Application of graphene oxide based Microfiber-Knot resonator for relative humidity sensing

S.R. Azzuhri\textsuperscript{a}, I.S. Amiri\textsuperscript{b,c,e,}, A.S. Zulkhairi\textsuperscript{d}, M.A.M. Salim\textsuperscript{d,e}, M.Z.A. Razak\textsuperscript{f}, M.F. Khyasudeen\textsuperscript{g}, H. Ahmad\textsuperscript{d}, R. Zakaria\textsuperscript{d}, P. Yupapin\textsuperscript{b}

\textsuperscript{a} Department of Computer System & Technology, Faculty of Computer Science & Information Technology, University of Malaya (UM), Malaysia
\textsuperscript{b} Computational Optics Research Group, Advanced Institute of Materials Science, Ton Duc Thang University, Ho Chi Minh City, Viet Nam
\textsuperscript{c} Faculty of Applied Sciences, Ton Duc Thang University, Ho Chi Minh City, Viet Nam
\textsuperscript{d} Photonics Research Centre, University of Malaya (UM), Malaysia
\textsuperscript{e} Laser Centre, Ibnu Sina Institute for Scientific & Industrial Research, Universiti Teknologi Malaysia (UTM), Malaysia
\textsuperscript{f} Institute of Microengineering and Nanoelectronics (IMEN), Universiti Kebangsaan Malaysia (UKM), Malaysia
\textsuperscript{g} Faculty of Electrical Engineering, Universiti Teknologi MARA (UITM), Malaysia

\textbf{ARTICLE INFO}

Keywords:
Microfiber
Resonator
Humidity sensor

\textbf{ABSTRACT}

A relative humidity (RH) sensor is proposed and demonstrated using a micro-knot resonator (MKR) enhanced with a layer graphene oxide (GO) coating. The MKR is fabricated by means of tapering a standard fiber, with the GO coating added by the drop-cast method. The proposed sensor is tested for an RH range of between 0\% and 80\% at 20\% intervals, and the configurations with and without the GO coating achieve sensitivities of 0.0104 nm/\% and 0.0095 nm/\%, respectively. The MKR configuration without the GO coating has a linear response correlation coefficient of 0.9908 and a resolution of 0.1\%, while the configuration with the GO coating has a linear response correlation coefficient of 0.9548 and a resolution of 0.096\% which is better. The proposed sensor has multiple applications, especially in the area of climate and atmospheric measurement and monitoring.

\textbf{Introduction}

Since its successful isolation from graphite in 2004 by mechanical exfoliation \cite{1}, graphene has become key material in realizing a multitude of new and novel photonics applications. These new abilities stem from the unique optical and electrical characteristics of graphene, such as the ability to exhibit the quantum Hall effect at room temperature \cite{2-4}, an ambipolar field effect and new optical properties \cite{5-8}. While graphene and its derivatives such as graphene oxide (GO) initially made in-roads in the field of communications, there has also been substantial interest in sensing applications \cite{9,10}. The extremely large surface area made possible by its 2-D structure is advantageous towards making these materials highly sensitive to changes in the surrounding environment. Furthermore, these materials can be easily produced on a large scale at low cost, making them even more advantageous for use in the sensing industry \cite{11-18}. GO in particular possess certain advantages over conventional graphene for sensing applications, as it retains its high sensitivity but is easier and less costly to fabricate. This is primarily due to the inclusion of oxygen functional groups such as carboxyl, hydroxyl and epoxy groups into the hexagonal crystalline structure of graphene, making it robust and easier to manage \cite{19}. This was exemplified by the work of Dikin et. al. who created GO paper with remarkable mechanical properties using stacks of single GO sheets \cite{20}. Furthermore, graphene oxide can easily be reduced in solution and as a thin film, which does not require an expensive equipment \cite{21}. GO also exhibits a strongly anisotropic complex dielectric function, which is beneficial to photonics waveguide based polariser and sensor applications \cite{22-26}.

The application of graphene and GO as a sensor in optical systems has been well demonstrated \cite{27,28}. Gaston et. al. \cite{29} proposed and successfully demonstrated a humidity sensor using a side-polished fiber (SPF) with a polyvinyl-alcohol (PVA) overlay. Yao et al. \cite{30} on the other hand reported a GO silicon with a bi-layer adaptable structure as a stress-based humidity sensor which exhibited minimal hysteresis, with a quick response and recovery time. However, both these approaches had certain limitations; the GO silicon bi-layer adaptable structure based sensor required a working voltage of 5 V, complicating the operation and overall effectiveness of the sensor, while the SPF based sensor can be difficult to fabricate, requiring significant time and care, as well as having a limited exposure interaction length \cite{31}.

