

Highly efficient sers performance from the silver nanoparticles/graphene nanoribbons/ cellulose paper

Tieu Tu Doanh^{1,2}, Thai Duong², Nguyen Cong Danh², Ton Nu Quynh Trang¹, Ngo Vo Ke Thanh²,
Vu Thi Hanh Thu^{1,*}, Nguyen Van Cattien^{3,*}



Use your smartphone to scan this QR code and download this article

¹Faculty of Physics and Engineering Physics, VNUHCM-University of Science

²Saigon Hitech Park Labs

³Ntherma Corporation

Correspondence

Vu Thi Hanh Thu, Faculty of Physics and Engineering Physics, VNUHCM-University of Science

Email: vththu@hcmus.edu.vn

Correspondence

Nguyen Van Cattien, Ntherma Corporation

Email: cattien.nguyen@ntherma.com

History

- Received: 2020-05-09
- Accepted: 2020-08-18
- Published: 2020-09-04

DOI : 10.32508/stdj.v23i3.2390



Copyright

© VNU-HCM Press. This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International license.



ABSTRACT

Introduction: Metal/graphene heterojunction structure has been one of the most crucial tools in the growth of high-performance Surface-enhanced Raman spectroscopy (SERS) platform, which is appropriate for sensing applications. In this research, we developed a SERS platform, graphene nanoribbons (GNRs) decorated silver nanoparticles (AgNPs) on cellulose paper substrate, in which GNRs was synthesized by wet chemical based on unzipping process of Multi-walled carbon nanotubes (MWCNTs), then GNRs hybridized with AgNPs through magnetron sputtering method.

Methods: The morphology of graphene nanoribbons coated-Ag (AgNPs@GNRs) was analyzed by field emission scanning electronic microscopy (FESEM) and transmission electron microscopy (TEM). The quality and thermal stability of GNRs were characterized using thermogravimetric analysis. Besides, its structure and quality were also characterized by Raman spectroscopy.

Results: The results show that the unzipping process of MWCNTs to form GNRs was strongly affected by dispersing time and stirring temperature. The suitable condition creating the Graphene Nanoribbons using MWCNTs was the dispersing time of 10 mins in an acid environment, stirring in 30 mins at room temperature and in 45 min at 100°C. Moreover, SERS platform of AgNPs@GNRs exhibit the outstanding SERS signal enhancement with rhodamine 6G (R6G) low concentration of 10^{-5} M compared with pristine graphene and silver thin film. This could be attributed to the synergistic effect between AgNPs, GNRs and analyze molecules based on the enhancement of electromagnetic mechanism (EM) and chemical mechanism (CM), which plays a vital role in promoting the improvement SERS behavior.

Conclusion: Ag NPs assembled onto graphene nanoribbons/ cellulose paper substrate could also serve as SERS active substrates for practical applications in various fields at trace levels.

Key words: Surface-enhanced Raman spectroscopy (SERS), Electromagnetic Mechanism, Chemical Mechanism, Carbon nanotubes, Graphene, Graphene Nanoribbons, Ag nanoparticles

INTRODUCTION

Raman spectroscopy is one of the techniques to detect molecules and provide their structural information based on the vibration energy levels of analytical molecules. However, it has some drawbacks such as the low scattering cross-section, the weak intensity of Raman signals, resulting in limitation in applications for detecting the low concentration of species. Many pathways which could improve the Raman signals have been proposed. In the 1970s, the Surface-Enhanced Raman Scattering (SERS) was developed^{1,2} and became one of the important analytical tools involving in the interaction between analytical molecules and a typical rough noble nanostructured metal surface leading a significant enhancement of the Raman scattering intensity.

Although the significant SERS substrate enhancement has been observed, however, there are now still arguments about their enhancement mechanism^{3,4}. In re-

cent years, SERS performance based on the enhancement of electromagnetic mechanism (EM) and chemical mechanism (CM) has attracted the widespread concern of researchers. First, EM plays a significant role in enhancing the sensitivity of SERS platform based on the localized surface plasmon resonance (LSPR) when the incident light interacts with the nanostructured metal surface, which generates the local electromagnetic field. It is usually called a “hot spot” and is one of the most factors for being responsible for improving the SERS performance. As a result, the enhancement factor (EF) for this mechanism can reach $10^8 - 10^{14}$ ⁵⁻⁸. The LSPR depends on the size, shape of nanoparticles, density, and gaps, which will affect the cross-section of Raman scattering^{9,10}. Second, the CM is associate with the charge transfer between the adsorbed molecule and the SERS substrate. In the charge transfer process, if the increasing separation of positive and negative charge in the molecule

