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Arc-discharge-assembled CNT/MoO₃ nanohybrids for ultra-sensitive and selective sub-ppm ethanol detection at room temperature

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In this study, one-dimensional carbon nanotube (CNT) and zero-dimensional MoO₃ nanohybrids were synthesized using a simple arc-discharge method for ethanol gas sensor applications. MoO₃ nanoparticles were uniformly distributed on the surface of mesoporous CNTs, which increased the specific surface area and the availability of active sites for charge carriers within the nanohybrid. MoO₃ functions as the receptor, while the CNTs serve as the transducer, leading to the modification in the depletion region at the hybrid surface, followed by enhancement of the sensing performance. The CNT/MoO₃ sensor exhibited the highest response of 76.5% to 1 ppm ethanol even at room temperature operation (30 °C), significantly outperforming CNT (12.5%) and MoO₃ (2.5%). Additionally, the CNT/MoO₃ sensor revealed rapid response and recovery time, excellent selectivity, and minimal humidity dependence. SEM, TEM, XRD, XPS, and BET analyses confirmed that the improved gas sensitivity of the CNT/MoO₃ nanohybrid is attributed to the increased active sites for charge carriers, abundant surface vacancies, and modification in the depletion region.

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1. Introduction

Volatile organic compounds (VOCs) have long been recognized as significant threats to environmental safety and human health.¹ Among these, ethanol, the most common VOC, poses a serious health risk at room temperature (RT) when prolonged exposure to ambient air exists.^{2–4} Therefore, there is a strong demand for the development of highly sensitive, small-sized, fast-response materials with enhanced selectivity and long-term stability at RT to protect human health.⁵ Metal oxide semiconductors (MOSSs) are widely used as the primary sensing materials due to their excellent stability, low cost, and good

response characteristics.^{6,7} Therefore, studies on MOS-based gas sensors, such as ZnO,^{8–10} In₂O₃,^{11,12} Fe₂O₃,¹³ and MoO₃,^{14,15} have been extensively reported. Among these, MoO₃, a novel n-type semiconductor, offers advantages such as low production cost and high electron transfer efficiency, making it a promising candidate for use in gas sensors, batteries, and other related applications.^{16,17} For instance, Wang *et al.* synthesized MoO₃/SnO₂ core-shell heterostructures that exhibited a high response value toward 100 ppm ethanol at 200 °C (ref. 18) and MoO₃ prepared by Li *et al.* demonstrated a response of 3.86 toward 20 ppm ethanol exposure at 240 °C, with excellent ethanol selectivity.¹⁹ However, the high operating temperature hinders the applicability of ethanol detection because of the significant risk of explosion.^{20,21} Therefore, reducing the operating temperature of MoO₃-based sensors while maintaining good response characteristics, fast response/recovery kinetics, and high selectivity is critical.

In the meantime, carbon nanotubes (CNTs) have been widely explored in the low-temperature sensor application since their discovery by Iijima *et al.* in 1991 (ref. 22) due to their high surface area, large active sites for gas contact, and low operating temperatures.²³ Hybridization of CNTs with MOSSs could introduce synergistic effects,^{24,25} further enhancing the gas sensitivity of the composite material.^{7,26,27} CNTs, which are known to exhibit p-type semiconductor properties,²⁸ can form p-n

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heterojunctions with n-type semiconductors like MoO_3 . Such heterojunctions oftentimes lead to improved gas sensitivity and selectivity while lowering the operating temperature of the sensor.

However, only a few studies about p–n heterojunctions involving CNT/ MoO_3 for ethanol detection at room temperature have been reported. In this study, we successfully fabricated CNT/ MoO_3 hybrid structures with various CNT contents by the arc-discharge method. The hybrid structures were characterized using SEM, TEM, XRD, Raman, XPS, and BET analyses to investigate their morphology, electronic state, specific surface area, surface features, pore volume, and gas adsorption properties. The CNT/ MoO_3 nanohybrid exhibited a higher specific surface area, thereby providing more surface-active sites for ethanol adsorption. The charge carrier transfer mechanism is modified by the formation of a depletion layer in the nanohybrid, leading to high sensing capability. Furthermore, we unveiled the sensing mechanism and the origin of the highest sensing capability in our nanohybrid materials.

2. Experimental section

2.1. Fabrication of CNT/ MoO_3 hybrid sensors

CNT/ MoO_3 hybrid sensors were fabricated on SiO_2/Si substrates *via* a co-arc discharge method. The substrates (dimensions: $5 \text{ mm} \times 0.65 \text{ mm}$, oxide thickness: 300 nm) underwent ultrasonic cleaning in acetone, methanol, and deionized water for 15 minutes each, followed by drying with nitrogen gas. Titanium and platinum layers with 30 nm- and 120 nm-thickness, respectively, were sequentially deposited to the substrate *via* DC magnetron sputtering, employing a shadow mask to form the parallel-electrode configurations. These electrode-patterned substrates were then placed on the inner wall of the arc-discharge chamber, following the procedure detailed in previous studies.^{7,8,26} The active sensing area was defined using Scotch tape as a mask. The arc discharge was conducted at

a current density of 45 A cm^{-2} in a hydrogen (H_2) atmosphere with a partial pressure of $6.0 \times 10^3 \text{ Pa}$. Five molybdenum wires served as the catalyst source were inserted into a pure hollow graphite tube (inner diameter: 3 mm; outer diameter: 6.4 mm; length: 160 mm). The thickness of the CNT/ MoO_3 nanohybrid was adjusted by varying the co-arc discharge duration to 5, 15, or 30 minutes. To enhance adhesion and ensure reliable electrical contact between the CNT/ MoO_3 nanohybrid and the Si/SiO_2 substrate, the as-deposited films were immersed in methanol. The resulting films went through a $400 \text{ }^\circ\text{C}$ heat treatment for 2 hours in dry air to eliminate the remaining amorphous carbon and fully oxidize the molybdenum to MoO_3 .²⁷ The samples with 5, 15, and 30 min were designated as CM5, CM15, and CM30, respectively. A schematic flowchart illustrating the CNT/ MoO_3 hybrid sensor synthesis is presented in Fig. 1a. In addition, chemical treatment with a mixture of nitric and sulfuric acid (1 : 3 ratio) was carried out for the CM15 sample to isolate bare CNTs (b-CNT). CM15 samples were also annealed at $700 \text{ }^\circ\text{C}$ for 3 hours to yield bare MoO_3 (b- MoO_3) by burning off CNTs.

2.2. Morphological and structural characterization

The morphological characteristics of the CNT/ MoO_3 network structures were analyzed using field-emission scanning electron microscopy (FE-SEM; JSM 700F, JEOL, Japan) and high-resolution transmission electron microscopy (HR-TEM; Tecnai G2 F30 S-TWIN, FEI, Netherlands). The X-ray diffraction was conducted using X'pert PRO-MPD (PANalytical, Netherlands) with $\text{Cu K}\alpha$ radiation. To explore the chemical bonding states, we employed the X-ray photoelectron spectroscopy (XPS; Thermo Scientific MultiLab 2000, Pittsburgh, PA, USA) with monochromatic $\text{Al K}\alpha$ radiation and Raman spectroscopy (UniRAM spectrometer equipped with a 532 nm excitation laser and a cooled CCD detector). Additionally, nitrogen adsorption/desorption analysis was conducted to evaluate the surface area

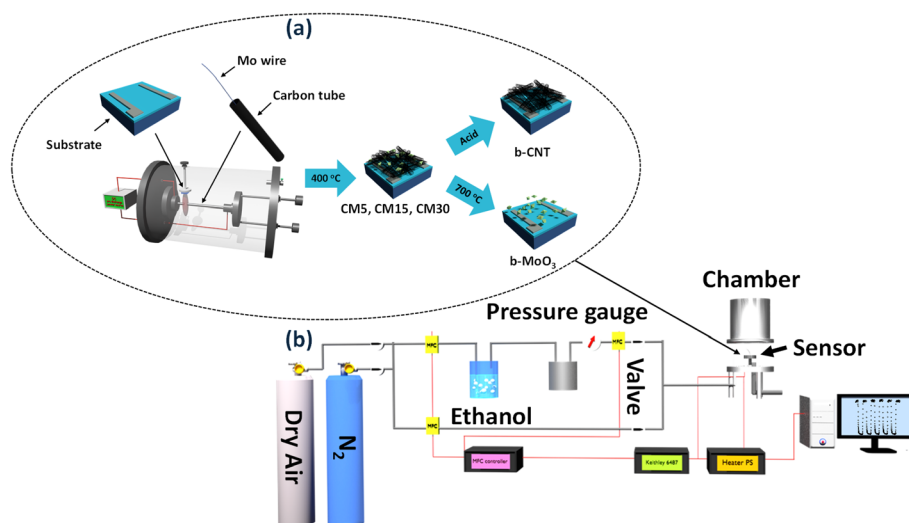


Fig. 1 (a) The fabrication flowchart of the CNT/ MoO_3 composite sensor on the Si/SiO_2 substrate. (b) Experimental setup for gas sensing measurement.

and pore characteristics of the hybrid structure using the Autosorb-1-C analyzer (Quantachrome).

