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Ultrasonication assisted fabrication of a tungsten sulfide/tungstite heterostructure for ppb-level ammonia detection at room temperature†

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A heterostructure of $WS_2/WO_3 \cdot H_2O$ has been prepared by partial oxidation of WS_2 nanosheets by exposing bulk WS_2 micron powder to ultrasonic waves in a bath sonicator. The as-prepared nanomaterial was used as a sensing film in an interdigitated electrode-based gas detecting device. The device was found to be specific towards ammonia gas among a group oxidizing and reducing gases. In particular, a response of as high as 11.36–254.66% was recorded for ammonia concentrations of 50 ppb to 2 ppm with excellent repeatability and reproducibility at room temperature. The response time and recovery time of the device was found to be a few tens of seconds suggesting its practicability. A plausible mechanism based on different active sites present in the receptor film is proposed and a logical reason behind its specificity towards ammonia gas is also inferred based on the Lewis acidic centers on the nano-surfaces. Overall, this proposed nanomaterial has very high potential for practical use as a room temperature ammonia sensor.

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1. Introduction

Detection of ammonia gas at low concentrations is very important for quality life as this is the most abundant air pollution element contributed by industry, vehicles, and food practices. Apart from this, there are several places in industry processes, breath analysis and other clinical diagnosis where it is necessary to ascertain ammonia concentration with highest possible accuracy.^{1–3} Several nanomaterials including metal oxides, semiconducting polymers, carbon nanotubes *etc.* are well established gas sensing receptors films for chemiresistive device applications. With the rise in two-dimensional (2D) material research, several exciting graphene analogous nanomaterials have emerged as an alternative to these existing materials. These materials provide excellent surface-to-volume ratio because of the weak interlayer van-der Waals attraction which allow exfoliation of ultrathin layers with nearly all the atoms exposed to the toxic analyte molecules. This weak interlayer physical forces in these materials (*i.e.* transition metal dichalcogenides (TMDs), hexagonal boron nitride (h-BN) *etc.*) allow easy fabrication of vertical heterostructure of 2D materials by simply stacking one over other by physical or chemical techniques as the epitaxial constraints are relaxed for these materials.⁴ However, the relative orientation control of layers

with respect to each other is an important aspect to be considered for vertical heterostructure. The vertically stacked heterogeneous nanomaterials provides a unique opportunity to tune the surface properties *via* work function tuning. For example, the work function of graphene sheets can be tuned by stacking them on any TMD nanosheets.⁵ By this stacking, a dipole moment is created across the two planes due to the interlayer charge transfer which could be used for fine control of the chemical reactivity on the surface. The neat interfaces at the heterojunction provide much reliable device performance. Unfortunately, fabrication of lateral heterostructure (with two materials linked by covalent bonding) remains challenging as the lateral growth requires lateral lattice matching. However, Ling *et al.* demonstrated a modified chemical vapor deposition (CVD) technique to grow lateral heterostructure (or parallel stitching) of 2D materials with even large lattice mismatch.⁶ In this technique, they sowed aromatic molecules on the substrate as seed to facilitate growth of the two different layers parallelly. A similar approach has also been reported by Zhang *et al.* to fabricate lateral heterostructure of TMDs.⁷ Zhou *et al.* synthesized 2D lateral heterostructure of tungsten sulfide (WS_2)/tungstite ($WO_3 \cdot H_2O$) by supercritical carbon dioxide ($sc-CO_2$) assisted exfoliation of WS_2 and then partially converting them into tungsten oxide monohydrate ($WO_3 \cdot H_2O$) through oxidation of these sheets under high pressure.³²

In this work, we have synthesized $WO_3 \cdot H_2O$ lateral heterostructure by simply bath sonicating the bulk WS_2 micron powder dispersed in ethanol–water (EtOH) and allowed local heating by avoiding any ice addition to the bath. This heating allowed partial oxidation of nanosheets with some water molecule being trapped in the crystal resulting in lattice H_2O .

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Several material characterization tools including XRD, Raman, UV/Vis, SEM, TEM, AFM *etc.* were utilized to characterize the phase, morphology, and heterogeneous nature of this nanomaterial. This material was further utilized in sensing toxic gases. The device showed excellent selectivity towards ammonia gas at room temperature.

2. Experimental section

2.1. Materials synthesis and characterization

The bulk WS₂ micron powder (>99.9% pure (Sigma Aldrich)) was used as the precursor material. Pure ethanol was obtained from Merck while deionized (DI) water of resistivity 18.2 MΩ cm was obtained from a Millipore system. This bulk WS₂ micro-powder was dispersed in 40 ml EtOH (ethanol: 65% water: 35% v/v) with an initial concentration of 6 mg ml⁻¹. The solution was ultrasonicated in a low power bath sonicator (PC analytics average power = 120 watt and frequency = 50 kHz) similar to our previous report, except that no ice was added to the bath.⁸ In place of ice addition, the sonicator was switched off for 1 minute after running for every 15 minutes to save the instrument from overheating. This cycle was repeated for about 72 hours. The dispersion was allowed to settle down for next 24 hours following which, it was centrifuged at 2000 rpm and top one-third of solution was collected for further characterization and device fabrication.

The morphology of bulk WS₂ and as-obtained nanomaterial was characterized using field-emission scanning electron microscopy (FE-SEM, ULTRA 55, Carl Zeiss AG). Low magnification and high-resolution transmission electron microscope (Low-Mag. TEM, and HR-TEM), high-angle annular dark-field

scanning transmission electron microscopy (HAADF-STEM) and corresponding energy dispersive spectroscopy (EDS) mapping analyses were done using a FEI transmission electron microscope (Titan Themis 300 kV) with sample dropcasted on a copper (300 mesh) grid. A Bruker (Dimension Icon) atomic force microscope (AFM) operating in tapping mode was utilized to understand the thickness profile of the nanomaterial. X-ray diffraction (XRD, Rigaku SmartLab) pattern was used to compare the bulk and as-fabricated nanomaterial crystal structure. Raman data was collected using Horiba (LabRAM HR) high-resolution Raman spectrometer equipped with 532 nm laser. Shimadzu UV-3600 UV-VIS-NIR spectrophotometer was employed to collect ultraviolet-visible (UV/Vis) spectrum.