Cite this article : Doanh T T, Duong T, Danh N C, Trang T N Q, Thanh N V K, Thu V T H, Cattien N V. **Highly efficient sers performance from the silver nanoparticles/graphene nanoribbons/ cellulose paper**. *Sci. Tech. Dev. J.*; 23(3):684-693.

is observed, resulting in increasing the cross-section of Raman scattering. The CM is often difficult to observe because it exhibits in a short time, and the EF can obtain from $10^1 - 10^3$ ⁶⁻⁸.

Graphene is a monolayer of sp^2 bonded carbon atoms packed into a honeycomb-like crystalline structure. It has wonderful physical and chemical properties such as thermal, chemical, electrical, mechanical and considers as a potential candidate for practical applications. It has a large surface area, superior molecule adsorption ability¹¹. Furthermore, it can quench the photoluminescence of fluorescent dyes, indicating the fluorescence background of analyzing molecules in SERS measurement can eliminate. Graphene nanoribbons (GNRs) are a member of the graphene family, 1D ribbons with widths in the nanometer range and length in the micrometer, exhibits a variety of electronic properties based on its structure¹². Itself graphene does not consider as a good candidate for SERS because it has the low cross-section Raman scattering¹³. The first observation of graphene material for SERS was reported by Ling and coworkers in 2010¹⁴. He reported that the GERS (Graphene-based SERS) showed an excellent charge transfer in SERS substrate. The phthalocyanine (Pc), rhodamine 6G (R6G), protoporphyrin IX (PPP) were deposited on graphene as a submonolayer. The highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) of these molecules stand on the two sides of the Fermi levels of graphene. Therefore, the CM is generated due to the occurrence of charge transfer between graphene and the molecules. However, the EF of CM is small to improve the SERS performance. This means graphene should cooperate with the noble metal nanoparticles (Au, Ag) for improving the Raman intensity^{15,16}. Furthermore, Ag NPs incorporating graphene can be avoided by the corrosion and oxidizing process, suggesting that the stability of SERS substrate could improve significantly.

Up to now, a lot of graphene-based substrates have been developed and used widely for the SERS platform with ultrasensitive, reproducible, and stable as well. Depend on the role of graphene material, we can classify graphene-metal substrates into four categories: a) graphene as matrix-supported metal nanostructures substrates, b) graphene as shield covered metal nanostructures substrates, c) graphene as sub-nanospacer separated metal nanostructures substrates and d) graphene as both the bottom platform and top shield sandwiched metal nanostructures substrates¹³. Among them, the first approach, which involves metal nanoparticles is inexpensive and

convenient, is directly grown onto graphene sheets and has been attracted widespread attention. Hsu and Chen et al.¹⁷ fabricated Ag NPs/rGO (reduced Graphene Oxide) using a microwave-assisted technique to detect 4-aminothiophenol (4-ATP) with a low concentration of 10^{-10} M, and the EF value obtained 1.27×10^{10} . While Caires et al. reported that the Au NRs (nanorods)/GO hybrids structure could analyze the Cresyl Violet perchlorate (CVP) molecule with a detection limitation of 10^{-11} M and the EF value of 10^6 obtained¹⁸. Liang et al. showed that Au NPs sputtered onto the Graphene layer grown on by CVD method to detect adenine molecule with a limitation of 10^{-7} M¹⁹. Fu et al. reported that the sensitivity of Ag NPs deposited onto rGO was more 10-fold higher than that of the flat graphene-Au NPs for detecting Rhodamine 6G (R6G) molecule²⁰. Although the approaches, as mentioned above, have good results for SERS performance. However, these substrates almost have been experienced the multiple steps to fabricate, time-consuming. Particularly, they are difficult to large scale samples that meet the requirement of promising applications.