2.3. Sensing measurement

Fig. 1b shows the schematic of the gas detection system using a pico-ammeter/voltage (Keithley 6487). The sensor substrates were placed on the top of the chamber in which the mass flow controllers (MFCs) control the sensor operating temperature and gas transfer rate. By applying a 1 V bias between the sensor electrodes, the corresponding resistance and conductance were recorded. Saturated ethanol vapors were produced using a bubbler system, in which dry air was passed through pure ethanol to generate a controlled vapor stream. The Antoine equation was used to determine the vapor pressure, which is given by: $\log_{10}(p_s) = A - \frac{B}{C + T}$, where p_s is the saturated vapor pressure, T is the temperature of the gas-liquid system²⁹ and A , B , and C are component-specific constants. The relative substance concentration in the gas phase (x_s) was determined by $x_s = P_s/P$, where P is the total pressure of the entire system. The ethanol gas or humidity concentration, C_g and C_H , respectively, was calculated *via* the equation:

C_g (or C_H) = $C_{\text{cyl}} \times \frac{f}{f + F}$, where f and F are the flow rates of

analyte gas (ethanol) and carrier gas, respectively, while C_{cyl} is the analyte gas concentration in the cylinder (or $C_{\text{cyl}} = x_s$ for humidity exposure).³⁰ The sensor response, S (%), is calculated using the equation: $(R_g - R_o)/R_o \times 100$, where R_o and R_g represent the baseline resistance in dry air before ethanol exposure and the resistance measured during ethanol exposure, respectively. For the gas selectivity test, the sensor was exposed toward 3 ppm of H_2S , NH_3 , H_2 , CO , CH_4 , methanol, acetone, and acetic acid gases. To operate the system under the humidity effect, water molecules were similarly pumped through the bubbler with the flow rate ratio controlled by the MFCs. The relative humidity index was determined by the Testo 635 Temperature and Moisture meters.

3. Results and discussion

3.1. Morphology and structure

The co-arc discharge process with a molybdenum-embedded graphite tube facilitates the simultaneous evaporation of molybdenum and graphite, leading to the formation of Mo nanoparticles and the growth of CNTs, which is primarily driven by the catalytic activity of molybdenum. Molybdenum in the graphite tube resulted in the formation of Mo nanoparticles, with CNT bundles interweaving through these particles. Subsequent oxidation of the as-synthesized Mo-embedded CNTs at 400 °C produced the CNT/MoO₃ nanohybrid structure. CNT/MoO₃ nanohybrids with various deposition times (5, 15, and 30 min) were denoted by CM5, CM15, and CM30, respectively, and their SEM images are presented in Fig. 2a, b, and c. The concentration of the CNT/MoO₃ nanohybrid on the substrate increased progressively with deposition time. SEM images confirmed the porous structure of the CNT/MoO₃

nanohybrid network, characterized by CNT bundles interwoven with MoO₃ nanoparticles. A continuous layer covering the entire sensor surface was not achieved even in the CM30 sample. The size of the MoO₃ particles ranges from several nanometers to tens of nanometers regardless of the deposition time. The SEM images of bare CNT (b-CNT) and bare MoO₃ (b-MoO₃) are also presented in Fig. 2d and e, respectively. In Fig. 2d, MoO₃ particles were completely leached out by acid treatment while preserving only CNTs on the substrate. Meanwhile, CNTs were completely combusted at 700 °C, leaving only MoO₃. High-resolution TEM images (Fig. 3a and b) further visualize the nanohybrid morphology, clearly confirming entangled MoO₃ nanoparticles interconnected with the CNT bundle network. TEM analysis also revealed the presence of single-walled CNTs due to the catalytic role of Mo.

The XRD patterns of the CNT/MoO₃ nanohybrids, along with the bare CNT (b-CNT) and bare MoO₃ (b-MoO₃) structures, are presented in Fig. 2f. The characteristic peaks observed at 26.5° and 43° are attributed to the (002) and (100) lattice planes of CNTs, respectively (JCPDS card no. 75-1621). Meanwhile, the peaks at 25.53°, 33.76°, 38.89°, and 55.84° correspond to the (040), (111), (060), and (042) lattice planes of the orthorhombic MoO₃ phase, respectively (JCPDS card no. 05-0508). Other crystalline phases were not detected. Raman spectra of b-CNT, b-MoO₃, and CNT/MoO₃ composite (CM15) between 250 and 2000 cm⁻¹ are presented in Fig. 3c. The quality of the CNTs was analyzed by Raman spectroscopy. The sharp peaks indicate that the vibrational modes predominantly result from a well-ordered, high-degree crystalline structure. The Raman corresponding spectrum of CNTs generally consists of two major peaks at ~1340 cm⁻¹ (D band) and ~1595 cm⁻¹ (G band).³¹ The D-band originated by sp³ electronic states is well-known for defective, glassy carbon, and disordered graphite, while the G-band is related to the two-dimensional hexagonal lattice sp² vibration in the graphite. The typical high G-band/D-band intensity ratio is a signature of the high crystallinity single-wall CNTs.³² The G-band to D-band peak intensity ratios (I_G/I_D) of the CNTs are ~2 in CM15, demonstrating the good quality of the as-synthesized CNTs. These results demonstrate the structural integrity of sp²-hybridized carbon atoms of the single-wall CNTs. The D- and G-band peaks of the b-MoO₃ disappeared due to burning CNTs out completely at 700 °C. Six typical peaks at 283, 334, 375, 665, 819, and 994 cm⁻¹ were observed, which is indicative of the crystal structure of α -MoO₃. The Raman band observed at 283 cm⁻¹ (B_{2g}, B_{3g}) corresponds to the wagging vibrational modes of the terminal oxygen atoms (Mo=O vibration). The band at 334 cm⁻¹ (B_{1g}, A_g) is attributed to the Mo₃-O bending mode, while the band at 375 cm⁻¹ (A_g) is assigned to the bending mode of the Mo=O bond.³³ The MoO₆ octahedral building units contain three nonequivalent oxygen atoms: terminal Mo=O (O₁), twofold-bridged Mo-O-Mo (O₂), and threefold-bridged O-Mo₃ (O₃). The Raman shifts at 994 cm⁻¹ (O₁), 819 cm⁻¹ (O₂), and 665 cm⁻¹ (O₃) are associated with the stretching vibrations of these respective oxygen sites, characteristic of the α -MoO₃ crystal structure.³⁴ In addition, b-CNT did not show any appearance of peaks related to the α -MoO₃, confirming the purity of CNTs after acid treatment.



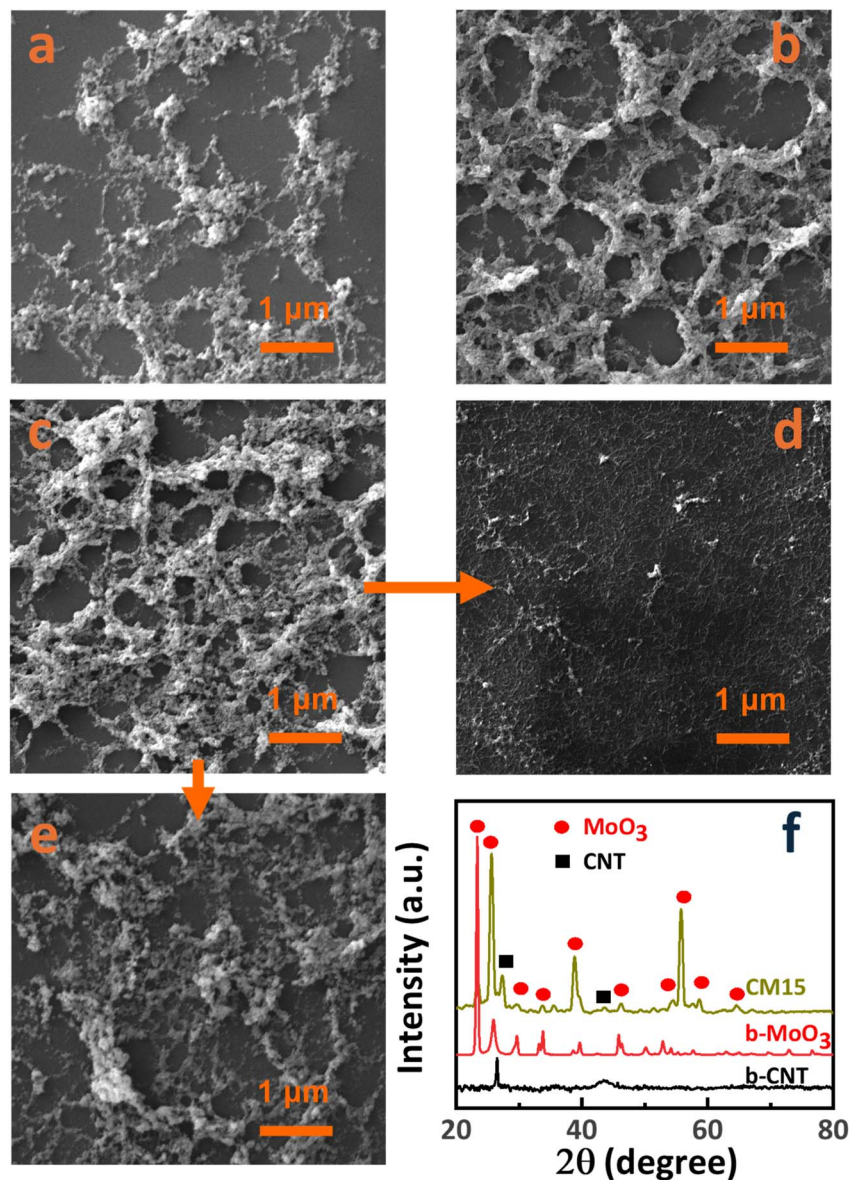


Fig. 2 SEM illustration of CNT/MoO₃ hybrids (a) CM5, (b) CM15, (c) CM30, (d) b-CNT, and (e) b-MoO₃. (f) XRD patterns of b-CNT, b-MoO₃, and CM15.