2.2. Fabrication of gas sensor and their sensing measurements

The interdigitated electrode (IDE) with 5 μm width and gap was fabricated by standard photolithography technique as reported in our previous report.⁹ In brief, 90 nm of platinum (Pt) was deposited over 10 nm titanium (Ti) for IDE fingers and contact pads. 2.5 micro-liter (μL) of the as-prepared nanomaterial was dropcasted manually on the IDE's active area and used as sensing device after drying at 100 °C for half-an-hour in ambient without any post-processing step. This device was probed inside a custom-built gas sensing set-up as reported earlier.¹⁰ Commercial synthetic air (standard composition: N₂ = 79%, and O₂ = 21% by volume) was used to dilute analytes to obtain required concentration. It was also used to attain baseline of the device. A biasing voltage of 1 V was provided to the device during sensing measurements.

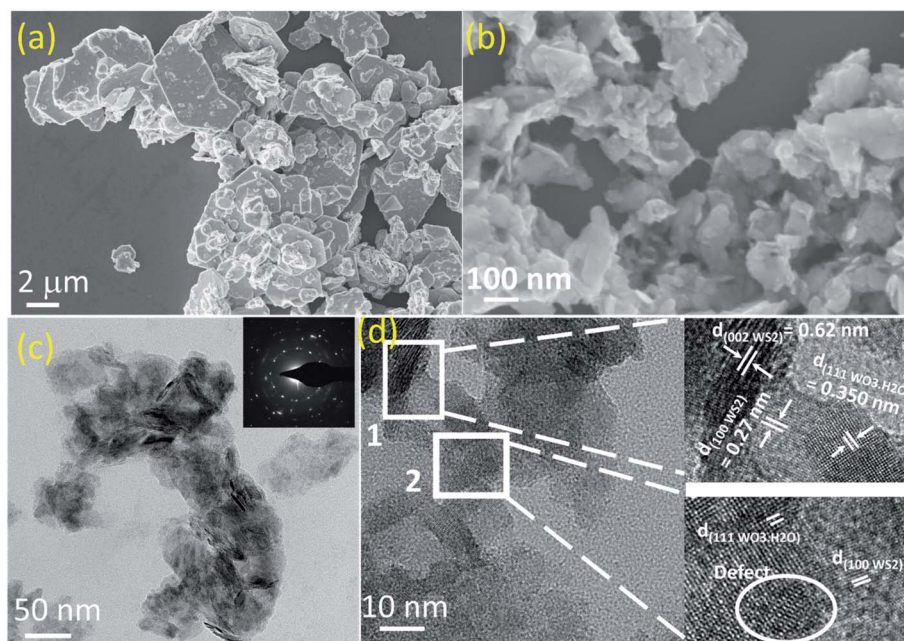


Fig. 1 FESEM image of (a) bulk WS₂ and (b) as-obtained nanomaterial after ultrasonication for 72 hours and centrifugation at 2000 rpm for 30 minutes; (c) low-mag. TEM image of the as-obtained nanomaterial (inset: SAED pattern) (d) HR-TEM image of the as obtained nanomaterial and zoomed image for two different area from this figure.



3. Results and discussion

3.1. WS₂/WO₃·H₂O synthesis

Bulk WS₂ powder (as shown in Fig. 1(a)) dispersed in EtOH was subjected to ultrasonic waves for long term (~72 hours). It should be noted that the bulk WS₂ was not able to disperse in pure water and thus, ethanol-water mixture was used as a convenient mixed solvent strategy.¹¹ Generally, to fabricate nanosheets of WS₂ (or any other TMDs) this method is taken into consideration and is called liquid phase exfoliation (LPE) as discussed in literature in detail.^{12–14} However, these experiments are performed in ice-cooling condition to avoid any oxidation which could take place due to local heat generation by strong ultrasonic waves.^{9,15} In this experiment, the bath sonicator chamber was not cooled (neither ice was added, nor any chiller was connected) and the intentional heat generation due to continuous bubble formation and collapsing due to ultrasonication was allowed. After long hours of sonication, the sample was allowed to settle down for 24 hours before collecting the supernatant of centrifuged sample. The FESEM image of the supernatant nanomaterial is shown in Fig. 1(b). It is evident that tiny nanoflakes have formed by the processing, which is also confirmed by low-mag. TEM images in Fig. 1(c). However, it also shows that many nanoflakes are oriented in vertical direction. It should be noted that we have done a thorough TEM measurements at various places and similar vertically oriented flakes were present along with the flat nanoflakes. This may be due to presence of different nanostructures with distinct surface energy in the liquid processed samples which require specific relaxation energy. The SAED pattern (inset of Fig. 1(c)) also confirmed the polycrystallinity in the sample. The HR-TEM image (as shown in Fig. 1(d)) of the sample clearly shows the heterogeneousness of the sample. The marked area 1 clearly shows the presence of lattice fringes due to (100) and (002) planes of WS₂ along with (111) plane of WO₃·H₂O. Area 2 also reveals these co-existing phases in the sample, along with some defects which may have been resulted from the long hours of sonication. However, both the areas clearly reveal lattice merging and a good interface at the lateral heterojunction. This neat interface is formed due to partial *in situ* oxidation of WS₂ nanosheets which has exfoliated during the sonication.

We further characterized the nanomaterial *via* different diffraction and spectroscopy techniques. The XRD pattern of bulk WS₂ powder and as-obtained nanomaterial is shown in Fig. 2(a). XRD peak corresponding to (002) peak for the bulk WS₂ sample (JCPDS card no. 08-0237) appears at a two-theta (2θ) value of 14.36° which shifts by 0.06° after nanostructuring towards higher 2θ values and appears at 14.42° in the as-obtained nanomaterial.