For the above reasons, a simple approach fabricating the AgNPs/Graphene Nanoribbons/Cellulose paper substrate with a large scale, uniform surface, and flexibility for detecting Rhodamine 6G (R6G) molecule at low concentration has been processed. Therefore, in this research, the Graphene Nanoribbons are made via wet chemical method using raw MWCNTs, then they are deposited onto cellulose paper via vacuum filtration. Finally, Ag NPs were decorated on Graphene Nanoribbons/Cellulose paper substrate by magnetron sputtering method controlling the distance, distribution, and size of these Ag NPs. Their SERS performance was carried out by cutting it into small pieces to evaluate the detection capability of R6G molecules.

EXPERIMENT

Materials

The multi-walled carbon nanotubes (MWCNTs, Ntherma Corp, USA), Potassium permanganate ($KMnO_4$, Sigma-Aldrich, $\geq 99\%$), Sulfuric acid (H_2SO_4 99.999%, Sigma-Aldrich), Hydrogen peroxide (H_2O_2 , 50%, Solvay, Thailand), Isopropanol (IPA, $(CH_3)_2CHOH$, 99.5%, Sigma-Aldrich), Ag target (99.99% pure, Singapore Advantech), Whatman Grade 589 Cellulose paper (Sigma-Aldrich) and Rhodamine 6G dye (R6G, $C_{28}H_{31}N_2O_3Cl$, 99%, Sigma-Aldrich) were used. Analytical reagents were used as received without any further purification. All of the aqueous solutions were prepared using de-ionized (DI) water.

Fabrication

Fabrication the Graphene Nanoribbons (GNRs)

GNRs were successfully fabricated via a wet chemical method. The detailed preparation process is described in **Figure 1**.

Firstly, 1.5 g MWCNTs powder was added into 400 ml of H₂SO₄ 98%. The mixture was stirred in 20 minutes. Then, 7.5 g KMnO₄ was added to the solution and continuously stirred in 1 hour at room temperature. After that, the solution was cooled down under 10°C, followed by the addition of H₂O₂ solution (20 ml, 50%) and stirred progressively. When the reaction was finished, this solution was washed with DI water and filtered using the filter paper. Finally, the sample was dried in a vacuum oven in 12 hours.

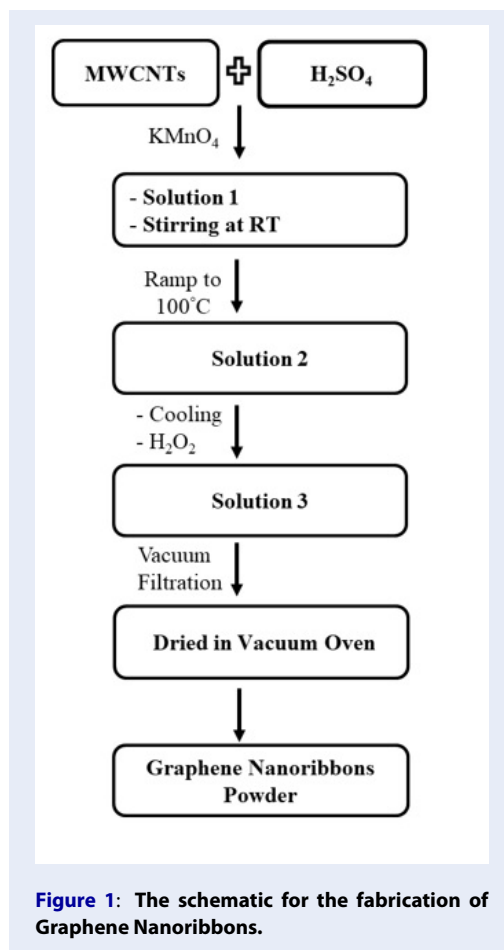


Figure 1: The schematic for the fabrication of Graphene Nanoribbons.

Preparation of Ag NPs/ Graphene Nanoribbons/ Cellulose paper substrate (Ag/GNRs/CP)

0.1g Graphene Nanoribbons were dispersed into 500 ml of IPA by high power ultrasonication. This solution was poured into the vacuum filtration system to make Graphene paper and then coated Ag NPs onto it by a sputtering technique. Sputtering time was 10 seconds in Ar environment with a total pressure of approximately 2.2 mTorr and a power 11W as our previous work²¹. The preparation process is described in **Figure 2**.

Characterization

The surface morphology and structure of the samples were characterized by Field-Emission Scanning Electron Microscopy (FESEM, Hitachi, S-4800), Transmission Electron Microscopy (TEM, Jeol JEM1400 and JEM1010). The Thermo Gravimetric Analysis (TGA, TG-DSC 1600, Labsys Evo) technique was used to define the quality and thermal stability of Graphene Nanoribbons. The characteristic peaks of Graphene Nanoribbons were evaluated by the Raman spectroscopy (Labram 300, Horiba Jobin Yvon).