The XPS spectra of b-CNT, b-MoO₃, and the nanohybrid CM15 are displayed in Fig. 3d, e, and f, respectively. In Fig. 3d, the typical doublet peaks at binding energies of ~232 eV and ~235 eV match with 3d_{5/2} and 3d_{3/2} states in Mo⁶⁺, respectively.³⁵ The core-level spectrum of the O 1s state consists of two chemical states: the lower binding energy region (529.8–530.1 eV) corresponding to bulk oxygen anions within the MoO₃ matrix and the higher binding energy region (530.5–531.0 eV) associated with accumulated oxide impurities. The C 1s spectrum, Fig. 3f, exhibits an asymmetric feature, which can be deconvoluted into two peaks. The dominant peak at ~284.5 eV stems from carbon-carbon interactions, including sp² hybridized C=C bonds, while the relatively weaker peak at ~286.6 eV is attributed to the carbon-oxygen interactions, including C-O bonds.³⁶

3.2. Electrical and gas sensing property

The current-voltage (*I*-*V*) characteristics of b-CNT, b-MoO₃, and CNT/MoO₃ nanohybrid sensors, Fig. 4a, were evaluated at room temperature (30 °C) under a dry air environment. To ensure accurate measurement, the sensing chamber was pre-heated to 300 °C in dry air to remove any residual water molecules from the sensor surfaces, followed by cooling to room temperature. The *I*-*V* curves, plotted in a logarithmic scale, exhibit consistent ohmic behavior across all sensors. The temperature-dependent conductance (*G*) of the sensors was assessed within the range of 30 °C to 300 °C. Conductance values were calculated using Ohm's relation $I = G \times V$, as depicted in Fig. 4b. The conductance of all sensors increased with temperature, corroborating the semiconducting properties of the materials. Among the samples, b-CNT showed the highest conductance or lowest

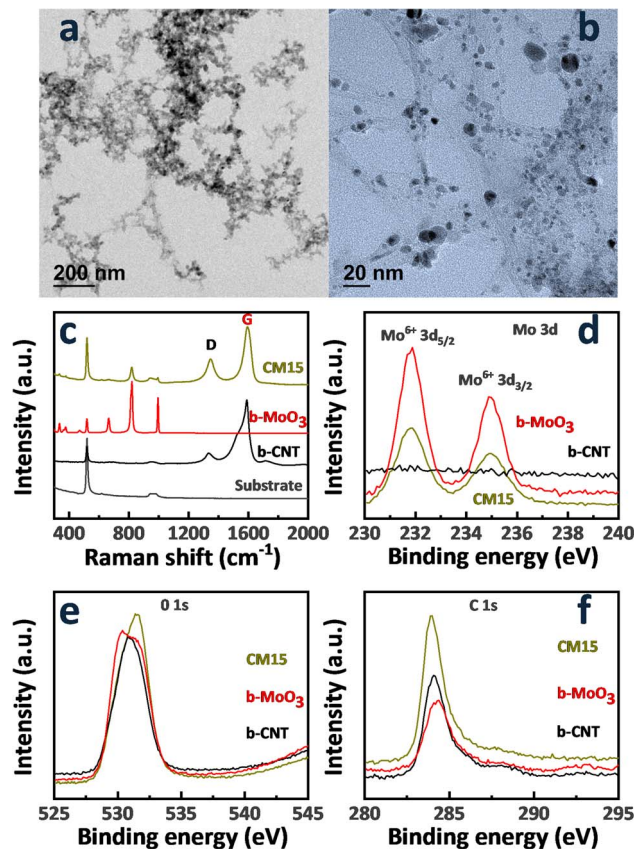


Fig. 3 (a) Low and (b) high-magnification TEM images of the CNT/MoO₃ hybrids (CM15). (c) Raman spectra of b-CNT, b-MoO₃, and CM15. High-resolution XPS spectra of (d) Mo 3d, (e) O 1s, and (f) C 1s states of b-CNT, b-MoO₃, and CM15.

resistance (Fig. 4d), resulting from the intrinsically high electrical conductivity of CNT as a well-established p-type semiconductor.^{37–39} Meanwhile, MoO₃ displayed n-type semiconducting behavior with a wide bandgap energy ranging from 3.11 to 3.22 eV.^{40,41} Therefore, in the CNT/MoO₃ nanohybrid, the conduction mechanism can be explained by a parallel circuit model, where the high-resistance MoO₃ component and the low-resistance CNTs act as parallel conductive pathways. The overall conductance of the hybrid structure was predominantly influenced by the intrinsically high hole-carrier concentration of CNTs. The incorporation of MoO₃ into the CNTs network reduced the overall conductance of the composite by several orders of magnitude. The decreasing conductance was likely attributed to the formation of the depletion region at the junction between MoO₃ particles and CNTs bundles, which effectively acted as an insulating layer and suppressed conductance.^{42,43} The conductance of the nanohybrid increased with greater deposition time, as shown in Fig. 4c, possibly owing to the increased material content. The CM5 sensor, with a thinner CNT/MoO₃ layer, exhibited higher conductance than the bare MoO₃ sensor, further underscoring the critical role of CNTs in dictating the electrical behavior of the composite.

The ethanol sensing behaviors of the sensors were investigated from 30 °C to 300 °C. The response and recovery kinetics

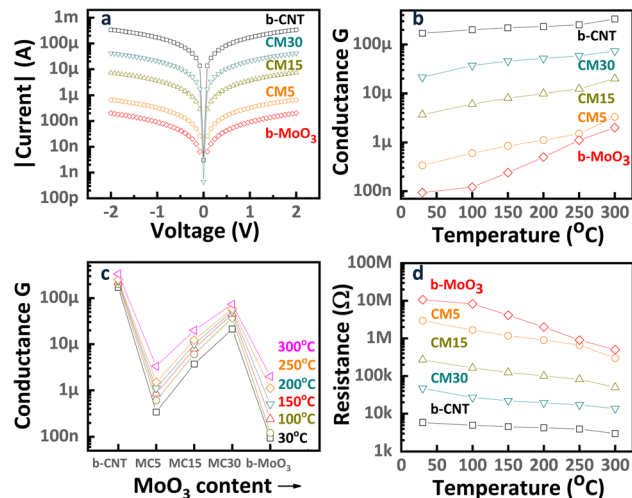


Fig. 4 (a) Current–voltage (I – V) characteristics of b-CNT, b-MoO₃, CM5, CM15, and CM30 sensors at RT. (b) The conductance of the sensors from 30 °C to 300 °C. (c) The conductance of the sensors is displayed as a function of increasing MoO₃ content and measuring temperatures. (d) The resistance behavior of the sensors from 30 °C to 300 °C.

of b-CNT, b-MoO₃, and CNT/MoO₃ nanohybrids upon exposure to 1 ppm ethanol in dry air are depicted in Fig. 5. Considering the sensor response S , samples can be categorized by the positive response group (Fig. 5a–d for b-CNT, CM5, CM15, CM30) and the negative response group (Fig. 5e for b-MoO₃). Since the major charge carriers determine the sensing polarity, the former group shows a *CNT-dominated* conduction mechanism with holes as major charge carriers (p-type). Meanwhile, *MoO₃-dominated* conduction with electrons as main charge carriers (n-type) occurs in the latter group. After exposure to ethanol (a reducing gas), the increasing resistance is considered to be a p-type semiconducting behavior. Therefore, the b-CNT, CM5, CM15, and CM30 sensors reveal *CNT-dominated* conduction. Whereas the resistance of the b-MoO₃ sensor decreases upon exposure to ethanol gas, indicating that the n-type conduction process is predominantly governed by the MoO₃ component. The sensor responses exhibit rapid kinetics, which reaches sensing saturation within 5 minutes. The behavior of response dependence was replotted in Fig. 5f and 6a to analyze the effect of MoO₃ content on response systematically. As the b-CNT and CNT-MoO₃ hybrid sensors operate at a higher temperature, less response can be achieved (Fig. 6b). In contrast, the MoO₃-dominated sensor, b-MoO₃, showed the opposite tendency. This difference originates from the difference in sensing mechanisms of CNTs and MoO₃.

However, the highest sensitivity of the b-MoO₃ was recorded at 250 °C, which is a general observation for most oxide semiconductors. As shown in Fig. 6a and b, b-CNT and b-MoO₃ revealed a small response of 12.5% and 4% at RT, respectively. Interestingly, forming the nanohybrids with MoO₃ and CNTs led to a synergistic effect with the highest response level obtained for the CM15, with a response level of 76.4% at RT, which is much higher than the b-CNT and b-MoO₃ sensors. The



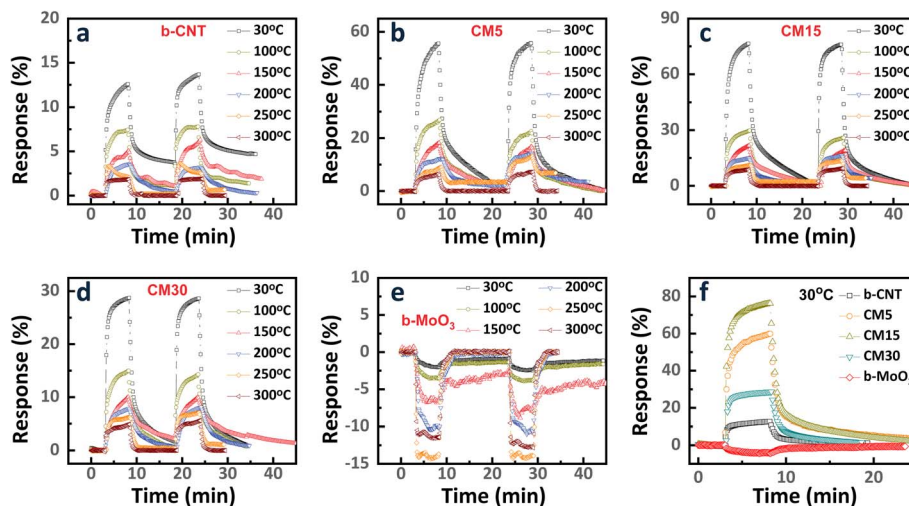


Fig. 5 The response and recovery kinetic curves of (a) b-CNT, (b) CM5, (c) CM15, (d) CM30, and (e) b-MoO₃ sensors to 1 ppm ethanol were measured at varying operating temperatures ranging from 30 °C to 300 °C. (f) The sensor response to 1 ppm ethanol as a function of MoO₃ content at 30 °C.