This is unusual in exfoliation as the nanostructuring usually results in shifting to lower 2θ value due to enhanced interplanar crystal spacing.^{16–18} Therefore, this shift is attributed to the planar tensile stress caused by the partial oxidation of nanosheets.¹⁹ Major peak corresponding to WO₃·H₂O appeared at 16.6° which is 0.1° offset (towards higher 2θ) value from the standard WO₃·H₂O XRD data (JCPDS card no. 43-0679) again

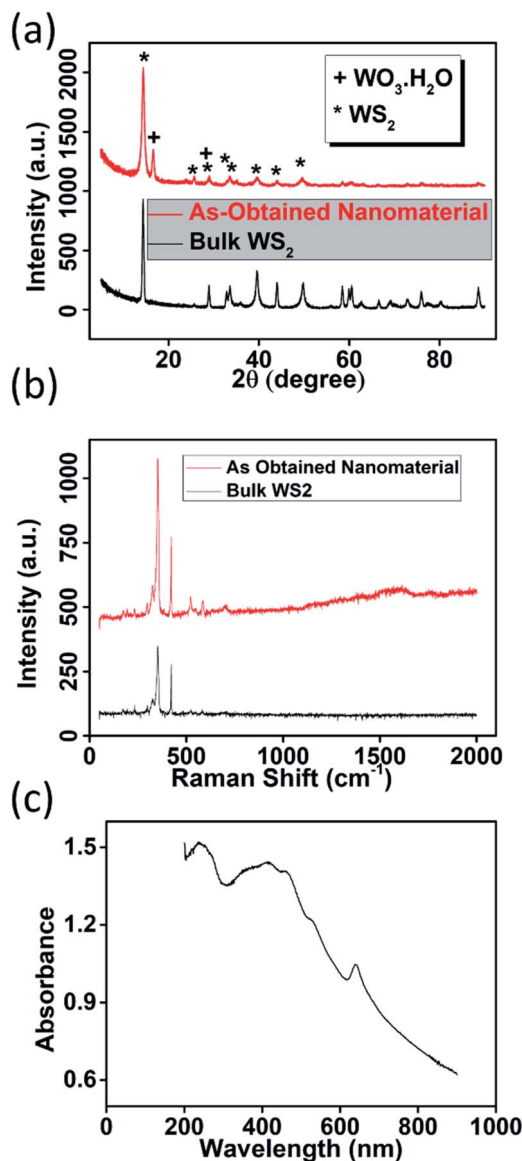


Fig. 2 (a) XRD pattern of bulk WS₂ and as-obtained nanomaterial (b) Raman spectroscopy data for bulk and as-obtained nanomaterial (c) UV/Vis spectra of as-obtained nanomaterial.

confirming the overall tensile stress in the nanomaterial.²⁰ The Raman data for the exfoliated nanomaterial is compared with the bulk WS₂ in Fig. 2(b).

The A_{1g} (out-of plane vibration) peak and E_{2g}¹ (in-plane vibration) peak of WS₂ still dominate in the exfoliated nanomaterial, confirming the partial oxidation of the nanosheets. The position of E_{2g}¹ peak in bulk WS₂ appears at 352.71 cm⁻¹ which remain intact in the exfoliated sample, however, A_{1g} peak in exfoliated nanomaterial appears to be blue shifted by 0.46 cm⁻¹ as compared to bulk sample. In the higher wavenumber regime, broad hump appears in the Raman signature of the as-obtained nanomaterial which is again due to partial oxidation of the WS₂ nanosheets during ultrasonication.²¹ The UV/Vis data for the exfoliated nanomaterial is shown in Fig. 2(c). The peaks at 239 nm and 406 nm correspond to the two

