge carrier traps. An increase in the annealing temperature leads to ordering of the system and structural perfection, leading to a decrease in the concentration of traps.

![Figure 6](image-url)

**Figure 6.** Change in the conductivity of pure In$_2$O$_3$ with different grain sizes (a) and hybrid materials (b) under dark and irradiation conditions.
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Figure 7. Photoresponse of synthesized samples.

3.3. Investigation of the Gas Sensor Properties of Hybrid Materials in Interaction with Nitrogen Dioxide in Dry Air

The illumination of the sensor was carried out in a pulsed mode with periods of 2 min by switching on and off the LED, as a result of which the change in the resistance of the sensor with time also acquires a periodic character and is an alternation of the curves of the decay and rise of the photoconductivity. The repeated cycle of turning on and off the light leads to the change in the resistance of the sensor in each of the subsequent cycles close to the previous one. In this steady state, the minimum resistance $R_{\text{light}}$ is achieved during the illumination of the sensor and the maximum resistance $R_{\text{dark}}$ is achieved in the dark conditions. The ratio (10) varies with the composition of the atmosphere and has been used as a “sensor signal” for this measurement technique.

$$S = \frac{R_{\text{dark}}(\text{gas})}{R_{\text{dark}}(\text{air})} \quad (10)$$

$$S_{\text{ph}} = \frac{R_{\text{dark}}}{R_{\text{light}}} \quad (11)$$

Figure 8 shows the change in the resistance of sensors based on pure oxides and hybrid materials under periodic blue LED illumination in the presence of various concentrations of NO$_2$ (0.1–0.25–0.5–1.0 ppm). As can be observed, the sensors show a periodic change in resistance under the influence of pulsed illumination, both in air and in the presence of NO$_2$. On the logarithmic scale of resistance, it can be observed that with an increase in the concentration of NO$_2$, the ratio of the photoresponse (11) increases for all sensors sensitized with Ru(II) heteroleptic complex. Moreover, with an increase in the concentration of NO$_2$, the average resistance of the sensors increases. The sensor resistances at the same NO$_2$ concentrations, obtained both with an increase and a decrease in the detected gas in air, are very close, which indicates a good ability of these sensors to operate reversibly.
Table 6 represents the average values of the sensor resistances under dark conditions and constant light illumination in dry air. The resistances were measured during 1 h for each step, in order to obtain a relatively stable state. The first observation is that the resistance of all sensors under the LED illumination is much less, which is due to the photodesorption of oxygen from the surface that was adsorbed in the air flow in dark conditions. It can also be noticed that the higher the annealing temperature of the sample (resulting in larger In₂O₃ grain size), the lower the baseline resistance during measurements. This can be associated with the height of potential barriers at the grain boundary: the smaller the crystallite size, the higher the barrier.

Table 6. Baseline resistance in dark conditions and under LED illumination in dry air at room temperature.

| Sample                  | R (Dark), Ohm | R (Light), Ohm |
|-------------------------|---------------|---------------|
| In₂O₃ (300)             | 7.6 x 10⁳     | 10³           |
| In₂O₃ (500)             | 2.6 x 10³     | 4.6 x 10²     |
| In₂O₃ (700)             | 7.2 x 10²     | 4.0 x 10²     |
| In₂O₃ (300) + RuTT      | 1.6 x 10⁶     | 6.5 x 10⁵     |
| In₂O₃ (500) + RuTT      | 5.3 x 10⁵     | 9.4 x 10⁴     |
| In₂O₃ (700) + RuTT      | 4.2 x 10⁵     | 8.6 x 10⁴     |

In our previous works, it was demonstrated that the energy of the blue LED ($\lambda_{\text{max}} = 470$ nm) excitation is equal to the absorption band of the Ru(II) heteroleptic complex in the visible range of the spectrum [8,9]. During light-activation, the electron-hole pairs are generated in it and then electrons can be transferred to the In₂O₃ conduction band facilitating to increase conductivity. NO₂ being a stronger electron acceptor than O₂, can attract the electrons from the conduction band of the semiconductor:

$$\text{NO}_2(\text{gas}) + e^- \leftrightarrow \text{NO}_2(\text{ads})$$  \hspace{1cm} (12)

$$\text{NO}_2(\text{gas}) + \text{O}_2(\text{ads}) \leftrightarrow \text{NO}_2(\text{ads}) + \text{O}_2(\text{gas})$$  \hspace{1cm} (13)
On the other hand, the chemisorbed particles (both \( \text{NO}_2^{-} \text{(ads)} \) and \( \text{O}_2^{-} \text{(ads)} \)) can be converted into physically sorbed particles by interaction with photogenerated holes:

\[
\text{NO}_2^{-} \text{(ads)} + h^+ \leftrightarrow \text{NO}_2 \text{(gas)}
\]  

(14)

The values of the sensor signal were calculated using formula (10) and the photoresponse—using formula (11). The concentration dependences of the sensor signal and photoresponse of semiconductor oxides and hybrids are shown in Figure 9.