SERS measurements

Raman spectra were collected using a Horiba XploRA PLUS Raman system with a 532 nm laser, power of 1.5 mW, and an objective with 100x magnification. The Rhodamine 6G (R6G) molecule is chosen to analyze and dissolved into DI water. Then the R6G solution was dropped onto GNRs, Ag NPs and Ag NPs/GNRs substrates, respectively, and dried in air. Each spectrum was obtained with acquisition time 1 second on 2 accumulation spots and repeated three times in various positions in the wavenumber range 500 – 2000 cm⁻¹.

RESULTS

Figure 3 (a and b) showed the FESEM and TEM images of raw MWCNTs from the supplier with the diameter in the range of 8-15 nm. Furthermore, they exhibited Raman peaks characteristic at D-peak (1350 cm⁻¹) and G-peak (1580 cm⁻¹) (**Figure 3c**) and were extremely purity (+ 99% C) (**Figure 3d**) which played an important factor in affecting the final graphene Nanoribbons product.

After being dispersed the MWCNTs in acid solution, the morphological characteristics of MWCNTs remained stable without damaging, as shown in **Figure 4**. Besides, these sample's purity was at + 99%

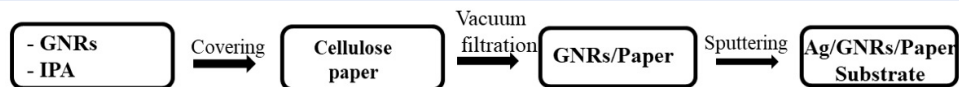
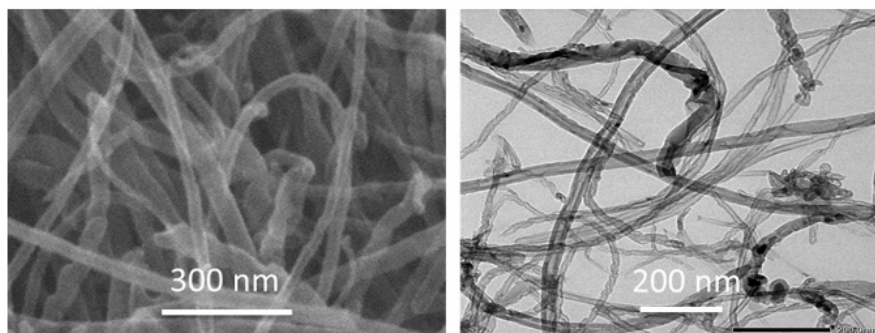
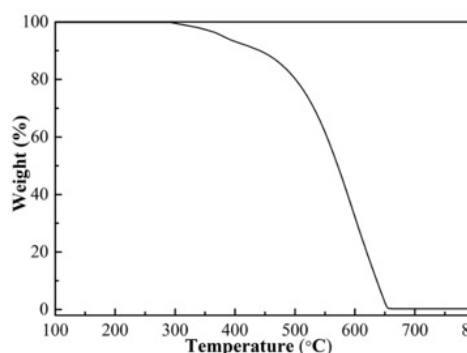


Figure 2: The process fabrication of AgNPs/ Graphene Nanoribbons/ Cellulose paper.

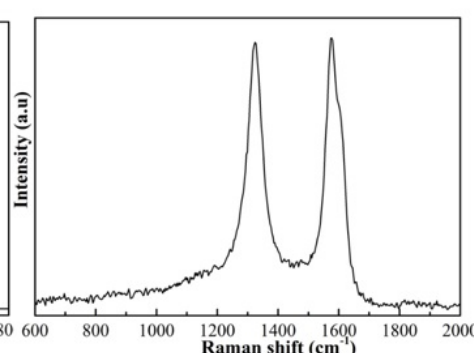


a)

b)



c)



d)

Figure 3: (a, b) the FESEM – TEM images, (c, d) TGA spectroscopy and Raman spectroscopy of raw MWCNTs.