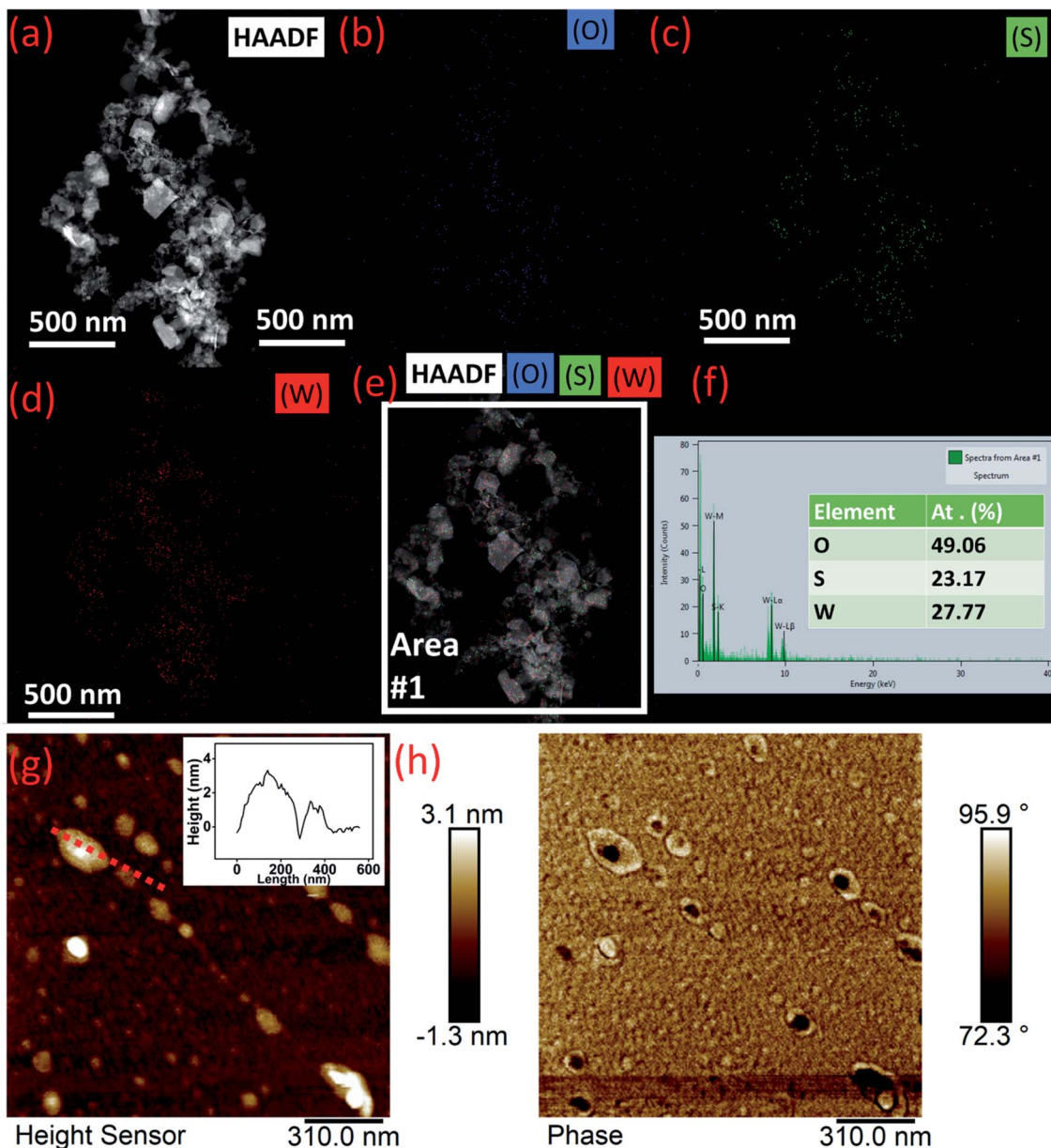


Fig. 3 (a) HAADF image of as-synthesized nanomaterial, elemental mapping in the sample corresponding to (b) oxygen (O), (c) sulfur (S), (d) tungsten (W), (e) elemental mapping imposed on HAADF image and (f) corresponding EDAX data (g) AFM image and its height-profile (inset) corresponding to a nanostructure and corresponding (h) AFM phase data.

unoccupied anti-bonding components of $\text{WO}_3 \cdot \text{H}_2\text{O}$, while peaks at 639 nm, 525 nm, and 459 nm correspond to characteristic excitonic A, B, and C band of WS_2 .^{22,23} HAADF-STEM image of the nanomaterial is shown in Fig. 3(a) with corresponding elemental mapping of elements in Fig. 3(b–d). These images clearly reveal the presence of oxygen in abundance in the sample. EDAX analysis of the corresponding area (Fig. 3(e) and (f)) clearly indicates the oxygen richness in sample due to partial oxidation. The AFM height profile (Fig. 3(g) and (h)) analysis of the samples showed ultrathin nature of the nanomaterial with thickness in the range of 1–4 nm and lateral dimension in the range 100–200 nm. The corresponding phase data of the AFM image shows nanomaterials with black spot in

the center with light yellowish shade at the outer part. This indicates that the WS_2 nanosheets are partially oxidized from edges towards center as depicted in schematic diagram in Fig. 4.

3.2. Gas sensing results

We utilized this nanomaterial as a sensing film in an IDE based chemiresistive device by simply dropcasting certain amount of the nanomaterial on IDE fingers. The device at a biasing voltage of 1 V shows stable baseline current in the range of few micro-ampere (μA). Then the device was subjected to several oxidizing and reducing gases and the ‘specificity (S)’ of the device was compared as per the histogram in Fig. 5(a). It



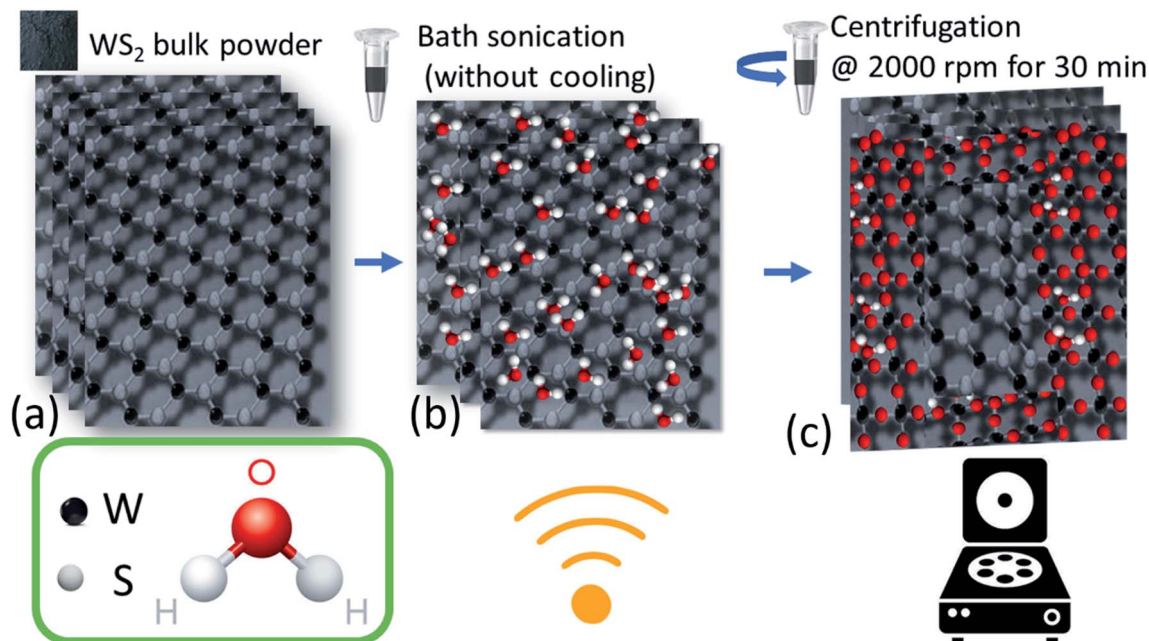


Fig. 4 Schematic of (a) bulk WS₂ powder existing in the form of layers (b) bath sonication in EtOH (only water molecules shown) without any additional effort to cool down the chamber and (c) sample centrifuged at 2000 rpm for 30 minutes.

should be noted here that the specificity (S_i) of a chemiresistive device is the measure of how selectively a sensor can identify a particular analyte 'i' among a set of 'n' gases. Mathematically, it can be calculated as per eqn (2), where, I_i is

the current in presence of analyte 'i' and I_a is the baseline current and numerical value of S_i ($0 \leq S_i \leq 1$) varies between zero to one (for $1 \leq j \leq n$).²⁴