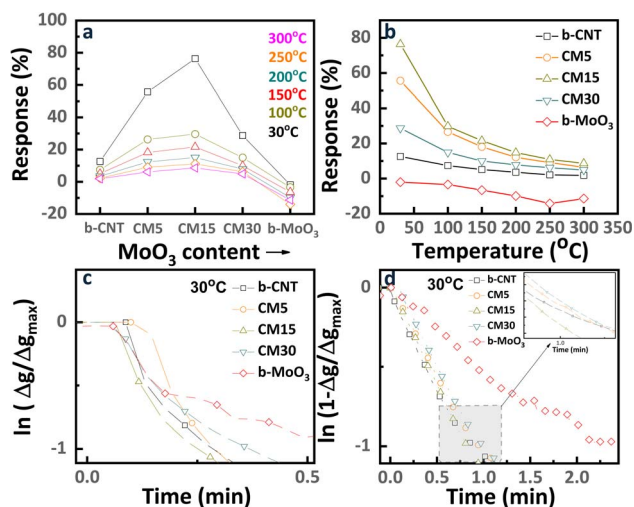


Fig. 6 (a) Response/recovery kinetics at 30 °C and (b) summary of response levels at different operating temperatures from 30 °C to 300 °C of b-CNT, CM5, CM15, CM30, and b-MoO₃ sensors to 1 ppm ethanol. Plots of (c) response time and (d) recovery time following the exponential rise and decay behaviors of the response and recovery kinetics.^{44,45}

summary of response/recovery curves of the sensors at RT is plotted in Fig. 5f.

The response time and recovery time of the five sensors are displayed in Fig. 6c and d. The CM15 sensor shows the fastest behavior of 14.4 seconds and 66 seconds in both response and recovery time, respectively, compared with other sensors. In addition, Table 1 lists the comparison between this study and other materials along with their corresponding response toward ethanol exposure.

Given that the CM15 hybrid sensor demonstrated the highest ethanol sensing performance at RT, its sensing behavior was further analyzed across various ethanol gas concentrations (0.1,

0.2, 0.5, and 1 ppm) under the same conditions. As depicted in Fig. 7 and its inset, the sensor exhibited a linear response with respect to ethanol concentration, validating the reliability and consistency of the sensing performance. The linear response can be provided due to enough active surface sites to facilitate the adsorption of ethanol molecules and ensure proportionality in the sensing mechanism.⁵² In addition, Fig. 7b shows the multiple repeatable cycles of the CM15 sensor upon 100 ppb of ethanol. The good repeatability of the CM15 sensor was attained even upon exposure to the low concentration of ethanol, 100 ppb. Finally, the response kinetics of the CM15 sensor to 3 ppm of H₂S, NH₃, H₂, CO, CH₄, methanol, acetone, and acetic acid gases at RT, and a comparison of its response levels with ethanol sensing are presented in Fig. 7c and d, respectively. The CNT/MoO₃ nanohybrid exhibits good selectivity for ethanol gas detection. The detection performance of ethanol is governed by factors such as dissociation energy, operating temperature, and the properties of the sensing material, including surface defects, morphology, and chemical composition. Accordingly, further optimization of the CNT/MoO₃ nanohybrid sensor can lead to enhanced performance by fine-tuning these critical parameters.

3.3. Gas sensing mechanism

The sensing signal is determined by the chemical adsorption reaction between the ethanol molecules and the pre-adsorbed oxygen ions on the sensor material surface. This interaction modulates the carrier concentration, resulting in changes to the electrical conductivity of the sensors. Initially, negative-charged chemisorbed oxygen ions (O₂⁻) can be formed through the adsorbed oxygen molecules on the surface of the materials by capturing free electrons from the conduction band of the sensing materials (eqn (1)). Thereby, a depletion layer is generated on the surface of MoO₃ nanoparticles due to low surface electron density while the oxygen adsorption renders an



Table 1 Gas sensitivity of different sensing materials toward ethanol exposure in literature and this work

Sensor	Concentration (ppm)	Operating temperature (°C)	Response (%)	Response time/recovery time (s)	Ref.
MoO ₃ /rGO nanoflakes	100	310	3700	4/13	46
Zn/MoO ₃ nanobelts	250	240	4900	48/76	47
MoO ₃ /MXene nanocomposites	50	100	453	46/276	48
MoO ₃ nanoplate	100	400	1200	22/15	49
MoO ₃ micro rods	500	332	750	10/10	50
MoO ₃ nanorods/SnO ₂ nanosheets	100	200	4764	65/230	18
MoO ₂ /MoO ₃ /MXene nanocomposites	200	RT	1870	42/297	51
CNT/MoO ₃ nanohybrids	1	RT	76.5	14/66	This work

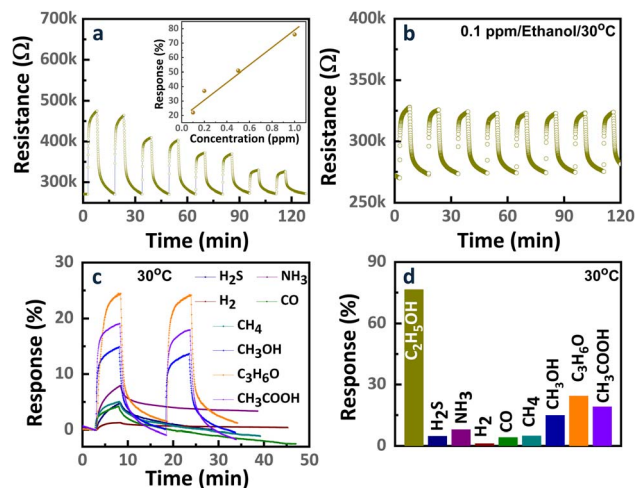
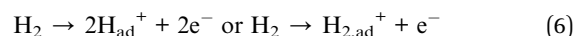
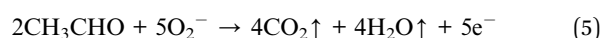
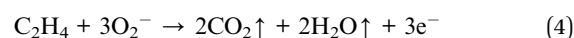
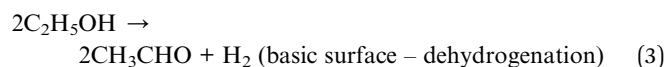
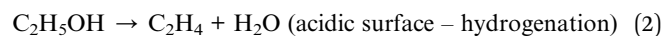
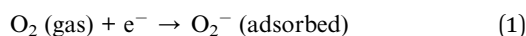


Fig. 7 (a) Sensing response/recovery curves of the CM15 sensor against various ethanol concentrations (0.1, 0.2, 0.5, and 1 ppm), and summary of response levels in the inset. (b) Repeatability for 8 cycles of the CM15 sensor for 0.1 ppm ethanol at 30 °C. (c) Response kinetics of the CM15 sensor to 3 ppm of H₂S, NH₃, H₂, CO, CH₄, methanol, acetone, and acetic acid gases at RT, and (d) comparison of response levels with ethanol sensing.

increase of hole concentration on CNTs surface, which leads to a boost in the conductivity of CNTs. Furthermore, after reducing gas (ethanol) exposure, the removal of chemisorbed oxygen causes the resistance of the p-type dominant CNT-MoO₃ sensor to increase. As a product of these redox reactions, the trapped electrons released back to the material conduction band and reduced the hole carrier concentration, resulting in a dramatic increase in resistance. Ethanol, a volatile organic compound, can be decomposed by dehydrogenation and dehydration on basic and acidic surface materials,^{53–55} respectively. The methodical decomposition reaction leads to the generation of CO₂ and H₂O as presented in eqn (2)–(5).^{56–58} However, the decomposition reactions on the acidic and basic surfaces result in the formation of intermediate products, alkenes, and aldehydes. They originate from bond strength, polar surface, nonpolar surface, anions vacancies, stoichiometric sites, and steric effects of C–H and C–O in ethanol.⁵³



The released electrons from redox reactions then recombine with CNT holes thus lowering the conductance of the CNT. The charge modulation accompanied by the ethanol adsorption formed the depletion region, as shown in Fig. 9. Upon exposure to 1 ppm ethanol, the resistance of CNTs increases, as indicated by a 12.5% sensing signal at RT. Hence, the resistance further decreases at the elevated temperature (Fig. 5f) owing to the higher rate of ethanol desorption. Meanwhile, b-MoO₃ shows negligible response to ethanol at room temperature, as illustrated in Fig. 8a. The graph demonstrates that the agglomerated MoO₃ has neutral cores with minimal surface modification following ethanol exposure, resulting in negligible detectable signal.

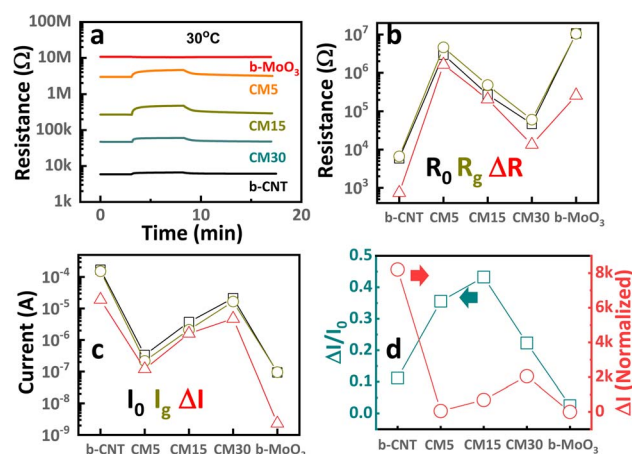


Fig. 8 (a) Transient resistance curves of the sensing structures toward 1 ppm ethanol exposure at 30 °C. (b) The corresponding R_0 , R_g , and ΔR . (c) The corresponding I_0 , I_g , and ΔI was converted from (b). (d) The $\Delta I/I_0$ values and the ΔI normalized to that of b-MoO₃.



