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ABSTRACT

The detection and monitoring of harmful toxic volatile organic compound (VOC) gases liberated from various domestic activities is necessary for environmental concern. In this regard, VOC sensors have taken greater demand in domestic environmental monitoring because of their low operating cost, cheap, reliable, low power consumption, improved energy efficiency and reduced emission. In this regard, metal oxide materials in their nano form show growing strength in many research laboratories and public health and so on with their effective chemical, physical, and electronic properties. Therefore, an attempt has been made to study the nano metal oxides for different toxic gases basing on overall research trend like, a good correlation between sensing materials and sensing properties to improve the sensitivity and selectivity of VOC sensors. This paper tries to accumulate the findings of different metal oxide semiconductor nanomaterials used in various types of VOC sensors.

Keywords: Nano materials; metal oxide semiconductors; gaseous pollutants; sensor.

1. INTRODUCTION

To support the life system on Earth there is atmosphere present, which constitutes of a complex dynamics of gaseous system. Thus, any unwanted chemical, particulates, organic or biological material causing damage to the living organism by damaging natural environment or atmosphere is called an air pollutant [1-4].

Air pollution is causing stratospheric ozone layer depletion causing life threats not only to human beings but also to the ecosystem of Earth. Besides this indoor air pollution and urban air quality are listed as two of the world's worst pollution problems [5-8].

There are different sources of pollutants present indoors and outdoors either naturally or due to human activities for different types of gaseous pollutants like Nitrogen oxide (NO₂) [9], carbon monoxide (CO), carbon dioxide (CO₂) [10], formaldehyde (HCHO) [11], methane (CH₄) [12], triethylamine (TEA) [13], Acetone (C_3H_6O) [14], sulphur oxide (SO_2), chlorofluorocarbon (CFCs), odours from garbage, sewage, industrial wastes etc. Amongst these pollutants volatile organic compounds (VOCs) are found more in air of our homes, schools and offices. Basically, it has been estimated that about 50-300 different types of VOCs are found. Some of the high volume of VOC emitting sources include a wide range of personal care products, vent gases during cooking, while making fire, painting the house, cutting the grass, using pesticides, surfactants used during cleaning cloths, dishes and floor etc, while cutting grass during gardening or simply during breathing too. Out of so many VOCs some are found to be highly toxic or carcinogenic in nature and some may have either short term or long term effect both on human health and also on the natural ecosystem. Therefore, extensive research is going on a wide variety of VOC sensors to monitor the indoor and outdoor air quality as it is becoming very essential for today's date.

The word "sensor" finds its origin from the Latin word "sentire" which basically means 'to identify' [15]. A sensor is a device that is capable of identifying a physical quantity and then converting it into a signal type that can be read by an observer or For by an instrument. accuracy purpose sensors are calibrated against known standards. Nowadays sensors are used everywhere like inside cars, machines, aeroplanes, homes, offices, robotics etc. A sensor is a device, which generates a functionally related output usually in the form of an electrical or optical signal corresponding to an input quantity [16, 17].

The sensitivity indicates how much the sensor's output changes when there is a small change measured in input quantity. Sensors that are capable of measuring very small changes have very high sensitivities. Demand for the variety of sensors and the capabilities of technological advancements allowing more and more sensors to be manufactured in different areas. Among all, gas sensing technology has received increasing attention and has become more significant because of its widespread and common applications on the detection and monitoring of flammable and toxic VOCs in domestic as well as in commercial and industrial environment. Thus, the detection of VOC gas molecules with conversion of their concentration information into electrical signals needs the use of sensing technologies in different fields. Among the different active sensing materials, metal oxides have shown effective results towards VOC sensing application. A lot of research studies have been done on VOC sensing applications with the help of metal oxides.

Therefore, in this paper effort has been made to gather findings of different researchers about different metal oxide semiconductor nano-materials used to detect few popular VOCs that are toxic in nature and the most popular pollutants available in and around human surroundings.

2. ROLE OF NANO METAL OXIDES IN VOC SENSING.

Nano metal oxides play a significant role in the gas sensing properties. Mostly the transition metals and their oxides from periodic table III-VI with electronic configuration (d0-d10) mainly exhibit the sensing property. The synthetic procedure used in the material science shows a remarkable change in the properties of nanomaterials and shows a better performance in different applications towards gas sensing. The properties are i) high surface-to-volume ratio facilitates more gas to adsorb onto the surface of nanomaterial for better sensing, ii) introducing defects in the nanomaterial, functionalization are more beneficial in the absorption of gas molecule, iii) proper charge transfer property because of semiconducting nature, iv) catalytic combustion or resistance modulation make nanomaterial suitable candidate for gas sensing.