\textsuperscript{*} Corresponding author at: Ton Duc Thang University, Ho Chi Minh City, Viet Nam.

\textbf{E-mail address:} irajsadeghamiri@tdt.edu.vn (I.S. Amiri).


https://doi.org/10.1016/j.rinp.2018.05.009
Received 25 March 2018; Received in revised form 1 May 2018; Accepted 2 May 2018
Available online 05 May 2018
2211-3797/ © 2018 The Authors. Published by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/BY-NC-ND/4.0/).
An interesting alternative to this would be the micro-knot resonator (MKR), which can be a prime candidate in the development of a cost-effective, easy to fabricate and simple to use optical sensors for climate and environmental monitoring. The MKR is essentially a tapered fiber with a waist region of several hundred nanometers to several micrometers in diameter that is formed as a loop [32]. This configuration allows the evanescent field of a propagating signal to interact with the surroundings of the MKR. The MKR possesses a number of key advantages over similar narrow-waist fibers, in particular its excellent diameter uniformity and sidewall smoothness [8]. Yu Wu et al. have fabricated fiber-optic interferometric humidity sensors based on silica/polymer MKR. The silica microfibers used in this work were fabricated by flame-heated taper-drawing of a single-mode fiber. The authors have reported a large humidity sensitivity and ~88 pm/10% RH for the silica and polymer MKR, at the range from 20%-RH to 96%-RH, and 17%-RH to 98%-RH respectively [33]. Jong Cheol Shin et al. have fabricated the MKR with tie and loop diameters of 2 and 70 µm respectively with a polyvinyl alcohol overlay, which is highly sensitive to the concentration of RH. They have reported that the RH sensitivity of the proposed MKR-based RH sensor strongly depends on the mode order (m), which were measured to be $-0.22$, $-0.44$, $-0.65$, and $-0.87 \mu$m$^{-1}$, respectively, where the multiple interference fringes depending on the mode orders are superimposed in the transmission spectrum [34]. In reference [35], the authors have improved the few modes 4 µm MKR sensitivity by coating the device with a polyvinyl alcohol (PVA) which can absorb the humidity efficiently. The improvement could be obtained via optimizing the diameter of the device.

By combining the MKR with a sensitive 2-D material coating, the sensitivity and responsivity of the MKR based sensor can be increased substantially. In this work, an MKR sensor utilizing a GO coating for the measurement of relative humidity (RH) is proposed and demonstrated. Humidity sensing is an integral part of many activities, including the monitoring of structures and manufacturing processes [36]. While humidity sensing and measurement can be made through conventional electronic sensors, certain limitations, including high fabrication and operation cost as well as maintenance requirements and an inability to be deployed in corrosive and explosive environments which restricts its use [37]. In this regard, optical based sensors such as the MKR can be used to overcome these limitations, and therefore offer significant potential for the development of large-scale sensor grids for environmental and climate monitoring.

**Experimental setup**

The MKR used in this work is fabricated from a tapered fiber, which is obtained using the flame-heating technique on a single-mode fiber (SMF). The schematic of the flame-brush assembly is shown in Fig. 1(a), while Fig. 1(b) provides the image of the actual setup. The SMF is commercially available, with the center region stripped of its protective polymer coating and cleaned thoroughly with propanol. The SMF is then secured by fiber holders onto two translation stages on a motorized rail, with both translation stages allowed to move along the rail opposite to each other. The center region of the SMF is placed such that it lies directly above the mouth of the torch. As the SMF is heated by the flame, it becomes soft and stretchable. At this juncture, the SMF is slowly stretched by the two translation stages as they move in opposing directions, creating a narrow waist region. The stretching process is monitored by measuring the loss of power of an amplified spontaneous emission (ASE) signal travelling through the SMF. The ASE source is assembled from a 980 nm laser diode and a 12 m long erbium doped fiber with an absorption rate of 5.0 dB/m at 1531 nm. The ASE source is then connected to one end of the SMF, with other end connected to a Thorlabs PM100USB optical power meter (OPM).

Flame brushing technique is easy to setup, provide high accuracy and does not require an expensive equipment or tedious setting. It is found that the shape of the taper should follow the adiabaticity criteria and has a longer transition length to achieve low loss characteristic. As the flame heats and softens the SMF, the translation stages make micro-transitions of 0.25 mm in opposing directions, gently stretching the fiber before returning to the start position. This is done for 20 iterations, after which the process repeats, only this time with the stages stretching further to 50 mm for 40 iterations. The process repeats for a final time, with the stages moving 1.0 mm in each direction for 40 iterations. Fig. 2 shows the Graph of number of iterations versus SMF stretching, while Fig. 3 shows the schematic of the SMF before and after the stretching process.