As confirmed by our observation, the sensing signal of the hybrid is enhanced with the addition of MoO₃ on CNTs, especially represented by ~76.2% in the highest response with CM15 at RT (Fig. 6a). The ethanol sensing signal is defined as the ratio of the sensor resistances before and after ethanol molecule adsorption, denoted as R_0 and R_g , respectively. Fig. 8b presents the raw data of R_0 , R_g , and $\Delta R = R_g - R_0$. Alternatively, the data can be expressed in terms of currents I_0 and I_g , and the change in the current $\Delta I = I_0 - I_g$, where I_0 and I_g correspond to the current in the absence and presence of the analyte gas, respectively. This approach is useful because the current is the flow rate of the main charge carriers resulting from the gas-surface interactions. Fig. 8c is derived from resistance (Fig. 8b) using the corresponding current values measured at 1 V. The current variation can be analyzed in two forms: $\Delta I/I_0$, which reflects the carrier charge ratio and serves as a measure of sensitivity, and ΔI , which indicates the absolute number of carrier charges transferred during the response, normalized to that of b-MoO₃. The $\Delta I/I_0$ value is significantly low for CNTs compared to the hybrid samples. Assuming all charges are consumed for the surface reaction with ethanol, upon ethanol exposure, only 12% of the holes in the CNT participate in the surface reaction. Meanwhile, 35%, 43%, and 22% of the holes in the hybrid materials (CM5, CM15, and CM30, respectively) are involved in the reaction (Fig. 8d). The corresponding number for b-MoO₃ is only 2.5%. The standby condition of the CM15 with the electron-depleted region of MoO₃ nanoparticles after contact with CNTs is schemed in Fig. 9b. The improved response of CM15 is attributed to the reduced conductivity of the CNTs, followed by the significant increase in the ethanol adsorption sites at the interference are toward the outer surface. As more MoO₃ particles contact CNTs, the further expanded junction area results in smaller conductivity in CNTs. In turn, the number of ethanol adsorption sites increases proportionally, ultimately resulting in the complete depletion of CNT holes. Ethanol adsorption is determined by the availability of

holes in the CNTs, which restricts further adsorption. Therefore, the higher sensing response of CM15 is attributed to the optimal ratio of conductance before and after ethanol exposure, which corresponds to the highest difference between the initial and post-exposure conductivity states (depicted on the I_0 and I_g , ΔI schematics of Fig. 9b). In summary, the incorporation of MoO₃ into CNT networks enhances the number of active adsorption sites while simultaneously reducing the overall conductivity of the hybrid, achieving the maximum response in CM15. However, with further MoO₃ additions like CM30, the excessive MoO₃ content renders adsorption sites over-saturated. The excessively large numbers of MoO₃ particles are electrically isolated from the CNTs network, thereby disturbing the charge conduction of the hybrids. Moreover, the complete depletion of holes from the CNTs under this condition results in a significant drop in sensing performance. Only ~20% sensing response could be attained with the CM30 hybrid. Moreover, after ethanol exposure, the released electrons recombined with the holes in CNTs (process A) or moved through the surface route to Pt electrodes (process A') (Fig. 9a). However, process A hardly occurred due to the restricted number of available holes in CNTs and the low probability of intensive migration through particles. Hence, only 12% of the holes in CNT could be depleted during the gas reaction. In the recovery state, the electrons will be provided either by the electron-hole pair generation process (B) or by transporting electrons from Pt electrodes (process B'). Both processes reveal the high energy barrier and thus lead to slow recovery. Meanwhile, the MoO₃ nanoparticles are attached to the CNT bundles in CM15 as presented in Fig. 9b, in which released electrons from the reactions as well as from MoO₃, can easily recombine with the holes of the CNTs. During the recovery process, the electron transfer rate to the CNT/MoO₃ interference area is also enhanced through band bending and good heterojunction formation between CNT and MoO₃. Therefore, both the response and recovery behavior of CM15 are significantly

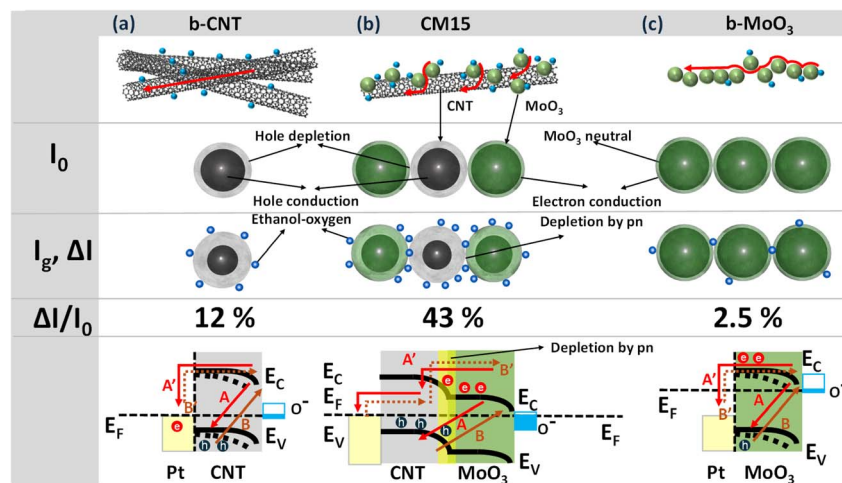


Fig. 9 Schematics and energy band diagrams of (a) b-CNT, (b) CM15, and (c) b-MoO₃ with the modified conductivity and depleted region before (I_0) and after (I_g) 1 ppm ethanol exposure at RT. Blue dots represent the ethanol reaction, while red arrows correspond to the travel distance of main charge carriers.



improved, as depicted in Fig. 6d. According to reported values, the work function of CNTs (4.95 eV) exceeds that of MoO₃ (3.95 eV), facilitating electron transfer from MoO₃ to CNTs at their interface. This electron transfer further enhances the functional properties of hybrids, particularly for applications involving gas sensing.

To further elucidate the sensing mechanism, the ethanol detection ability of CM15 was investigated upon exposure to 1 ppm ethanol diluted in N₂ (in the absence of oxygen) at RT. The response curve, illustrated in Fig. S1, showed a response of ~57%, confirming that ethanol adsorption follows the mechanism described in eqn (6). Additionally, this observation corroborates that ethanol adsorption reveals the oxygen-independent adsorption pathway in CNT/MoO₃ nanohybrids for ethanol sensing at room temperature. Without the above-mentioned methodical decomposition reaction, enhanced ethanol detection is still clearly observed, which is due to the following reasonable explanation.

Firstly, of particular note is that the CNT-dominated and MoO₃-dominated sensors show a decrease and an increase in sensing response, respectively, as the temperature increases. At RT, the CNTs adsorb ethanol *via* eqn (4)–(6) while MoO₃ reveals negligible adsorption. However, at the elevated temperature, the ethanol adsorption and the oxygen ionosorption process of the CNTs are small (as illustrated in Fig. 6a and b) while ethanol dominantly reacted with ionosorbed oxygen on MoO₃. The simultaneous combustion reaction can occur at the elevated temperature as followed by eqn (4) and (5).^{59–61} Thus, the optimum response of b-MoO₃ can be achieved at 250 °C, which indicates that the hybrid formation with CNTs in MoO₃-dominated sensors does not result in any significant synergy effect at a high temperature. However, the combustion reaction contributed to the enhancement of the hybrid sensing behavior in the CNT-dominated sensors with an appropriately small number of MoO₃ nanoparticles. Since b-CNT does not show any response at the elevated temperature, our observation confirms that electrons released during the combustion reaction at the MoO₃ nanoparticles are transported to the surface of CNTs, subsequently exhibiting p-type semiconductor behavior.

Secondly, Fig. 10 presents the nitrogen adsorption–desorption isotherms to analyze the pore size distribution and specific surface area of b-CNT, b-MoO₃, and CM15. The adsorption–

desorption curves exhibit type IV isotherms,⁶² being indicative of mesoporous structures. Based on the Brunauer–Emmett–Teller (BET) equation, the specific surface areas of b-MoO₃, b-CNT, and CM15 are determined to be 23.1 m² g⁻¹, 52.4 m² g⁻¹, and 72.12 m² g⁻¹, respectively. Additionally, the pore volume of CM15 is measured at 0.073 cm³ g⁻¹, which is significantly higher than that of b-MoO₃ (0.013 cm³ g⁻¹) and b-CNT (0.057 cm³ g⁻¹). The enhanced specific surface area and pore volume of CM15 suggest an abundance of active sites for oxygen adsorption owing to the hybrid 0D/1D structure of CM15, which is advantageous for gas interaction on the surface. Such improvement directly contributes to enhanced gas detection capabilities.⁶²

Thirdly, ethanol exhibits relatively high adsorption energy ($E_{\text{ads}} = -0.96$ eV mol⁻¹) compared to the adsorption energy of surface-adsorbed oxygen species ($E_{\text{ads}} \sim -0.35$ eV mol⁻¹).⁶³ This significant difference facilitates preferential interaction between ethanol molecules and the adsorbed oxygen species on the material surface. Such interaction plays a critical role in enhancing the gas detection performance by promoting more effective surface reactions.

Fourthly, the semiconducting characteristics of MoO₃ on the surface of CNTs serve as an effective catalyst for oxygen dissociation. This catalytic activity boosts the diffusion kinetics of ethanol molecules toward surface vacancies, thereby facilitating rapid surface reactions. As a result, the hybrid material exhibits good sensitivity and fast response to ethanol detection even at room temperature.⁶⁴

And lastly, the exceptional electrical conductivity of CNTs significantly lowers the operating temperature of sensors, enhancing their energy efficiency. Furthermore, the lowest unoccupied molecular orbital (LUMO) energy of ethanol (0.13 eV),^{65,66} accelerates electron transfer between the nanohybrid material and ethanol gas molecules. This facile electron interaction underpins the high sensing selectivity for ethanol compared to other gases.