C, and their structure was consistent with two Raman peaks at D-peak (1350 cm^{-1}) and G-peak (1580 cm^{-1}) as shown in Figure 5 (a and b), respectively. The acid solution is a good environment for dispersing them and reacting in the next steps. Thus, dispersing time was 10 mins to get a well-dispersed solution. After dispersing MWCNTs into the acid environment, KMnO_4 salt was added slowly and constantly stirred. Oxidation reaction occurred, and Graphene Nanoribbons were gradually formed (Figure 6). However, MWCNTs were unzipped at some outer walls and reached the unzipping saturation at a reaction time of 30 mins. Then the stirring temperature was increased to accelerate the reaction to get full unzip of MWCNTs.

The reaction was performed at temperatures 100°C in 1 hour to investigate the ability to unzip MWCNTs to form Graphene Nanoribbons. The experiments did not make over 100°C to avoid evaporating the solution in the reaction. At high temperature, the unzipping MWCNTs increases (Figure 7). This is consistent with the results^{22,23}. Especially, the reaction at 100°C in time conditions of 15, 30, and 45 mins, the Graphene Nanoribbons images are very clear, almost walls of MWCNTs were unzipped while in 60 mins the Graphene Nanoplatelets were formed (Figure 7B). All these samples are high purity (+ 99% C) (Figure 8a), showing their quality is good and satisfiable for Graphene paper and SERS application as well. In stirring time 60 mins, in Raman spectrum, the ratio $I_{2D}/I_D > 1$ demonstrating that this sample is

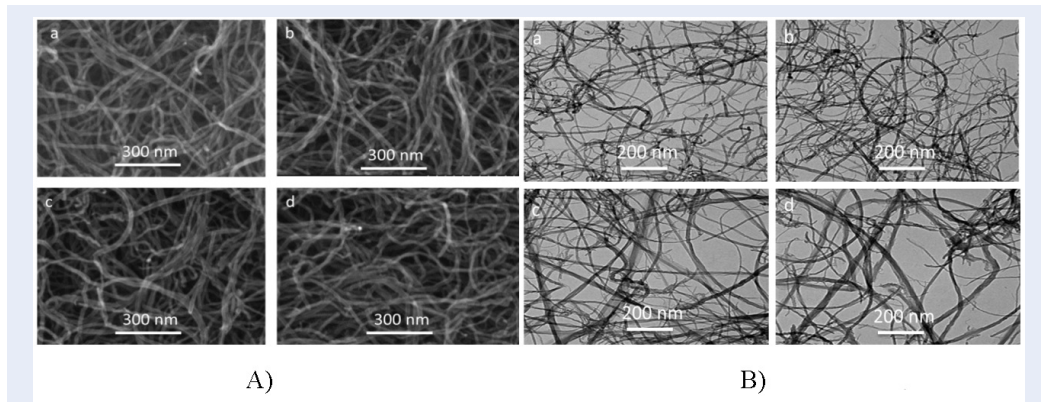


Figure 4: (A, B) the FESEM - TEM of MWCNTs dispersing in acid in different times a) 5 mins, b) 10 mins, c) 15 mins and d) 20 mins, respectively.

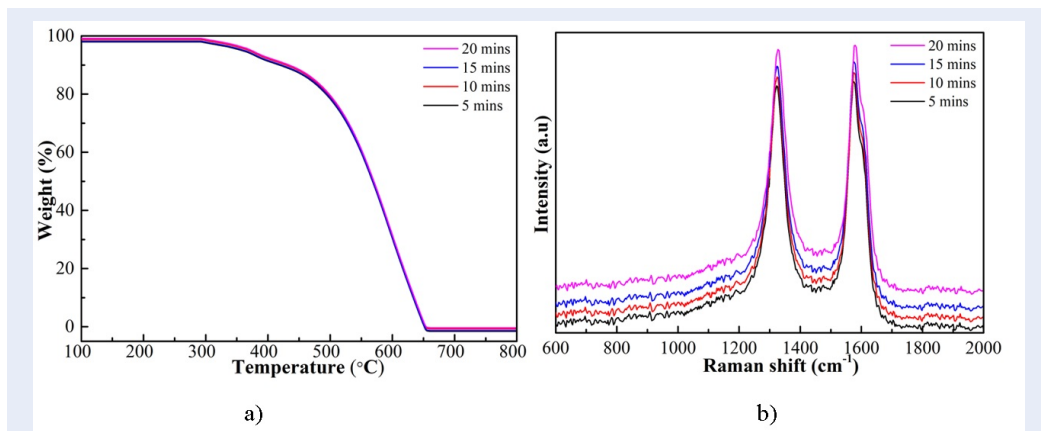


Figure 5: (a, b) the TGA and Raman spectroscopies of MWCNTs dispersing in acid at different times.