It is important to note that in the fabrication of the fiber, the core diameter remains unchanged, and only the cladding stretches until it becomes so thin that the core is essentially exposed at the tapered waist region. The cladding diameter at the untapered regions remains unchanged, while the cladding in the transition region forming a slope. The region near the tapered waist and nearest the untapered region are seen to have diameters of 8.0 and 125.0 µm respectively, as shown in the figure. The tapered SMF structure allows the evanescent field that emits from the signal propagating through the core to interact with the surrounding environment, as the cladding region is now too narrow to confine the evanescent field. The evanescent field can now interact with...
its surrounding, and any change in the propagating signal can be related to the changing parameter in the environment. The MKR itself is created by simply looping the fabricated tapered fiber so that it forms a ring at the tapered region. It must be noted that although the tapered fiber is still structurally strong, significant care needs to be taken to ensure that the tapered fiber does not break. In this work, a ring with a diameter of 3.02 µm is obtained. The microscopic image capture of the MKR is given in Fig. 4. The untapered part of the MKR assembly is spliced to pigtails and can then be connected to conventional fiber components and devices.

The radius of tapered fiber can be approximated by the equation:

$$ r(x) = r_0 \exp \left( \frac{1}{2} \int \frac{dx}{L} \right) $$

(1)

where $r_0$ is the radius of the fiber core before tapering and $L$ is the hot zone length with a distance of $x$, where $x$ can be calculated from Eq. (2):

$$ L(x) = L_0 + \alpha x $$

(2)

where the $\alpha$ is constant which varies between $-1$ and $1$, and $L_0$ being the length of the fiber before tapering. The radius of the tapered can be obtained as in reference [38] as follows:

$$ r(x) = r_0 \left( 1 + \frac{\alpha x}{L_0} \right)^{-\frac{1}{\alpha}} $$

(3)

The GO nanoparticles used in this work are obtained commercially in the form of an aqueous solution. The Raman spectrum of the GO nanoparticles is given in Fig. 5. Three Raman peaks are obtained at 1353 cm$^{-1}$, 1605 cm$^{-1}$ and 2715 cm$^{-1}$, corresponding to the D, G and 2D peaks. These peaks verify that the sample consists of GO nanoparticles, with little to no impurities present. The ratio of $I_D$ to $I_G$, at approximately 1:4 indicates that the GO nanoparticles are stacked up in multiple layers [39,40].

The setup of the MKR as a humidity sensor is given in Fig. 6. Once formed, the MKR is left in place and the translation stages locked to prevent accidental movements that might damage the MKR. The ASE source is now replaced with a Yokogawa AQ2200-136 tunable light source, and the OPM is substituted with an Anritsu model MS9740A optical spectrum analyzer (OSA). A simple, homemade containment chamber is assembled around the MKR, and water vapor is introduced into the chamber by injecting nitrogen gas into a container containing water, which is then channelled into the chamber. The RH range is measured from an approximate value of 20% to 80% at 20% intervals using a hygrometer, which also acts as the reference source.

**Results and discussion**

The MKR fabricated in this work is tested as a humidity sensor with and without the GO coating. This is done to ascertain the performance of the MKR under both configurations, as well as to gauge the improvement in performance as a result of the GO layer. In order to ensure that as little as possible changes or modifications are made to the experiment setup, which could in turn affect the measurements obtained, the MKR is first tested without the GO coating. Subsequently, the GO coating is added by simply drop-casting the GO solution on the MKR, and the entire experiment repeated. This ensures that the two configurations are as similar as possible, and thus provide a more accurate comparison.
Fig. 7 shows the response of the MKR in both configurations against an increasing RH, from 20% to 80% at step-jumps of 20% each time. It can be seen from the figure that the extinction ratio of the transmission spectrum for the MKR with the GO coating increases from around $-41$ to $-27$ dBm as the RH value increases as compared to an increase of only 10 dB for the MKR with no coating. This is attributed to the GO layer exhibiting high levels of hygroscopity [41], thus as the RH increases more and more water molecules are absorbed by the GO matrix. This in turn causes the refractive index of the GO layer to increase, such that it can better confine the evanescent field of the test signal propagating through the core of the MKR. In the case of the MKR with no coating, the change in the refractive index of the air as a result of the rising RH is still detectable, but as this change is less distinct only a small shift is observed in the extinction ratio. This clearly indicates that the addition of the GO coating increases the MKR sensitivity significantly, and therefore enhancing its performance substantially. The transmission spectrum obtained when using the MKR with no coating has a Q-factor of approximately 6307 and a free spectral range (FSR) of about 0.24 nm, while the GO coated MKR has a transmission spectrum Q-factor and FSR approximately 10,365 and 0.156 nm respectively.