3.4. Sensor performance against humidity effect

For the gas sensor operation at RT, the effect of ambient moisture must be considered. Fig. 11a depicts the influence of relative humidity (RH) on the resistance of sensors, which

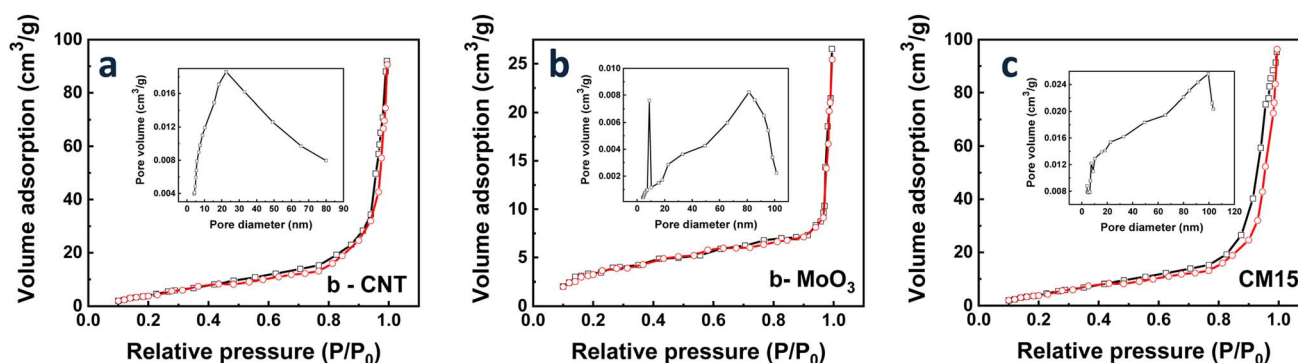


Fig. 10 The nitrogen adsorption and desorption isotherms of (a) b-CNT, (b) b-MoO₃, and (c) CM15 (the insets are the pore size distribution).



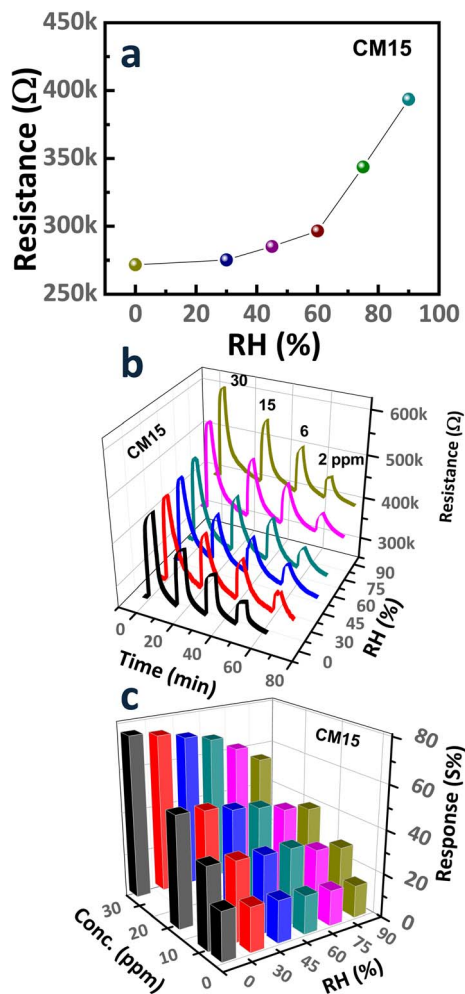


Fig. 11 (a) Resistance of the CM15 sensor under varying RH conditions. (b) Transient response curves and (c) corresponding response levels of the CM15 sensor to ethanol concentrations (0.1–1 ppm) at different RH levels (0%, 30%, 45%, 60%, 75%, and 90%) at RT.

exhibits a progressive increase in resistance as RH rises from 0% to 90%. Specifically, the baseline resistance of the sensor increased from ~ 271.7 k Ω at 0% RH to ~ 393.6 k Ω at 90% RH, thereby deteriorating the sensor response. This humidity effect is ascribed to the n-type doping effect or the adsorption of water molecules on the sensor surface, which alters the surface charge dynamics and interferes with gas adsorption processes.⁶⁷ Water molecules undergo dissociation on the sensor surface, leading to the generation of hydroxyl species accompanied by the donation of electrons. The electron donation reduces the hole concentration in the p-type CNT/MoO₃ nanohybrid, which decreases its electrical conductivity.^{68,69}

The transient response curves and corresponding response levels of the CM15 sensor toward various ethanol concentrations (0.1–1 ppm) under different RH levels (0%, 30%, 45%, 60%, 75%, and 90%) at RT are presented in Fig. 11b and c. Water adsorption on the sensor surface alters the adsorbed species and lowers the response level. Specifically, the response of the CM15 sensor to 0.1 ppm ethanol gradually decreased

from 22% to 14.6% as RH increased from 0% to 90%. These observations highlight that the sensor exhibits dominant ethanol-sensing characteristics at lower humidity levels. However, the dependence on humidity remains relatively weak for the nanohybrid sensor operating at RT, which is enabled by CNTs. CNTs demonstrate immunity to water molecule adsorption, contrasting with the pronounced humidity sensitivity observed in metal oxides as reported in the literature.^{8,30,70–73} This relatively low dependence on humidity represents an additional advantage of the CNT/MoO₃ nanohybrid sensor for ethanol detection at ambient conditions.

4. Conclusion

The CNT/MoO₃ nanohybrid was successfully synthesized using an arc-discharge method. The CNT content in CNT/MoO₃ nanohybrids was tuned by adjusting the deposition time. Characterization through SEM, TEM, XRD, and BET analyses confirmed that the composites reveal a mesoporous structure, consisting of 0D MoO₃ particles encapsulated on 1D CNTs. The CM15 sensor demonstrated the highest sensing performance in terms of response time, recovery time, and selectivity. The charge carrier transfer mechanism, involving both electrons and holes, is modified by the formation of a depletion layer in the nanohybrid. Compared to bare CNT and bare MoO₃, the CNT/MoO₃ nanohybrid exhibited a higher specific surface area, providing more surface-active sites, which enhanced its sensitivity. We believe that our nanohybrid materials show significant potential for the development of advanced ethanol detection sensors.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information is available. See DOI: <https://doi.org/10.1039/d6ra00372a>.

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References

- 1 Y. Wang, H. Li, D. Huang, X. Wang, L. Cai, Y. Chen, W. Wang, Y. Song, G. Han and B. Zhen, A high-performance ethanol gas sensor based on Ce-doped SnO₂ nanomaterials prepared by the Pechini method, *Mater. Sci. Semicond. Process.*, 2022, **137**, 106188, DOI: [10.1016/j.mssp.2021.106188](https://doi.org/10.1016/j.mssp.2021.106188).
- 2 J. Zhang, S. Ma, B. Wang and S. Pei, Hydrothermal synthesis of SnO₂-CuO composite nanoparticles as a fast-response