Scheme 1. Schematic diagram VOC sensing mechanism.

3. DIFFERENT POLLUTANTS AND THEIR DETECTION 3.1. Formaldehyde (CH₂O).

Presence of CH₂O in indoor air within the range (0.2-50 ppm) may cause a serious health problem. The maximum exposure level of CH₂O is 80 ppb per 30 minute as the direction of world health organization (WHO). So, the early detection of CH₂O is highly necessary. In this regard, the development of nano based material plays a key role in the selective detection of CH₂O. Till date a handful of nano based metal oxides are used in the detection of CH₂O. The mechanism of CH₂O mainly depends on chemisorbed oxygen on the surface of the catalyst. The oxygen molecule present in air, first chemisorbed on the catalyst. Again this oxygen species (O₂⁻) react with CH₂O and simultaneously releases of electrons to the catalyst. This leads to a decrease in electron depletion layer (ELD) thickness and thereby increasing the sensing efficiency. The mechanism was shown as follows: $O_2 + 2e^- \rightarrow 2O^-$

 $CH_2O + 2O^-_{(ads)} \rightarrow CO_2 + H_2O + 2e^-$

 $CH_2O + O^-_{(ads)} \rightarrow CH_2O_2 + e^-$

Zhang *et al.* synthesized NiO doped SnO₂nanosphereand utilized the material for better sensing for CH₂Oat different temperature range 50° C to 400° C than neat SnO₂ and NiO. The selectivity of the material NiO doped SnO₂ was again compared with ethanol, acetone, methylbenzene, chloroform and 2cholorophenol. It was seen that 50 ppm CH₂O showed enhanced sensing (26.03) at optimum temperature (100^oC). Chemisorbed oxygen plays an important role in sensing work. The mechanism behind VOC sensing using metal oxide nanomaterials at a particular temperature is basically based on the phenomenon of adsorption of oxygen from the surrounding atmosphere. This process involves a change in density of the conducting electrons on the surface of the sensing element as shown in scheme 1. This change is due to series of chemical reactions as follows:

(i) Pre-adsorption of oxygen on the surface of metal-oxide semiconducting nanomaterial.

(ii) Adsorbs the specific VOC on test.

(iii) Reaction takes place between oxygen and the adsorbed VOC.

(iv) Desorption of reacted VOC on surface.

These steps of reactions generate three types of stable oxygen anions namely O_2^- , O^- and O^{2-} during the electron delivery process that takes place between the VOCs and the surface of the metal-oxide semiconductor nanomaterial. During the reaction with VOC under test, a conductivity increase means the adsorbed oxygen plays the role of acceptor whereas a conductivity decrease means the adsorbed oxygen playing the role of a donor. Thus, the sensing mechanism of metal-oxide semiconductor gas nanomaterials for VOC detection is quite complex to explain and is based on the resistance change due to the chemical and electronic interaction between the test gas and the material. This complexity occurs due to the various influencing parameters like chemical composition, adsorption ability, surface properties, catalytic effect, temperature, humidity and modified surface morphology etc.

In case of SnO_2 nanosphere, only half-backed monolayer of oxygen was chemisorbed whereas in-case of NiO-SnO₂, doping of NiO on the surface of SnO_2 easily aggregate and chemisorbed the full monolayer oxygen ions and oxidised to higher oxidation state [18]. Kim and his group prepared multi-heterogeneous oxides NiO/Fe₂O₃ loaded SnO₂ for formaldehyde detection.

Catalytic hetero-junction effect plays a crucial role in high gas sensing. Apart from hetero-junction property, loading of catalyst and operating temperature also plays an important role in high sensing of formaldehyde. Herein, the establishment of multi hetero-junction between NiO, SnO_2 and Fe_2O_3 (p-n junction between NiO and SnO_2 and n-n hetero-junction between SnO_2 and Fe_2O_3). The migration of e's from NiO to SnO_2 and again to Fe_2O_3 decreased the EDL and increases the formaldehyde sensing of NiO/Fe₂O₃-SnO₂ composite [19].