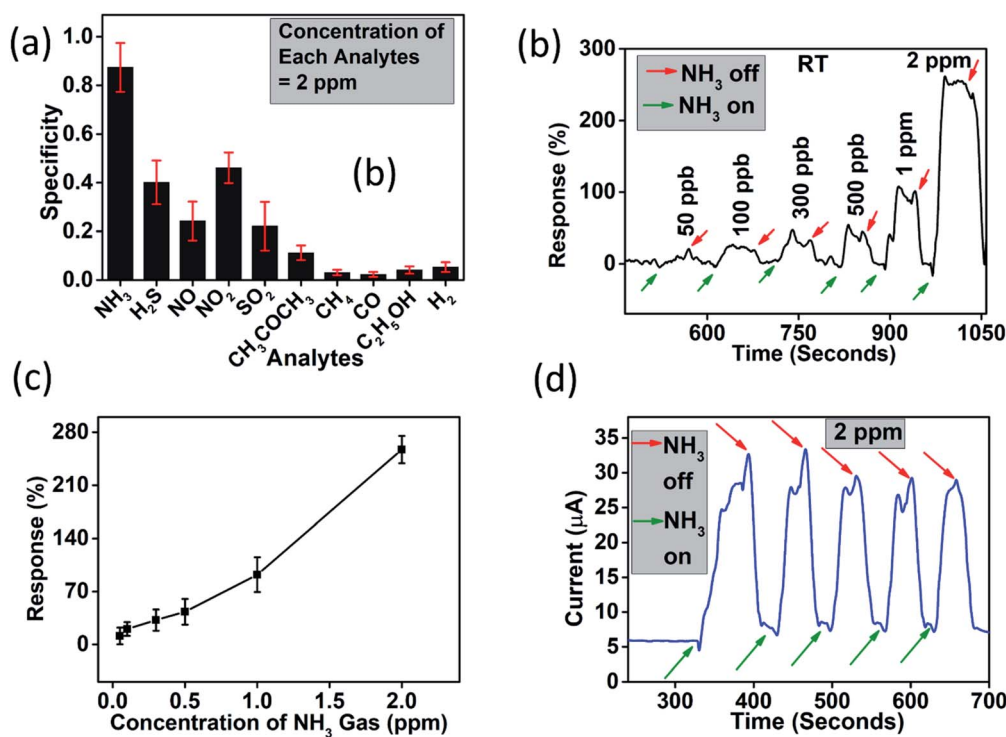


Fig. 5 (a) Specificity of the device for six different analytes for 1 ppm concentration (b) response (%) for ammonia concentration 50 ppb to 2 ppm at room temperature (c) response vs. concentration (ppm) plot with an exponential fit with equation $y = a/(1 + e^{-k(x - xc)})$ (d) repeatability of 2 ppm ammonia concentration for five consecutive on-off cycles.

$$S_i = \frac{\left(\frac{I_i}{I_a} - 1\right)}{\sum_{j=1}^n \left(\frac{I_j}{I_a} - 1\right)} \quad (1)$$

It is clearly evident from Fig. 6(a) that among ten ($n = 10$) different analyte (NH_3 , H_2S , NO , NO_2 , SO_2 , CH_3COCH_3 (acetone), CH_4 (methane), CO , $\text{C}_2\text{H}_5\text{OH}$ (ethyl alcohol), and H_2), the device has highest specificity towards ammonia gas.

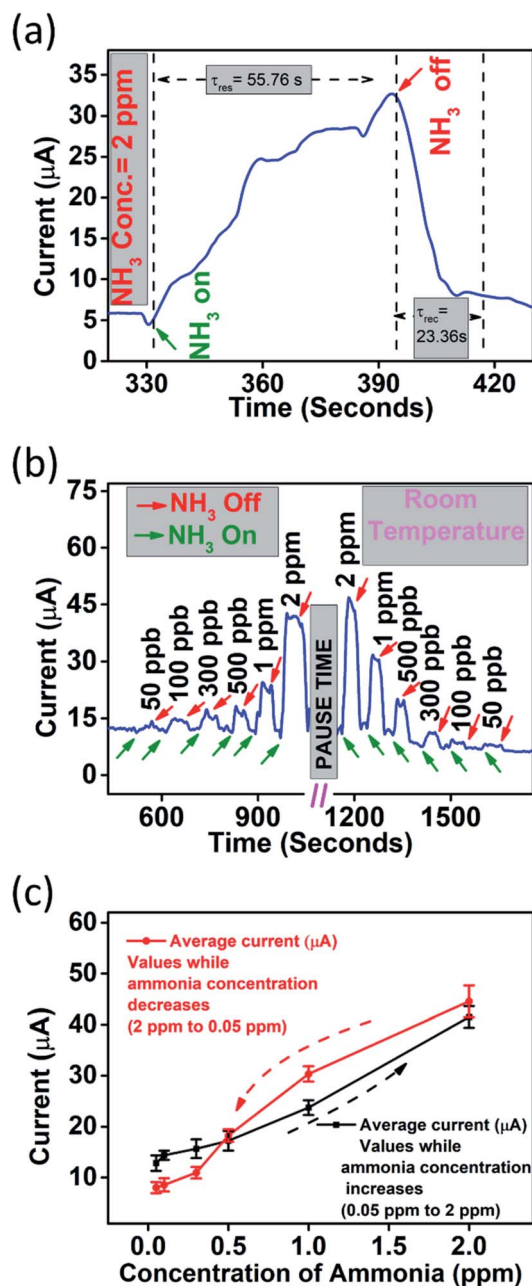


Fig. 6 (a) Extraction of response time (τ_{res}) and recovery time (τ_{rec}) for 2 ppm ammonia concentration (b) variation in current over time when exposed to ammonia gas from 50 ppb to 2 ppm and back to 50 ppb in synthetic air and (c) corresponding hysteresis.

Therefore, the device was subjected to different concentration of this analyte. Fig. 5(b) shows a dynamic response curve for when the device was subjected to ammonia concentration in the range 50 ppb to 2 ppm. The response (%) of the device was found to be 11.36–254.66% in this concentration range. It should be noted here that the response percentage of the sensor is calculated as

$$\text{Response}(\%) = \frac{I_g - I_a}{I_a} \times 100 \quad (2)$$

where I_g is the current of the device in presence of analyte and I_a is the baseline current in presence of synthetic air. The static curve for the response showing the device to device variation in three different devices is shown in Fig. 5(c). We went further to see the repeatability of the result when the device was subjected to same concentration. Fig. 5(d) shows the repeatability of the device when the device was subjected to five cycles of 2 ppm of ammonia. The device showed highly correlated results in all five cycles. The response time and recovery time of the device was found to be in the range of tens of seconds which indicates its practical usability (as shown in Fig. 6(a)).