![Figure 9. Dependence of the sensor signal (a) and photoresponse (b) of In\(_2\)O\(_3\)(300), In\(_2\)O\(_3\)(500), In\(_2\)O\(_3\)(700) samples and hybrid materials based on them on the concentration of NO\(_2\).](image)

The obtained data show that the introduction of a photosensitizer into nanocrystalline oxides, In\(_2\)O\(_3\), leads to an increase in sensor signal and photoresponse by 2–3 orders. For pure oxides, the sensor signal correlates with a specific surface area and is inversely with crystallite size. For hybrid materials, both the sensor signal and photoresponse increase with increasing matrix crystallinity. In this case, the main role is played by traps of different air humidity (relative humidity at 25°C). It can also be noticed that the higher the annealing temperature of the sample conditions compared to the photoresponse time. Moreover, the photogenerated electrons accelerated by the kinetics of photoconductivity.

On the one hand, nanocrystalline In\(_2\)O\(_3\) has a significant advantage over the sample with a large crystallite size, because nanosized crystalline grains have a large specific surface area, which leads to a more intense interaction of the semiconductor with the gas phase and, as a result, to an increase in the sensor signal. This was observed for pure oxides in NO\(_2\) detection. However, a consequence of a decrease in nanocrystals is an increase in the height of potential barriers at the grain boundary, which in turn can lead to a decrease in the mobility of free electrons and the probability of their tunneling [25,26]. The consequence of this was an increase in resistance with a decrease in the size of crystallites (Figure 8 and Table 6). The obtained result corresponds to the calculated values of the activation energy \( (W_{\text{hop}}) \) for the samples at room temperature: In\(_2\)O\(_3\)(300) (0.55 eV) > In\(_2\)O\(_3\)(500) (0.38 eV) > In\(_2\)O\(_3\)(700) (0.29 eV). On the other hand, the crystal structure of nanosized materials is more defective, and that has been proven above using various characterization methods. Acting as traps, these defects can reduce the number of nonequilibrium charge carriers and, as a result, increase the time of relaxation and recovery to a steady state of resistance in dark conditions compared to the photoresponse time. Moreover, the photogenerated electrons from the heterocyclic Ru(II) complex can also be trapped by this type of defect.

The humidity effect was studied for the samples based on In\(_2\)O\(_3\) annealed at 300 and 700 °C. Figure 10 shows the dependence of the sensor signal on the NO\(_2\) concentration at different air humidity (relative humidity at 25 °C RH = 25, 45, 65%).
As can be noticed, a monotonous increase in sensor signal is observed with increasing humidity for pure oxides. It was demonstrated that in humid atmosphere the conductivity of the n-type metal oxides increases, as dissociative adsorption of H₂O molecules or replacement of the chemisorbed oxygen ions can lead to release of the additional electrons into the conduction band [21,27]. The increase in the concentration of the electrons can stimulate the adsorption of the NO₂ molecules [6]. Sample In₂O₃(300) has the largest specific surface area and, consequently, more adsorption sites; therefore, this effect is more significant for it compared to sample In₂O₃(700).

The hybrid materials have specific interactions with NO₂ at the humid atmosphere. For the NO₂ concentration below 0.25 ppm, the sensor signal increases with increasing relative humidity because of additional electrons released by the interaction with water vapor. However, for higher NO₂ concentrations, the change in sensor signal in humid atmosphere is negligible comparing with the sensor signal in dry air. In this case, vacant surface active sites play vital roles as, due to their limited amount, the surface is saturated with adsorbed water and the proportion of “water electrons” becomes limited. Moreover, certain sites are already occupied by molecules of the Ru(II) heteroleptic complex in hybrid samples. Therefore, water and nitrogen dioxide molecules can compete for the same remaining adsorption sites. As the In₂O₃(700) sample has the smallest specific surface area, a very limited number of water molecules will be able to adsorb onto the free sites. This is additionally confirmed by the fact that with an increase in relative humidity up to 65%, the sensor signal begins to decrease and, compared to the sample In₂O₃(300), the sensor signal in a humid atmosphere is less.