Figs. 8 and 9 provide the response of the resonance wavelength coating, the change in the refractive index of the air as a result of the rising RH is still detectable, but as this change is less distinct only a small shift is observed in the extinction ratio. This clearly indicates that the addition of the GO coating increases the MKR sensitivity significantly, and therefore enhancing its performance substantially. The transmission spectrum obtained when using the MKR with no coating has a Q-factor of approximately 6307 and a free spectral range (FSR) of about 0.24 nm, while the GO coated MKR has a transmission spectrum Q-factor and FSR approximately 10,365 and 0.156 nm respectively.

Figs. 8 and 9 provide the response of the resonance wavelength
towards the changing RH levels for the uncoated and GO coated MKR configurations respectively. From Fig. 8, it can be seen that the uncoated MKR has a regression coefficient of determination ($R^2$) of 0.9098, with a computed sensitivity of 0.0095 nm/%RH. On the other hand, the GO coated MKR has an $R^2$ value of 0.9548 and a sensitivity of 0.0104 nm/%RH, as shown in Fig. 9 the MKR sensor with the GO coating in this work demonstrates the ability to obtain measurements along a smaller range, which has comparable sensitivity value with others previously observed [42–46]. In this manner, the measurement ranges demonstrated by Zheng et al. and Alvarez et al. [47] for similar works are 51–78% RH and 0–15% RH, respectively.

According to the definition of humidity sensitivity of the sensor, the resolution can be calculated using following equation as [48]

$$R_{\text{sensor}} = \frac{R_{\text{wavelength}}}{\text{Sensitivity}}$$

(4)

where, the $R_{\text{sensor}}$ is the resolution of the MKR sensor, the $R_{\text{wavelength}}$ is the wavelengths resolution due to the RH variations. As it can be seen from Figs. 8 and 9, the wavelength shift of 0.1 nm observed for the 100% RH changes. Therefore, the wavelength resolution ($R_{\text{wavelength}}$) is 0.001 nm. Using the Eq. (4), the sensor resolutions ($R_{\text{sensor}}$) for the MKR without and with the GO coating are 0.1% and 0.096% respectively, where the MKR coated with GO shows a higher sensitivity and therefore a better resolution.

Fig. 10 shows the response of the optical power towards the change in RH value.

It can be seen from the figure that for the MKR without a GO coating, there is only a gradual change, with the power barely dropping from just above −30.5 dBm at a RH value of 0% to slightly less than −30.5 dBm as the RH value approaches 90%. On the other hand, the MKR coated with the GO nanoparticles experiences a drastic reduction in the power of the propagating signal, with the power dropping rapidly from approximately −30.7 dBm at an RH value of 0% to around −31.3 dBm at an RH value of 60%. After this, the signal power reduces at a slower rate as the RH increases, with the power dropping only slightly to −31.4 dBm as the RH value approaches 90%. This behavior is attributed to the GO particles becoming saturated with water molecules, and thus not able to host more molecules even as the RH increases. Table 1 summarizes the performance characteristic of the proposed MKR humidity sensor for both configurations.

Overall, it can be seen that the use of the GO layer on the MKR significantly increases its performance as a sensor, in particular with an increase in sensitivity and better linearity. The GO coated MKR sensor has many potential sensing applications, most notably in climate and atmospheric parameter measurement.

**Conclusion**

In this work, an MKR based optical sensor is used as a RH sensor and its performance compared in two different cases, namely with and without a GO coating. The MKR is fabricated using a tapered fiber which is then looped into a knot. The MKR is first tested without the GO coating, and subsequently, the GO coating is added by the drop-casting liquid technique. The MKR is capable of detecting changes in the RH levels and has a regression coefficient of determination and sensitivity of 0.9098 and 0.0095 nm/%RH respectively for the uncoated MKR as well as a regression coefficient of determination and sensitivity of 0.9548 and 0.0104 nm/%RH respectively for the coated case. The proposed MKR sensor would have many potential sensing applications, particularly in measuring climate and atmospheric related parameters.

**Acknowledgements**

This work was supported by the research grants LRGS (2015)/NGOD/UM/KPT, GA010-2014 (ULUNG) and BR002-2016.

**Appendix A. Supplementary data**

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.rinp.2018.05.009.

**References**


---

**Table 1**

<table>
<thead>
<tr>
<th>Performance parameter</th>
<th>Without GO layer</th>
<th>With GO layer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>0.0095 nm/%RH</td>
<td>0.0104 nm/%RH</td>
</tr>
<tr>
<td>Linear response correlation coefficient</td>
<td>90.98%</td>
<td>95.48%</td>
</tr>
<tr>
<td>Resolution (%)</td>
<td>0.1</td>
<td>0.096</td>
</tr>
</tbody>
</table>

**Fig. 10.** Output optical power of MKR with and without graphene as a function of relative humidity.