Hysteresis is an important issue in chemiresistive gas sensing method. We investigated hysteresis error associated with the device. Hysteresis error (%) is calculated as

$$\gamma H(\%) = \pm \frac{\Delta I_{max}}{2F_{FS}} \times 100 \quad (3)$$

where, ΔI_{max} is the difference in current output during gas purging and gas desiccation while FFS is the full-scale output (*i.e.* the algebraic difference between the maximum current measured with maximum input stimulus and minimum input stimulus applied). The dynamic current change and corresponding memory effect is shown in Fig. 7(c) and (d) respectively. The average hysteresis error (%) when device was operated at room temperature is calculated to be 6.17%. We also investigated the long-term stability of the device by subjecting the same device (stored in desiccator) to various ammonia concentration after a month. The dynamic response of the device for 50 ppb to 2 ppm and back to 50 ppb of ammonia is shown in Fig. S1 (ESI).† The average hysteresis of the device has gone up by 4.21%. This increment may be due to the overall drift in baseline current. However, the current of the device and its response towards various concentration is preserved with average variation in observed response (after a month) is 3.84%.

We compared our result to some of the recent reported WS_2 based ammonia gas sensors as shown in Table 1. Although, most of the reported WS_2 and its hybrids/nanocomposites could detect ammonia at room temperature, but detecting below 1 ppm remains a challenge for them. $\text{WS}_2/\text{WO}_3 \cdot \text{H}_2\text{O}$ based device could detect much lower than this threshold at the same temperature. Also, the response and recovery time of our device is comparable to the reported results.

We also compared $\text{WS}_2/\text{WO}_3 \cdot \text{H}_2\text{O}$ based ammonia sensor with other nanomaterials-based room temperature chemiresistive NH_3 detectors (Table S1, (ESI)†) and found that our result is comparable to those.



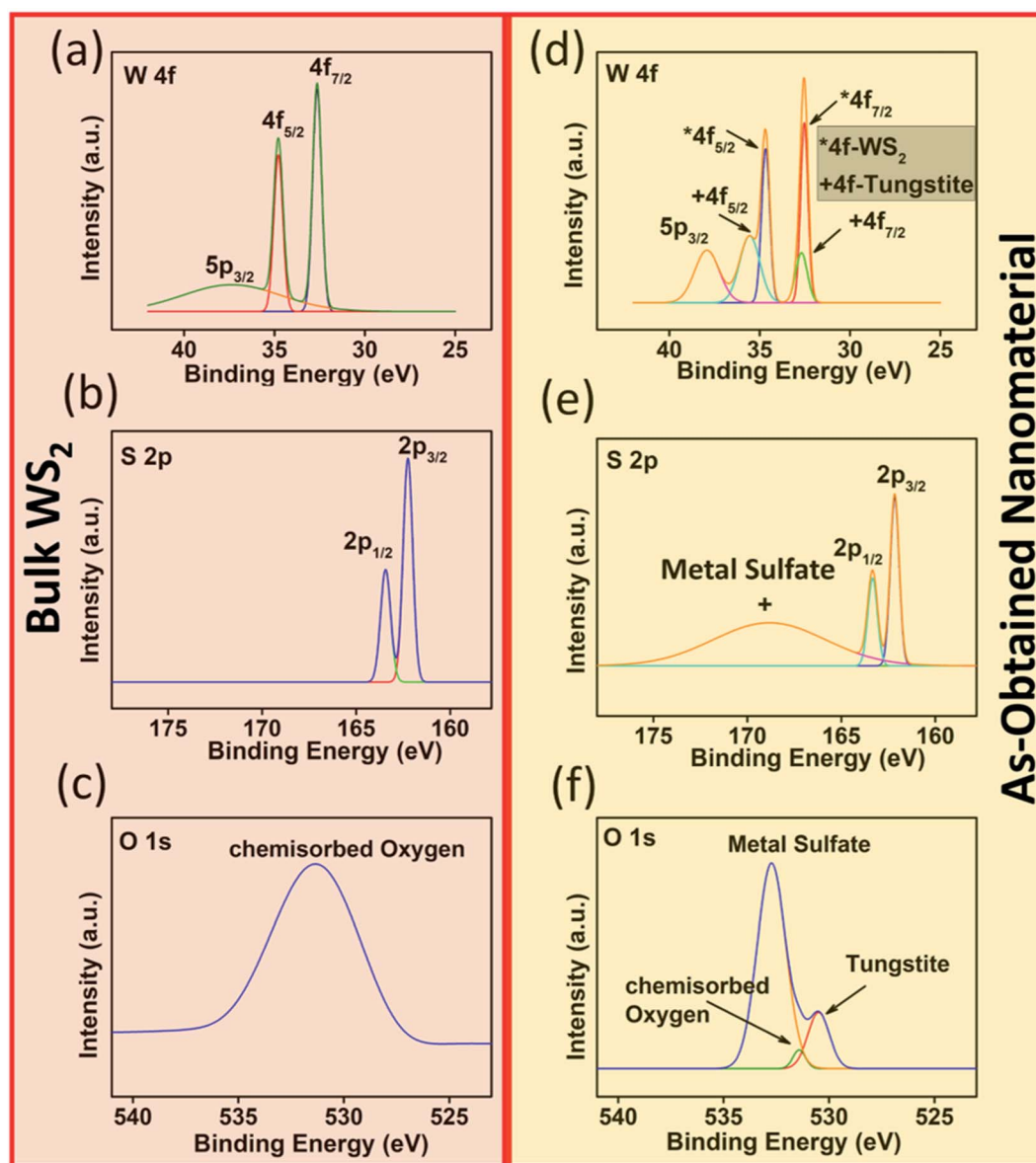


Fig. 7 (a) Tungsten (W 4f), (b) sulfur (S 2p), (c) oxygen (O 1s) XPS spectra of bulk WS_2 and (d) tungsten (W 4f), (e) sulfur (S 2p), (f) oxygen (O 1s) XPS spectra of as-obtained nanomaterial.