- ethanol gas sensor, *J. Alloys Compd.*, 2021, **886**, 161299, DOI: [10.1016/j.jallcom.2021.161299](https://doi.org/10.1016/j.jallcom.2021.161299).
- 3 N. Devabharathi, R. Parasuraman, A. M. Umarji and S. Dasgupta, Ultra-high response ethanol sensors from fully-printed co-continuous and mesoporous tin oxide thin films, *J. Alloys Compd.*, 2021, **865**, 158815, DOI: [10.1016/j.jallcom.2021.158815](https://doi.org/10.1016/j.jallcom.2021.158815).
 - 4 B. Jiang, J. Lu, W. Han, Y. Sun, Y. Wang, P. Cheng, H. Zhang, C. Wang and G. Lu, Hierarchical mesoporous zinc oxide microspheres for ethanol gas sensor, *Sens. Actuators, B*, 2022, **357**, 131333.
 - 5 M. Kočí, P. S. Wrobel, M. Godzierz, O. Szabó, S. Pusz, Š. Potocký, M. Husák and A. Kromka, Highly Sensitive Gas and Ethanol Vapor Sensors Based on Carbon Heterostructures for Room Temperature Detection, *ACS Appl. Mater. Interfaces*, 2025, **17**, 14703–14715, DOI: [10.1021/acsami.4c21591](https://doi.org/10.1021/acsami.4c21591).
 - 6 N. M. Hieu, H. Kim, C. Kim, S.-K. Hong and D. Kim, A hydrogen sulfide gas sensor based on pd-decorated ZnO nanorods, *J. Nanosci. Nanotechnol.*, 2016, **16**, 10351–10355.
 - 7 N. M. Hieu, C. Van Phuoc, T. T. Hien, N. D. Chinh, N. D. Quang, C. Kim, J.-R. Jeong and D. Kim, A separated receptor/transducer scheme as strategy to enhance the gas sensing performance using hematite-carbon nanotube composite, *Sensors*, 2019, **19**, 3915.
 - 8 N. Duc Chinh, Y. Haneul, N. Minh Hieu, N. Manh Hung, N. Duc Quang, C. Kim and D. Kim, pn-Heterojunction of the SWCNT/ZnO nanocomposite for temperature dependent reaction with hydrogen, *J. Colloid Interface Sci.*, 2021, **584**, 582–591, DOI: [10.1016/j.jcis.2020.10.017](https://doi.org/10.1016/j.jcis.2020.10.017).
 - 9 N. M. Hieu, T. T. Hien, N. D. Chinh, N. D. Quang, N. M. Hung, C. Van Phuoc, S.-M. Lee, J.-R. Jeong, C. Kim and D. Kim, ZnTe-coated ZnO nanorods: Hydrogen sulfide nano-sensor purely controlled by pn junction, *Mater. Des.*, 2020, **191**, 108628.
 - 10 N. M. Hieu, C. Van Phuoc, C. V. Anh, N. M. Hung, A. D. Phan, N. D. Chinh, S. Majumder, T. H. Cuong, H. G. Chuc and T. Nguyen, The kinetics of a light irradiation enhanced room temperature NO₂ gas sensor using hybrid ZnO/ZnTe nanorod structures, *RSC Adv.*, 2024, **14**, 39418–39428.
 - 11 N. D. Chinh, N. D. Quang, H. Lee, T. T. Hien, N. M. Hieu, D. Kim, C. Kim and D. Kim, NO gas sensing kinetics at room temperature under UV light irradiation of In₂O₃ nanostructures, *Sci. Rep.*, 2016, **6**, 1–11, DOI: [10.1038/srep35066](https://doi.org/10.1038/srep35066).
 - 12 N. M. Hung, N. M. Hieu, N. D. Chinh, T. T. Hien, N. D. Quang, S. Majumder, G. S. Choi, C. Kim and D. Kim, Rb₂CO₃-decorated In₂O₃ nanoparticles for the room-temperature detection of sub-ppm level NO₂, *Sens. Actuators, B*, 2020, **313**, 128001, DOI: [10.1016/j.snb.2020.128001](https://doi.org/10.1016/j.snb.2020.128001).
 - 13 Z. Wang, K. Zhang, T. Fei, F. Gu and D. Han, α -Fe₂O₃/NiO heterojunction nanorods with enhanced gas sensing performance for acetone, *Sens. Actuators, B*, 2020, **318**, 128191, DOI: [10.1016/j.snb.2020.128191](https://doi.org/10.1016/j.snb.2020.128191).
 - 14 Z. Tang, X. Deng, Y. Zhang, X. Guo, J. Yang, C. Zhu, J. fan, Y. Shi, B. Qing and F. Fan, MoO₃ nanoflakes coupled reduced graphene oxide with enhanced ethanol sensing performance and mechanism, *Sens. Actuators, B*, 2019, **297**, 126730, DOI: [10.1016/j.snb.2019.126730](https://doi.org/10.1016/j.snb.2019.126730).
 - 15 X. Hu, L. Chen and Y. Chen, Universal and Versatile MoO₃-Based Hole Transport Layers for Efficient and Stable Polymer Solar Cells, *J. Phys. Chem. C*, 2014, **118**, 9930–9938, DOI: [10.1021/jp501995t](https://doi.org/10.1021/jp501995t).
 - 16 M. M. Mohamed, T. M. Salama, M. Morsy, R. M. Abou Shahba and S. H. Mohamed, Facile strategy of synthesizing α -MoO_{3-x} nanorods boosted as traced by 1% graphene oxide: Efficient visible light photocatalysis and gas sensing applications, *Sens. Actuators, B*, 2019, **299**, 126960.
 - 17 S. Barazzouk, R. P. Tandon and S. Hotchandani, MoO₃-based sensor for NO, NO₂ and CH₄ detection, *Sens. Actuators, B*, 2006, **119**, 691–694.
 - 18 X. Wang, Y. Gao, Q. Zhang, X. He and X. Wang, Synthesis of MoO₃ (1D)@SnO₂ (2D) core-shell heterostructures for enhanced ethanol gas sensing performance, *Sens. Actuators, B*, 2023, **382**, 133484.
 - 19 H.-Y. Li, L. Huang, X.-X. Wang, C.-S. Lee, J.-W. Yoon, J. Zhou, X. Guo and J.-H. Lee, Molybdenum trioxide nanopaper as a dual gas sensor for detecting trimethylamine and hydrogen sulfide, *RSC Adv.*, 2017, **7**, 3680–3685.
 - 20 C. Su, L. Zhang, Y. Han, C. Ren, X. Chen, J. Hu, M. Zeng, N. Hu, Y. Su, Z. Zhou and Z. Yang, Controllable synthesis of crescent-shaped porous NiO nanoplates for conductometric ethanol gas sensors, *Sens. Actuators, B*, 2019, **296**, 126642, DOI: [10.1016/j.snb.2019.126642](https://doi.org/10.1016/j.snb.2019.126642).
 - 21 B. Sharma, K. Karuppasamy, D. Vikraman, E.-B. Jo, P. Sivakumar and H.-S. Kim, Porous, 3D-hierarchical α -NiMoO₄ rectangular nanosheets for selective conductometric ethanol gas sensors, *Sens. Actuators, B*, 2021, **347**, 130615, DOI: [10.1016/j.snb.2021.130615](https://doi.org/10.1016/j.snb.2021.130615).
 - 22 S. Iijima, Helical microtubules of graphitic carbon, *Nature*, 1991, **354**, 56–58, DOI: [10.1038/354056a0](https://doi.org/10.1038/354056a0).
 - 23 S. Iijima and T. Ichihashi, Single-shell carbon nanotubes of 1-nm diameter, *Nature*, 1993, **363**, 603–605.
 - 24 B.-Y. Wei, M.-C. Hsu, P.-G. Su, H.-M. Lin, R.-J. Wu and H.-J. Lai, A novel SnO₂ gas sensor doped with carbon nanotubes operating at room temperature, *Sens. Actuators, B*, 2004, **101**, 81–89, DOI: [10.1016/j.snb.2004.02.028](https://doi.org/10.1016/j.snb.2004.02.028).
 - 25 D. Eder, Carbon Nanotube–Inorganic Hybrids, *Chem. Rev.*, 2010, **110**, 1348–1385, DOI: [10.1021/cr800433k](https://doi.org/10.1021/cr800433k).
 - 26 S. Moon, N. M. Vuong, D. Lee, D. Kim, H. Lee, D. Kim, S.-K. Hong and S.-G. Yoon, Co₃O₄-SWCNT composites for H₂S gas sensor application, *Sens. Actuators, B*, 2016, **222**, 166–172, DOI: [10.1016/j.snb.2015.08.072](https://doi.org/10.1016/j.snb.2015.08.072).
 - 27 N. D. Hoa, N. Van Quy and D. Kim, Nanowire structured SnO_x-SWNT composites: High performance sensor for NO_x detection, *Sens. Actuators, B*, 2009, **142**, 253–259, DOI: [10.1016/j.snb.2009.07.053](https://doi.org/10.1016/j.snb.2009.07.053).
 - 28 S. B. Naghadeh, S. Vahdatifar, Y. Mortazavi, A. A. Khodadadi and A. Abbasi, Functionalized MWCNTs effects on dramatic enhancement of MWCNTs/SnO₂ nanocomposite gas sensing properties at low temperatures, *Sens. Actuators, B*, 2016, **223**, 252–260, DOI: [10.1016/j.snb.2015.09.088](https://doi.org/10.1016/j.snb.2015.09.088).