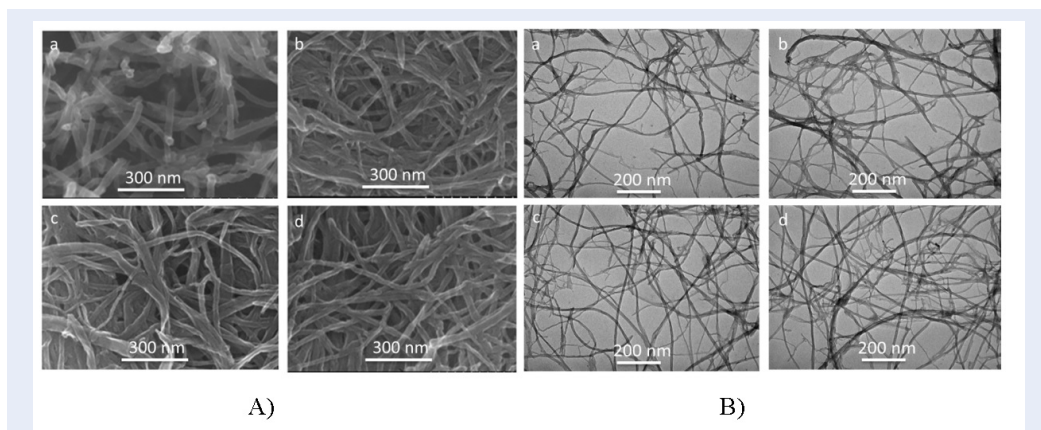


Figure 6: (A, B) the FESEM - TEM of Graphene Nanoribbons samples in reaction at room temperature in different times a) 15 mins, b) 30 mins, c) 45 mins and d) 60 mins, respectively.

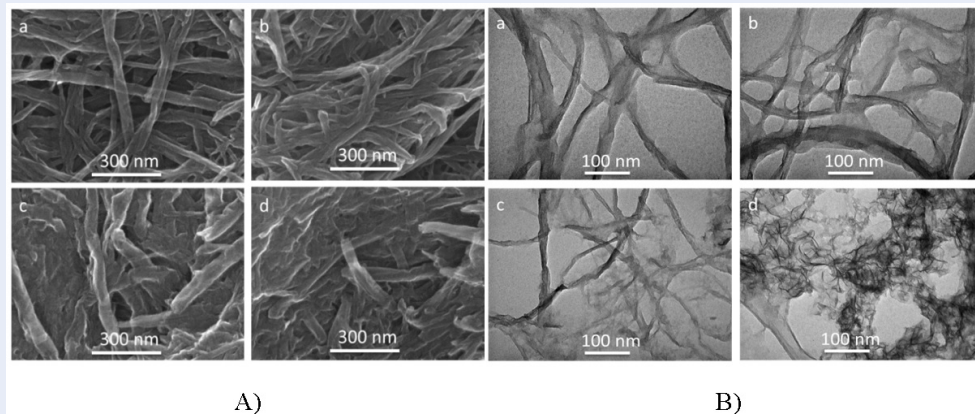


Figure 7: (A) the FESEM, (B) TEM of Graphene Nanoribbons samples in reaction at 100°C in different times a) 15 mins, b) 30 mins, c) 45 mins and d) 60 mins, respectively.

more defects, while in 15, 30, and 45 mins, the ratio $I_{2D}/I_D < 1$ showing the good crystallization (Figure 8 b). Furthermore, Graphene Nanoplatelets will not be suitable in the next step to make the free-standing Graphene paper because they are tiny flakes that can not grid together. The best reaction condition for Graphene Nanoribbons in this work is 45 mins and 100°C. It is also a condition for the next steps to make Graphene paper and decorate Ag NPs as well.

The Graphene paper is fabricated with the same process in our previous work²⁴. Besides, the procedure for Ag NPs was anchored onto the GNRs paper via the sputtering technique as in previous work²¹. The result shows that the Ag NPs distributed onto GNRs surface with a uniform diameter under 20 nm, no agglomeration with interparticle distances (nanogaps) about 2-5 nm (Figure 8). Especially, no agglomeration is observed in the SEM image, resulting in having the strong electric field distribution, which is important in the performance of SERS substrate. Furthermore, the surface of the sample is clean because there are not any extraordinary substances. It is ideal for identifying the typical signals of R6G molecule.