Table 1 Comparison of our $\text{WS}_2/\text{WO}_3 \cdot \text{H}_2\text{O}$ based ammonia gas sensor with recently reported chemiresistive sensing devices for NH_3 detection

| Material | Measured ammonia range (in ppm) | Response (%) | Response time/recovery time (seconds) | Synthesis technique | Operating temp. | Ref. |
|--|---------------------------------|-----------------|---------------------------------------|-------------------------------------|-----------------|-----------|
| WS_2 | 1–10 | ~225 @ 1 ppm | 120/150 @ 5 ppm | Ball milling of bulk WS_2 | RT | 25 |
| WS_2 | 1–5 | ~0.22 @ 5 ppm | ~300/600 @ 5 ppm | Plasma assisted synthesis at 500 °C | RT | 26 |
| WS_2 | 0.05–3 | 4.3% @ 0.05 ppm | 147/14 @ 3 ppm | Liquid exfoliation | 250 °C | 27 |
| WS_2/TiO_2 | 20–500 | 43.72 @ 250 ppm | –/174.43 @ 30 ppm | Chemical synthesis | RT | 28 |
| WS_2/RGO | 10–50 | 121 @ 10 ppm | 60/300 @ 30 ppm | Hydrothermal synthesis | RT | 29 |
| WS_2/Pt | 50–750 | 3 @ 50 ppm | –/– | Hydrothermal synthesis | RT | 30 |
| WS_2/WO_3 | 250–2000 | 150 @ 200 ppm | 160/310 @ 1000 ppm | Liquid processing method | 250 °C | 31 |
| $\text{WS}_2/$ | 0.05–2 | 11.36 @ 50 ppb | 55.76/23.36 @ 2 ppm | Liquid exfoliation method | | This work |
| $\text{WO}_3 \cdot \text{H}_2\text{O}$ | | | | | | |

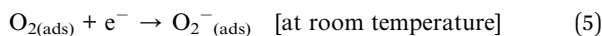


3.3. Plausible mechanism

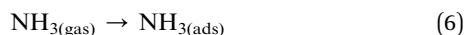
As it is predicted in the Fig. 4 as well as from experimental and theoretical predictions in earlier reports, the oxidation of the sheets happens from the edges, and hence, the edges of the nanosheets are considered to be tungstite for simplicity.³² Therefore, ideally there are three obvious sites where, the NH₃ molecules could interact with the receptor film, they are: (i) at the tungsten sulfide surface (ii) at tungstite surface and (iii) at their interface.

At the first interaction site, where NH₃ molecule interact with WS₂ surface, it is based on simple charge transfer mechanism as evident from the theoretical and experimental reports in literature.³³ NH₃ being a reducing gas, when it interacts with intrinsically n-type WS₂ nano-surface, it donates electron to it. These electrons populate the majority carrier in WS₂ which ultimately results in enhancement of current in presence of NH₃ molecule as evident from the gas sensing results (as shown in Fig. 6(b)).

Another active site is the WO₃·H₂O nano-surface, which interacts with any analyte at its surface through ionized oxygen (present due to adsorption from the ambient).³⁴ According to Wolkentein's model, the electron from the conduction band of WO₃·H₂O (which is a n-type semiconductor) is trapped by adsorbed surface O₂ molecules and form a metastable oxygen ion (O₂⁻_(ads)) as per eqn (4) and (5).



When NH₃ molecule interacts with these oxygen ions the following series of reactions occur as



The electrons released in eqn (7) are released back to the conduction band of tungstite which leads to the increase in the density of electrons which results in increase in current in the sensor film.

The third possible active site is the interface of tungsten sulfide/tungstite. Due to the formation of depletion region at the n-n lateral heterojunction of the nanomaterial, the overall resistance of the material goes high. However, when the material is exposed to reducing gas like NH₃, it transfers electron to the interface resulting in lowering the barrier potential and hence overall resistance.

Thus, interaction of NH₃ molecule at all available active sites in the receptor film results in increase in current which is reflected in the sensing data. To understand the selectivity of WS₂/WO₃·H₂O towards ammonia gas, we did chemical analysis of the sample using XPS as shown in Fig. 7. The tungsten (W 4f) XPS spectrum of exfoliated WS₂ shows additional peaks corresponding to tungstite while the sulfur (S 2p) spectrum shows additional peaks at higher binding energy corresponding to surface sulfate (SO₄²⁻) which could form in the process of

oxidation. The oxygen (O 1s) spectrum of the bulk WS₂ shows a single peak corresponding to chemisorbed oxygen while the exfoliated sample has additional peaks due to presence of sulfate as well as tungstite. The presence of SO₄²⁻ at the surface of nanomaterial, make it enriched in Lewis acid.²⁸ These Lewis acid centers make it specific to the Lewis base gases like ammonia.³⁵

4. Conclusions

To conclude, we have prepared WS₂/WO₃·H₂O lateral heterostructure nanomaterial by ultrasonication, by simply altering the sonication condition and subsequent centrifugation. This nanomaterial was used as a receptor film to detect NH₃ gas with high selectivity, in the concentration range of 50 ppb to 2 ppm. The response time and recovery time of the device was found to be of the order of tens of seconds. The repeatability, stability, and reproducibility of the device was found to be very good and comparable to the latest reports. We also proposed a sensing mechanism based on interaction of gas molecules at three different sites namely, the tungsten sulfide surface, tungstite surface and their interface. The proposed model is capable of explaining the sensing data, in the backdrop of material characterization results. This work will open a new possibility towards the fabrication of new heterostructure material based on metal sulfide/metal oxide for functional applications such as gas sensors.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- X. Xiao, X.-F. Cheng, X. Hou, J.-H. He, Q.-F. Xu, H. Li, N.-J. Li, D.-Y. Chen and J.-M. Lu, *Small*, 2017, **13**, 1602190.
- B. R. Varju, S. A. Wollschlaeger and D. B. Leznoff, *Chem.-Eur. J.*, 2019, **25**, 9017–9025.