Zhou and his team fabricated 3D hierarchical Co3O4 nanostructure by calcining Co5-based MOF microcrystals. As prepared Co_3O_4 -350 shows the best sensing towards formaldehyde at low temperature (170^oC), the low detection limit of 10 ppm and 30 day stability.

Here, 3D nanostructure, high surface area, more active active sites, high porosity, shorter ionic diffusion lengths made the material more effective for formaldehyde sensing [20].

3.2. Methane (CH₄).

Methane is one of the most significant constituents of natural gas. It shows hazardous because it considered a

greenhouse and also flammable. Therefore it is necessary to detect CH_4 at lower concentration for warning purposes when it leaks and also to prevent hazards. Nowadays a handful of works have been done by the scientists and researchers. Nasresfahani *et al.* was synthesized Pd-doped $SnO_2/PRGO$ and Pd-doped SnO_2/rGO and compared the methane sensing. The sensingperformance was evaluated from the concentration of methane gas which ranges between 800-16000 ppm at room temperature. In this regard Pd doped SnO_2/rGO shows the better result (9.8 % sensitivity) among all synthesized materials. This is because rGO nanosheets provide a favourable surface for nucleation and growth of nanoparticles and a conducting network that facilitate transfer of electron. High catalytic activities of Pd also play as an additive role which improves the methane sensitivity at a very low working temperature [21].

Microwave assisted spinel Co_3O_4 nanoparticles were fabricated by Shaalan and his team for methane sensing. Co_3O_4 is a p-type of semiconductor. As the holes are the major charge carriers in the p-type semiconductor, it dominates the passage of current through the boundary of semiconductor. Thus, there is a decrease in resistance occurs due to the formation of oxygen ions on the surface of the material by the trapping electrons. Again, during the mixing of CH₄ and O₂, the electrons come back to the material and increase the resistance because of electron-hole recombination [22].

3.3.Ethanol.

Ethanol is a reducing gas that used as the raw material for colors, antifreeze and formaldehyde. Direct methanol fuel cells (DMFC) are eco-friendly and considered as the important alternative fuel for automotive manufactures. However, 10 mL intake of methanol may cause total blindness and more than 30 mL, which may cause fatal diseases. So, the early detection of low concentration of methanol at low operating temperature is highly necessary.

Yang *et al.* adopted the hydrothermal method to synthesize SnO_2/Zn_2SnO_4 porous nanosphere for excellent ethanol gas sensing. The two main responsible factor for ethanol sensing are: i) porous structure and ii) the hetero-junction attained by SnO_2 and Zn_2SnO_4 . High surface area provides a large active sites which are essential for gas adsorption and diffusion. More is the surface active sites more is the adsorption between the adsorbed oxygen and methanol and more is the sensing. The hetero-junction play a significant role in sensing performance because of different band gap energy and work function of SnO_2 and Zn_2SnO_4 . Herein the conduction band edge potential of Zn_2SnO_4 is higher than the conduction band edge potential of SnO_2 [23].

Cheng *et al.* were synthesized pure and Pd-doped α -Fe₂O₃ porous nanotubes (PNTs) by a facile electrospinning approach. the results reveal that, compared with pure α -Fe₂O₃, the 3.0 wt% Pd-doped α -Fe₂O₃ PNTs sensors exhibit superior performance in ethanol detection (at 240 °C). The highest sensitivity reaches 65.4 for 50 ppm ethanol and the detection limit is as low as 0.1 ppm. It is excellent selectivity, and good sensor towards ethanol [24].

3.4. Methanol.

Methanol is an essential substance in daily life and in the industry. It is also considered as a raw material for colors, antifreeze and formaldehyde. Though it has advantages, the dietary intake of 10 mL may result in total blindness and exceeds that quantity may cause fatal diseases. So, this is necessary to quick detect methanol gas of low concentration at low oprating temperature. Li and his group fabricated Sm_2O_3 doped ZnO/SmFeO₃ microsphere for the high performance in methanol sensing.

The composite show excellent behaviour toward very low concentration of methanol (5 ppm) sensing at very low operating temperature (195[°]C) with 24 sec recovery time. Herein two main factors are responsible for methanol sensing are high surface area and p-n junction between ZnO (p-type) and SmFeO₃ (n-type).