Table 7 illustrates a comparative analysis of materials used for the detection of NO₂ gas by photoactivation. Using illumination at room temperature instead of heating for activation of physical and chemical processes on the surface of the sensitive layer will significantly reduce the power consumption of sensors in the case of their practical application. An analysis of the literature data demonstrated that UV sources are mainly used for photoactivated gas sensors. The use of low-power LEDs with radiation in the visible region of the spectrum, unlike UV sources, will further reduce energy consumption and allows one to embed such sensors into compact electronic devices. The periodic illumination with a duty light cycle of 50% used in this work will further reduce the power consumption by half. The highest sensor signal is established for CdSe QD@In₂O₃ composite material [5] and here the authors also used a periodic illumination approach. However, the specific
power was higher (20 mW/cm² vs. 8 mW/cm²) and the response differs by one order (10⁶ vs. 1.75 × 10⁵). Hereby, the hybrid materials based on nanocrystalline In₂O₃ and Ru(II) heteroleptic complex have good prospects, also like the CdSe QD@In₂O₃ composites, for further research and applications as low power consumption gas sensors.

Table 7. Comparison of the light-activated (with LED) NO₂ gas sensor performances at room temperature for different materials.

| Material | Wavelength, nm | Incident Irradiance, mW/cm² | NO₂ Concentration, ppm | Sensor Signal (a), % Ref. |
|----------|----------------|-----------------------------|------------------------|--------------------------|
| Mesoporous In₂O₃ | 400 | - | 5.0 | 900 [33] |
| In₂O₃ | 385 | 1.0 | 8.0 | 17,900 [34] |
| WO₃ | 590 | 340 | 0.16 | 820 [35] |
| Au/ZnO | 365 | 1.2 | 5.0 | 455 [36] |
| Al/TiO₂/Al₂O₃/p-Si | 254 | - | 20 | 11.5 [37] |
| ZnS-core/ZnO-shell nanowires | 254 | 1.2 | 1.0 | 339 [38] |
| Bi₂O₃-core/ZnO-shell nanobelt | 254 | 1.2 | 1.0 | 227 [39] |
| N-719 dye/ZnO hybrid | 480 | 370 | 1.25 | 143 [40] |
| CuO/ZnS nanowire | 365 | 2.2 | 5.0 | 955 [41] |
| CdSe QD@In₂O₃ | 535 | 20 | 1.6 | 10⁶ [5] |
| CdSe QD@ZnO | 535 | 20 | 1.6 | 3000 [5,42] |
| ZnO/In₂O₃ composite | 365 | 25 | 5 | 221 [43] |
| Au/MoS₂ | 365 | - | 2.5 | 30 [44] |
| ZnO | 455 | 5 | 0.025 | 20 [45] |
| WS₂-decorated rGO | 430 | 0.66 | 1.0 | 21 [46] |
| ZnO/(ZnSe(shell)@CdS(core)) composite | 535 | 20 | 2.0 | 6900 [6] |
| WO₃ | 365 | 8 | 5.0 | 11,300 [47] |
| Perylenediimide-sensitized SnO₂ | 400–700 | - | 0.5 | 12,900 [48] |
| Polypeptide-assisted ZnO nanorods | 365 | - | 10.0 | 400 [49] |
| Au-ZnO nanorods | 495 | 50 | 1.0 | 109 [50] |
| Ag-ZnO heterostructure | 470 | 75 | 1.0 | 150 [51] |
| MoS₂/ZnO nanohybrid | 365 | 0.3 | 0.5 | 2310 [52] |
| ZnO/CaPbBr₃ NCs | 470 | 8.0 | 2.0 | 30,000 [7] |
| In₂O₃–ZnO nanotubes | 365 | 1.95 | 0.5 | 3170 [53] |
| In₂O₃/Ru(II) heteroleptic complex | 470 | 8.0 | 1.0 | 1.75 × 10⁵ This work |

(a) The sensor signals were recalculated as \( S = \frac{R(\text{NO}_2) - R(\text{air})}{R(\text{air})} \times 100\% \).

4. Conclusions

Synthesized organic–inorganic hybrid materials based on the nanocrystalline In₂O₃ and Ru(II) heteroleptic complex demonstrated high sensitivity in NO₂ (0.1–1.0 ppm) detecting at room temperature under periodic illumination with blue light (\( \lambda_{\text{max}} = 470 \) nm). Surface modification of In₂O₃ with the heteroleptic complex leads to the growth of photosensitivity accompanied by the reproducible change in the photoresponse depending on the NO₂ concentration. For pure oxides, the sensor signal correlates with the specific surface area, while for hybrid materials, both the sensor signal and photoresponse increase with increasing the matrix crystallinity; the In₂O₃(700)+RuTT sample demonstrated the
highest values of response. In this case, the main role is played by traps of nonequilibrium charge carriers, which are structural defects in the matrix. An increase in the annealing temperature decreases their concentration, which makes it possible to accelerate the kinetics of photoconductivity.

**Supplementary Materials:** The following are available online at https://www.mdpi.com/article/10.390/chemosensors10040135/s1; Scheme S1: Synthesis of Ru (II) complex.

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