- 3 E. Rafatmah and B. Hemmateenejad, *Electroanalysis*, 2019, **31**, 632–638.
- 4 K. S. Novoselov, A. Mishchenko, A. Carvalho and A. H. Castro Neto, *Science*, 2016, **353**, aac9439.
- 5 D. Deng, K. S. Novoselov, Q. Fu, N. Zheng, Z. Tian and X. Bao, *Nat. Nanotechnol.*, 2016, **11**, 218–230.
- 6 X. Ling, Y. Lin, Q. Ma, Z. Wang, Y. Song, L. Yu, S. Huang, W. Fang, X. Zhang, A. L. Hsu, Y. Bie, Y.-H. Lee, Y. Zhu, L. Wu, J. Li, P. Jarillo-Herrero, M. Dresselhaus, T. Palacios and J. Kong, *Adv. Mater.*, 2016, **28**, 2322–2329.
- 7 X. Q. Zhang, C. H. Lin, Y. W. Tseng, K. H. Huang and Y. H. Lee, *Nano Lett.*, 2015, **15**, 410–415.
- 8 N. Sakhuja, R. Jha and N. Bhat, in *Proceedings of IEEE Sensors*, Institute of Electrical and Electronics Engineers Inc., 2018 October, vol. 2018.
- 9 R. K. Jha, J. V. D'Costa, N. Sakhuja and N. Bhat, *Sens. Actuators, B*, 2019, **297**, 126687.
- 10 R. K. Jha, V. Singh, J. Sinha, S. Avasthi and N. Bhat, *IEEE Sens. J.*, 2019, **19**, 11759–11766.
- 11 K.-G. Zhou, N.-N. Mao, H.-X. Wang, Y. Peng and H.-L. Zhang, *Angew. Chem.*, 2011, **123**, 11031–11034.
- 12 Y. Lin, B. Adilbekova, Y. Firdaus, E. Yengel, H. Faber, M. Sajjad, X. Zheng, E. Yarali, A. Seitkhan, O. M. Bakr, A. El-Labban, U. Schwingenschlögl, V. Tung, I. McCulloch, F. Laquai and T. D. Anthopoulos, *Adv. Mater.*, 2019, **31**, 1902965.
- 13 V. Nicolosi, M. Chhowalla, M. G. Kanatzidis, M. S. Strano and J. N. Coleman, *Science*, 2013, **340**(6139), 1226419.
- 14 J. N. Coleman, M. Lotya, A. O'Neill, S. D. Bergin, P. J. King, U. Khan, K. Young, A. Gaucher, S. De, R. J. Smith, I. V. Shvets, S. K. Arora, G. Stanton, H. Y. Kim, K. Lee, G. T. Kim, G. S. Duesberg, T. Hallam, J. J. Boland, J. J. Wang, J. F. Donegan, J. C. Grunlan, G. Moriarty, A. Shmeliov, R. J. Nicholls, J. M. Perkins, E. M. Grievson, K. Theuvsen, D. W. McComb, P. D. Nellist and V. Nicolosi, *Science*, 2011, **331**, 568–571.
- 15 C. Backes, B. M. Szydłowska, A. Harvey, S. Yuan, V. Vega-Mayoral, B. R. Davies, P. L. Zhao, D. Hanlon, E. J. G. Santos, M. I. Katsnelson, W. J. Blau, C. Gadermaier and J. N. Coleman, *ACS Nano*, 2016, **10**, 1589–1601.
- 16 G. Q. Han, Y. R. Liu, W. H. Hu, B. Dong, X. Li, Y. M. Chai, Y. Q. Liu and C. G. Liu, *Mater. Chem. Phys.*, 2015, **167**, 271–277.
- 17 R. K. Jha, D. Burman, S. Santra and P. K. Guha, *IEEE Sens. J.*, 2017, **17**, 7340–7347.
- 18 R. K. Jha and P. K. Guha, *Nanotechnology*, 2016, **27**, 475503.
- 19 B. D. Cullity and S. R. Stock, *Elements of X-ray diffraction*, Edison-Wesley Publ. Co. Inc, USA, 2nd edn.
- 20 L. Chen, T. Mashimo, H. Okudera, C. Iwamoto and E. Omurzak, *RSC Adv.*, 2014, **4**, 28673–28677.
- 21 A. G. Kelly, V. Vega-Mayoral, J. B. Boland and J. N. Coleman, *2D Mater.*, 2019, **6**, 45036.
- 22 M. Ahmadi and M. J. F. Guinel, *Acta Mater.*, 2014, **69**, 203–209.
- 23 G. Pagona, C. Bittencourt, R. Arenal and N. Tagmatarchis, *Chem. Commun.*, 2015, **51**, 12950–12953.
- 24 R. K. Jha, N. Sakhuja, S. Jakhar and N. Bhat, *IEEE Trans. Nanotechnol.*, 2019, **18**, 932–939.
- 25 X. Li, X. Li, Z. Li, J. Wang and J. Zhang, *Sens. Actuators, B*, 2017, **240**, 273–277.
- 26 M. O'Brien, K. Lee, R. Morrish, N. C. Berner, N. McEvoy, C. A. Wolden and G. S. Duesberg, *Chem. Phys. Lett.*, 2014, **615**, 6–10.
- 27 N. Sakhuja, R. K. Jha and N. Bhat, *IEEE Sens. J.*, 2019, **19**, 11767–11774.
- 28 Z. Qin, C. Ouyang, J. Zhang, L. Wan, S. Wang, C. Xie and D. Zeng, *Sens. Actuators, B*, 2017, **253**, 1034–1042.
- 29 X. Wang, D. Gu, X. Li, S. Lin, S. Zhao, M. N. Rumyantseva and A. M. Gaskov, *Sens. Actuators, B*, 2019, **282**, 290–299.
- 30 C. Ouyang, Y. Chen, Z. Qin, D. Zeng, J. Zhang, H. Wang and C. Xie, *Appl. Surf. Sci.*, 2018, **455**, 45–52.
- 31 R. K. Jha, M. Wan, C. Jacob and P. K. Guha, *IEEE Sens. J.*, 2018, **18**, 3494–3501.
- 32 P. Zhou, Q. Xu, H. Li, Y. Wang, B. Yan, Y. Zhou, J. Chen, J. Zhang and K. Wang, *Angew. Chem., Int. Ed.*, 2015, **54**, 15226–15230.
- 33 C. Zhou, W. Yang and H. Zhu, *J. Chem. Phys.*, 2015, **142**, 214704.
- 34 F. Li, S. Guo, J. Shen, L. Shen, D. Sun, B. Wang, Y. Chen and S. Ruan, *Sens. Actuators, B*, 2017, **238**, 364–373.
- 35 S. Kabcum, N. Kotchasak, D. Channei, A. Tuantranont, A. Wisitsoraat, S. Phanichphant and C. Liewhiran, *Sens. Actuators, B*, 2017, **252**, 523–536.