High surface area of $SmFeO_3$ (24 h) enhances the active sites which adsorb oxygen. This leads to releasing back of electrons to the semiconductor surface and decrease the conductivity. The reaction as follows:

$$CH_3OH + 3O^n \rightarrow CO_2 + H_2O + 3ne^{-1}$$

Secondly, the formation of p-n heterojuction between ZnO and SmFeO₃, the transfer of electron and hole takes place through the interface as long they balance the Fermi level. Methanol reacts with the adsorbed oxygen on the surface of ZnO, releasing electron back. $CH_3OH + O^- \rightarrow HCHO + H_2O + e^-$

$$HCHO + O^- \rightarrow CO_2 + H_2O + e^-$$

Again, on the surface of SmFeO₃, the hole react with the adsorbed oxygen

$$CH_3OH + h^+ + O^- \rightarrow HCHO + H_2O$$

HCHO + $h^+ + O^- \rightarrow CO_2 + H_2O + e^-$

That mean the two different type of metal oxide can easily reactive with oxidative and reductive gases leading to high methanol sensing [25].

3.5. Acetone.

Acetone is good for some synthetic fibres and it is water miscible, can be used in common cleaning purpose. Acetone poisoning occurs when there's more acetone in your body than your liver can break down. When exposed to the air, it quickly evaporates and remains highly flammable. Acetone is dangerous to use around an open flame.

He prolonged exposure to acetone i.e. more than 173 ppm is highly toxic and cause serious threat to human health. As per the literature survey, the excelled gas of a healthy man is less than 0.8 ppm and in case of diabetic person is more than 1.8 ppm. So, sensing acetone is the best way to diagnose as compared to other testing. Thus it is highly necessary to prepare a suitable material that can easily sense acetone at a lower concentration. Wu *et al.* synthesized $ZnFe_2O_4/rGO$ for acetone sensing (10 ppm) at the operating temperature ($150^{0}C-225^{0}C$).

When adsorbed oxygen on the surface of $ZnFe_2O_4$ surface capture electrons in the CB to form O2- and O-. This results in the increase in resistance because of electron depletion layer. When acetone is exposed to the surface of $ZnFe_2O_4$, O_2^- and O⁻ react acetone at the grain boundary of $ZnFe_2O_4$. At this time again electrons are released back to CB.

$$O_{2(gas)} \rightarrow O_{2(ads)}$$

$$ZnFe_2O_4/rGO + O_{2 (ads)} + e^- \rightarrow O_2^-$$

$$ZnFe_2O_4/rGO + O_2^- + e^- \rightarrow 2O^-$$

$$CH_3COCH_3 + 8O^- \rightarrow CO_2 + H_2O + 8e^-$$

Again, the enhancement of sensing of acetone occurred due to 2D-rGO nanosheet by the improvement of electron mobility

inside the composite, forming a heterojunction structure in the composite. [26].

Xu and his team fabricated hollow Pt functionalized tungsten oxide (W₁₈O₄₉) for acetone sensing. The combined effect of porous shell and properties of Pt enhances the sensing performances 20 times more than pure W₁₈O₄₉. The enhanced sensing properties mainly due to two factors, one is spreading of Ptnano particle over the surface of W18O49 and the second one is the lower work function of W₁₈O₄₉. The spillage of Pt onto the surface of W₁₈O₄₉ dissociates the O-H, CH, O-O= bond, so more oxygenated species are available on the surface of composite, which trap more electrons. More acetone molecule adsorbed on the surface, which reacts with the adsorbed oxygen. As the work function of W₁₈O₄₉ (4.6eV) is less than Pt (5.65 eV), the electron flows from W₁₈O₄₉to Pt nanoparticle till the Fermi level equalized, decreasing the electron depletion layer and simultaneously increasing the scottky barrier height at the interface of p-n junction $(W_{18}O_{49} \text{ and } PtO_x)$.

This results in increase the resistance and therby the sensing performance. Apart from these properties, the high surface are, hollow sphere provides a better active sites for gas diffusion and inhibits the excellent gas sensing [27].

3.6. Triethylamine (TEA).

TEA is a colourless volatile liquid widely used in chemical industries. It is also highly explosive. The maximum consumption limit of TEA is 10 ppm or less. Higher to this concentration level, it causes headache, lungs problem etc. Various traditional methods are used for TEA detection such as gel chromatography and gas tube detection. But these methods are tedious and also costly. To avoid these problems, sensing is the best way to detect very low concentration of TEA gas at the faster rate. In this regard, porous metal oxides are the best option to use as the sensing material. Highly effective SnO_2 hallow microfiber was prepared by Zhou and his co-workers for TEA sensing.