- 29 G. W. Thomson, The Antoine equation for vapor-pressure data, *Chem. Rev.*, 1946, **38**, 1–39.
- 30 N. D. Chinh, N. M. Hung, S. Majumder, C. Kim and D. Kim, Hole-supply-rate-controlled methanol-gas-sensing reaction over p-type Co_3O_4 /single-walled carbon nanotube hybrid structures, *Sens. Actuators, B*, 2021, **326**, 128956.
- 31 S. Lefrant, I. Baltog and M. Baibarac, Surface-enhanced Raman scattering studies on chemically transformed carbon nanotube thin films, *J. Raman Spectrosc.*, 2005, **36**, 676–698.
- 32 M. S. Dresselhaus, G. Dresselhaus, A. Jorio, A. G. Souza Filho and R. Saito, Raman spectroscopy on isolated single wall carbon nanotubes, *Carbon*, 2002, **40**, 2043–2061.
- 33 F. Ji, X. Ren, X. Zheng, Y. Liu, L. Pang, J. Jiang and S. F. Liu, 2D- MoO_3 nanosheets for superior gas sensors, *Nanoscale*, 2016, **8**, 8696–8703.
- 34 M. Diskus, O. Nilsen, H. Fjellvåg, S. Diplas, P. Beato, C. Harvey, E. van Schroyen, L. Lantman and B. M. Weckhuysen, Combination of characterization techniques for atomic layer deposition MoO_3 coatings: From the amorphous to the orthorhombic α - MoO_3 crystalline phase, *J. Vac. Sci. Technol., A*, 2012, **30**, 01A107.
- 35 R. Ji, J. Cheng, X. Yang, J. Yu and L. Li, Enhanced charge carrier transport in spray-cast organic solar cells using solution processed MoO_3 micro arrays, *RSC Adv.*, 2017, **7**, 3059–3065.
- 36 T. I. T. Okpalugo, P. Papakonstantinou, H. Murphy, J. McLaughlin and N. M. D. Brown, High resolution XPS characterization of chemical functionalised MWCNTs and SWCNTs, *Carbon*, 2005, **43**, 153–161.
- 37 L.-C. Wang, K.-T. Tang, I.-J. Teng, C.-T. Kuo, C.-L. Ho, H.-W. Kuo, T.-H. Su, S.-R. Yang, G.-N. Shi and C.-P. Chang, A single-walled carbon nanotube network gas sensing device, *Sensors*, 2011, **11**, 7763–7772.
- 38 B. Arash and Q. Wang, Detection of gas atoms with carbon nanotubes, *Sci. Rep.*, 2013, **3**, 1782.
- 39 Y. Wang and J. T. W. Yeow, A review of carbon nanotubes-based gas sensors, *J. Sens.*, 2009, **2009**, 493904.
- 40 S. S. Sunu, E. Prabhu, V. Jayaraman, K. I. Gnanasekar, T. K. Seshagiri and T. Gnanasekaran, Electrical conductivity and gas sensing properties of MoO_3 , *Sens. Actuators, B*, 2004, **101**, 161–174.
- 41 A. K. Prasad and P. I. Gouma, MoO_3 and WO_3 based thin film conductometric sensors for automotive applications, *J. Mater. Sci.*, 2003, **38**, 4347–4352.
- 42 R. Rohilla, J. Prakash and K. Dasgupta, Review of Carbon Nanotube/Graphene Nanosheet Chemiresistive Gas Sensors, *ACS Appl. Nano Mater.*, 2025, **8**, 23414–23465, DOI: [10.1021/acsanm.5c04068](https://doi.org/10.1021/acsanm.5c04068).
- 43 R. Rohilla, J. Prakash, P. T. Rao, A. Pathak and K. Dasgupta, Synergistic sensing properties of a standalone portable prototype using an integrated graphitic carbon nitride-carbon nanotube film for ultra-sensitive and selective acetone detection, *J. Mater. Chem. C*, 2025, **13**, 18350–18364, DOI: [10.1039/D5TC02355F](https://doi.org/10.1039/D5TC02355F).
- 44 N. M. Vuong, D. Kim and H. Kim, Surface gas sensing kinetics of a WO_3 nanowire sensor: part 1—oxidizing gases, *Sens. Actuators, B*, 2015, **220**, 932–941, DOI: [10.1016/j.snb.2015.06.031](https://doi.org/10.1016/j.snb.2015.06.031).
- 45 N. M. Vuong, D. Kim and H. Kim, Surface gas sensing kinetics of a WO_3 nanowire sensor: Part 2—Reducing gases, *Sens. Actuators, B*, 2016, **224**, 425–433, DOI: [10.1016/j.snb.2015.10.070](https://doi.org/10.1016/j.snb.2015.10.070).
- 46 Z. Tang, X. Deng, Y. Zhang, X. Guo, J. Yang, C. Zhu, Y. Shi, B. Qing and F. Fan, MoO_3 nanoflakes coupled reduced graphene oxide with enhanced ethanol sensing performance and mechanism, *Sens. Actuators, B*, 2019, **297**, 126730.
- 47 S. Yang, Y. Liu, T. Chen, W. Jin, T. Yang, M. Cao, S. Liu, J. Zhou, G. S. Zakharova and W. Chen, Zn doped MoO_3 nanobelts and the enhanced gas sensing properties to ethanol, *Appl. Surf. Sci.*, 2017, **393**, 377–384, DOI: [10.1016/j.apsusc.2016.10.021](https://doi.org/10.1016/j.apsusc.2016.10.021).
- 48 S. Zhang, P. Song, J. Sun, Y. Ding and Q. Wang, $\text{MoO}_3/\text{Ti}_3\text{C}_2\text{T}_x$ MXene nanocomposites with rapid response for enhanced ethanol-sensing at a low temperature, *Sens. Actuators, B*, 2023, **378**, 133216.
- 49 D. Chen, M. Liu, L. Yin, T. Li, Z. Yang, X. Li, B. Fan, H. Wang, R. Zhang, Z. Li, H. Xu, H. Lu, D. Yang, J. Sun and L. Gao, Single-crystalline MoO_3 nanoplates: topochemical synthesis and enhanced ethanol-sensing performance, *J. Mater. Chem.*, 2011, **21**, 9332–9342, DOI: [10.1039/C1JM11447F](https://doi.org/10.1039/C1JM11447F).
- 50 Y. Liu, S. Yang, Y. Lu, W. Chen and G. S. Zakharova, Hydrothermal synthesis of h- MoO_3 microrods and their gas sensing properties to ethanol, *Appl. Surf. Sci.*, 2015, **359**, 114–119.
- 51 S. Zhang, P. Song, Y. Zheng, Y. Ding and Q. Wang, $\text{MoO}_3/\text{MoO}_3/\text{MXene}$ ternary nanocomposites for high-performance ethanol detection at room temperature, *J. Alloys Compd.*, 2022, **925**, 166663.
- 52 T. Goto, Y. Shimizu, H. Yasuda and T. Ito, Photoexcited ZnO nanoparticles with controlled defects as a highly sensitive oxygen sensor, *Appl. Phys. Lett.*, 2016, **109**, 023104.
- 53 J. M. Vohs and M. A. Barteau, Dehydration and dehydrogenation of ethanol and 1-propanol on the polar surfaces of zinc oxide, *Surf. Sci.*, 1989, **221**, 590–608.
- 54 T. Jinkawa, G. Sakai, J. Tamaki, N. Miura and N. Yamazoe, Relationship between ethanol gas sensitivity and surface catalytic property of tin oxide sensors modified with acidic or basic oxides, *J. Mol. Catal. A: Chem.*, 2000, **155**, 193–200.
- 55 J. F. DeWilde, C. J. Czopinski and A. Bhan, Ethanol dehydration and dehydrogenation on γ - Al_2O_3 : mechanism of acetaldehyde formation, *ACS Catal.*, 2014, **4**, 4425–4433.
- 56 N. Yamazoe, G. Sakai and K. Shimanoe, Oxide semiconductor gas sensors, *Catal. Surv. Asia*, 2003, **7**, 63–75.
- 57 P. Shankar and J. B. B. Rayappan, Gas sensing mechanism of metal oxides: The role of ambient atmosphere, type of semiconductor and gases—A review, *Sci. Lett. J.*, 2015, **4**, 126.
- 58 L. Hu, D. S. Hecht and G. Grüner, Carbon Nanotube Thin Films: Fabrication, Properties, and Applications, *Chem. Rev.*, 2010, **110**, 5790–5844, DOI: [10.1021/cr9002962](https://doi.org/10.1021/cr9002962).



- 59 J. Kong, M. G. Chapline and H. Dai, Functionalized carbon nanotubes for molecular hydrogen sensors, *Adv. Mater.*, 2001, **13**, 1384–1386.
- 60 J. L. Johnson, A. Behnam, S. J. Pearton and A. Ural, Hydrogen sensing using Pd-functionalized multi-layer graphene nanoribbon networks, *Adv. Mater.*, 2010, **22**, 4877–4880.
- 61 T. Hübner, L. Boon-Brett, G. Black and U. Banach, Hydrogen sensors – A review, *Sens. Actuators, B*, 2011, **157**, 329–352, DOI: [10.1016/j.snb.2011.04.070](https://doi.org/10.1016/j.snb.2011.04.070).
- 62 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, Physisorption of gases, with special reference to the evaluation of surface area and pore size distribution (IUPAC Technical Report), *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 63 M. Sinha, R. Mahapatra, M. K. Mondal, S. Krishnamurthy and R. Ghosh, Fast response and low temperature sensing of acetone and ethanol using Al-doped ZnO microrods, *Phys. E*, 2020, **118**, 113868.
- 64 S. Dhall, M. Kumar, M. Bhatnagar and B. R. Mehta, Dual gas sensing properties of graphene-Pd/SnO₂ composites for H₂ and ethanol: Role of nanoparticles-graphene interface, *Int. J. Hydrogen Energy*, 2018, **43**, 17921–17927.
- 65 S. Pei, S. Ma, X. Xu, X. Xu and O. Almamoun, Modulated PrFeO₃ by doping Sm³⁺ for enhanced acetone sensing properties, *J. Alloys Compd.*, 2021, **856**, 158274, DOI: [10.1016/j.jallcom.2020.158274](https://doi.org/10.1016/j.jallcom.2020.158274).
- 66 Z. Wen and L. Tian-Mo, Gas-sensing properties of SnO₂-TiO₂-based sensor for volatile organic compound gas and its sensing mechanism, *Phys. B*, 2010, **405**, 1345–1348.
- 67 Z. Chen and C. Lu, Humidity sensors: a review of materials and mechanisms, *Sens. Lett.*, 2005, **3**, 274–295.
- 68 G. Heiland and D. Kohl, Physical and chemical aspects of oxidic semiconductor gas sensors, *Chem. Sens. Technol.*, 1988, **1**, 15–38.
- 69 Z. Bai, C. Xie, M. Hu, S. Zhang and D. Zeng, Effect of humidity on the gas sensing property of the tetrapod-shaped ZnO nanopowder sensor, *Mater. Sci. Eng., B*, 2008, **149**, 12–17.
- 70 N. D. Chinh, T. T. Hien, L. Do Van, N. M. Hieu, N. D. Quang, S.-M. Lee, C. Kim and D. Kim, Adsorption/desorption kinetics of nitric oxide on zinc oxide nano film sensor enhanced by light irradiation and gold-nanoparticles decoration, *Sens. Actuators, B*, 2019, **281**, 262–272, DOI: [10.1016/j.snb.2018.10.113](https://doi.org/10.1016/j.snb.2018.10.113).
- 71 N. M. Hieu, C. Van Phuoc, N. M. Hung, C. V. Anh, A. D. Phan, J. Nah, J.-R. Jeong, P. T. Huy and D. Kim, A highly stable humidity sensor based on a new Bi₂O₃/CNT hybrid nanostructure, *Sens. Actuators, A*, 2023, **351**, 114141, DOI: [10.1016/j.sna.2022.114141](https://doi.org/10.1016/j.sna.2022.114141).
- 72 N. M. Hieu, C. Van Phuoc, C. V. Anh, N. M. Hung, A. D. Phan, N. D. Chinh, S. Majumder, P. T. Huy, J.-R. Jeong, J. Nah and D. Kim, Humidity suppression in Bi₂O₃/SWCNT nanohybrid structures for room temperature acetone detection, *Ceram. Int.*, 2023, **49**, 16944–16950, DOI: [10.1016/j.ceramint.2023.02.057](https://doi.org/10.1016/j.ceramint.2023.02.057).
- 73 N. D. Chinh, C. Kim and D. Kim, UV-light-activated H₂S gas sensing by a TiO₂ nanoparticulate thin film at room temperature, *J. Alloys Compd.*, 2019, **778**, 247–255.