Sensing mechanism of hollow SnO₂microfibres is mainly due to the presence of oxidative radicals $(O_2^-, O^-, and O^{2^-})$ and formation of electron depletion layer on the surface of the material. Other than that, unique structural characteristics such as the small grain boundary of SnO₂, large active sites, 1D carbon morphology and hollow porous structure contribute the outstanding response to TEA sensing having concentration 100 ppm at operating temperature 270°C [28]. Song et al. synthesized Au decorated porous α -Fe₂O₃nanorod for very low ppm level detection of Triethyl amine sensor. As selectivity is the most important character for gas sensor, Au/a-Fe₂O₃ exhibit the highest response towards 50 ppm of TEA at 40° C as compared to neat α -Fe₂O₃. High response is mainly based on the bond energy. As bond energy of C-N (305 KJ/mol) is weakest among all other bonds, TEA is most unstable. For this reason the bond breaks easily lead to the high activity of TEA and response increases. When TEA is exposed on the surface of Au/ α -Fe₂O₃, there is an oxidation and reduction reaction occur between TEA and $O^{\delta-}$ asper the reaction:

 $N(C_2H_5)_3 + O^{\delta-}(O^{2-}, O^-, O_2^-) \rightarrow N_2 + H_2O + CO_2 + e^-$

In addition, the decoration of gold on the surface of α -Fe₂O₃ leads to extraction of more number of electrons from the conduction band of α -Fe₂O₃. High extent of chemisorbed oxygen adsorption capacity on the surface of Au/ α -Fe₂O₃ results in the high response by accelerating the reaction between the oxygen species and the reducing gas. This leads to decrease in recovery time of Au/ α -Fe₂O₃ sensors [29]. In addition, few other nano materials used in the sensing of various types ofVOCs at different concentratin and operating temperature are summerised in the Table 1.

SI.	Nanomaterials used	Type of	Concentration and operating	References
No.		VOCs	temperature	
1.	CdIn ₂ O ₄ Nano porous thin film	formaldehyde	100 ррт, 100 ⁰ С	30
2.	SnO ₂ hollow microspheres	formaldehyde	200 ppm, 300 ⁰ C	31
3.	Au nanoparticle modified ZnO	formaldehyde	10 ppb	32
4.	Co ₃ O ₄ derived from Co5-Based MOF	formaldehyde	10 ppm, 175 ⁰ C	33
	Microcrystals			
5.	ZnONanostructures	methane	200μL/L, 275 ⁰ C	34
6.	Ordered Mesoporous In ₂ O ₃	methane	600^{0} C	35
7.	SnO ₂ @rGOPANI	methane	100-10000 ppm, room	36
			temperature	
9.	Pd-doped α -Fe ₂ O ₃ porous nanotubes (PNTs)	ethanol	50 ppm, 240 ⁰ C	24
10.	TiO ₂ nanoflower	methanol	100 ppm, 200 ⁰ C	37
11.	2D nanosheet assembled	acetone	200 ppm, 370 ⁰ C	38
	pdZnOmicroflowers			
13.	NiO/ZnO hybrid	acetone	200 ppm, 260 ⁰ C	39
14.	ZnO/ZnFe ₂ O ₄ hollow spheres	acetone	50 pm, 280 ⁰ C	40
15.	ZnFe ₂ O ₄ -graphene quantumdots	acetone	10ppm, at room temperature	41
16.	1 wt% RGO-CdFe ₂ O ₄	Acetone	10 ppm, 270 ⁰ C	42
17.	SnO ₂ hollow microfiber	triethylamine	100ppm, 270 ⁰ C	43
18.	α - Fe ₂ O ₃ microrod	triethylamine	500ppm, 275 ⁰ C	44
19.	TiO ₂ /SnO ₂ nanosheet	triethylamine	50 ppm, 260 ⁰ C	45
20.	CuO sphere	triethylamine	100 ppm, 230 ^o C	46

Table 1. Different other materials utilized in the sensing of VOCs.

4. CONCLUSIONS

This paper addresses the detection and monitoring of harmful volatile organic compounds liberated mostly from household activities. Because of the adverse effects of these gaseous pollutants on human health and also on the surrounding environment, VOC sensors have been taken up for greater demand in environmental and also domestic monitoring. This paper is an attempt to study few well known VOC sensors based on metal oxide semiconductor nano materials.

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