Herein, the SERS behaviors of GNRs, AgNPs, and Ag/GNRs/CP, which were used as SERS substrates were examined using Rhodamine 6G (R6G) as a target molecule under the excitation wavelength of 532 nm. In Figure 10, the typical peaks of R6G onto GNRs, Ag NPs, and Ag NPs/ GNRs substrates were shown. They are 612 cm^{-1} , 775 cm^{-1} , 1181 cm^{-1} , 1311 cm^{-1} (1363 , 1511 , and 1651 cm^{-1}), and 1575 cm^{-1} could be due to C-C-C ring in-plane, C-H out-plane bending, C-H in-plane bending, C-O-C stretching, C-C

stretching of the aromatic ring, and $\text{C}\frac{1}{4}\text{O}$ stretching, respectively^{25,26}. To the GNRs substrate, the Raman signal of R6G at peaks 612 , 775 , 1181 , and 1651 cm^{-1} can be distinguishable while 1131 , 1363 , 1151 , and 1575 cm^{-1} peaks are weak signals. With Ag NPs substrate, the Raman signal of R6G at peaks 612 , 775 , 1181 , 1363 , 1151 , and 1651 cm^{-1} are clear while peaks 1131 and 1575 cm^{-1} are dim. However, when the Ag NPs/GNRs substrate is used, the Raman signal intensity of R6G at all peaks is increased and enhanced significantly. This is explained that there is a combination of both EM and CM. The EM occurs when the Ag NPs play as hot spots that generate the surface plasmon under the excitation of the incident light. While the CM associates the charge-transfer transitions between the Fermi level of the Ag, GNRs, and R6G molecules. This suggests that the Ag/GNRs/CP platform becomes a favorable substrate for SERS applications.

DISCUSSION

The GNRs formation from MWCNTs was investigated. Firstly, the MWCNTs were dispersed into H_2SO_4 acid, and this showed that the stirring time did not affect the quality of MWCNTs in the acid environment. It means that the MWCNTs did not be almost shortened or damaged (Figure 4). Then, at the room temperature, the MWCNTs initially seemed to be unzipped some outer layers (Figure 6).

Tour *et al.*²² gave the mechanism of unzipping CNTs based on the oxidation of alkenes by permanganate in acid medium. Firstly, the manganate ester was formed (2, Figure 11 b) and could induce the dione (a molecule containing the ketones group) in the dehydrating medium. Next, the ketones group distorted

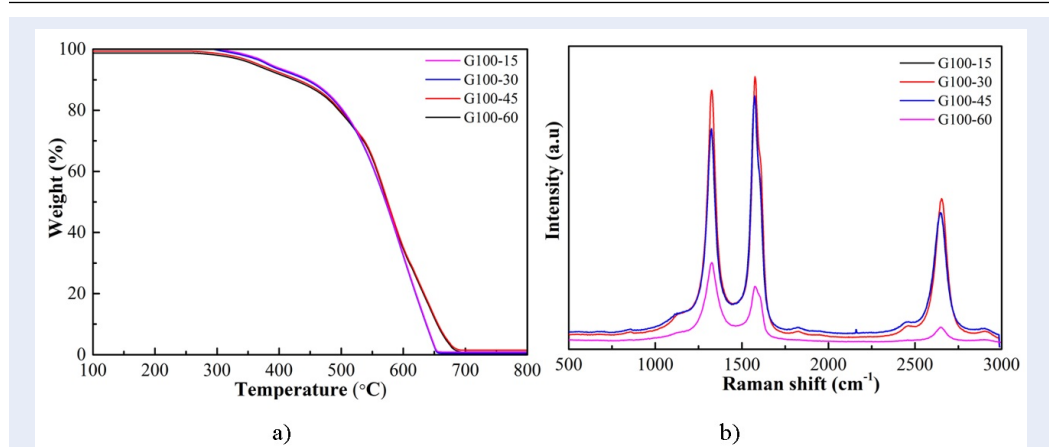


Figure 8: (a,b) The TGA and Raman spectroscopy of Graphene Nanoribbons samples in reaction at 100 °C at different times.

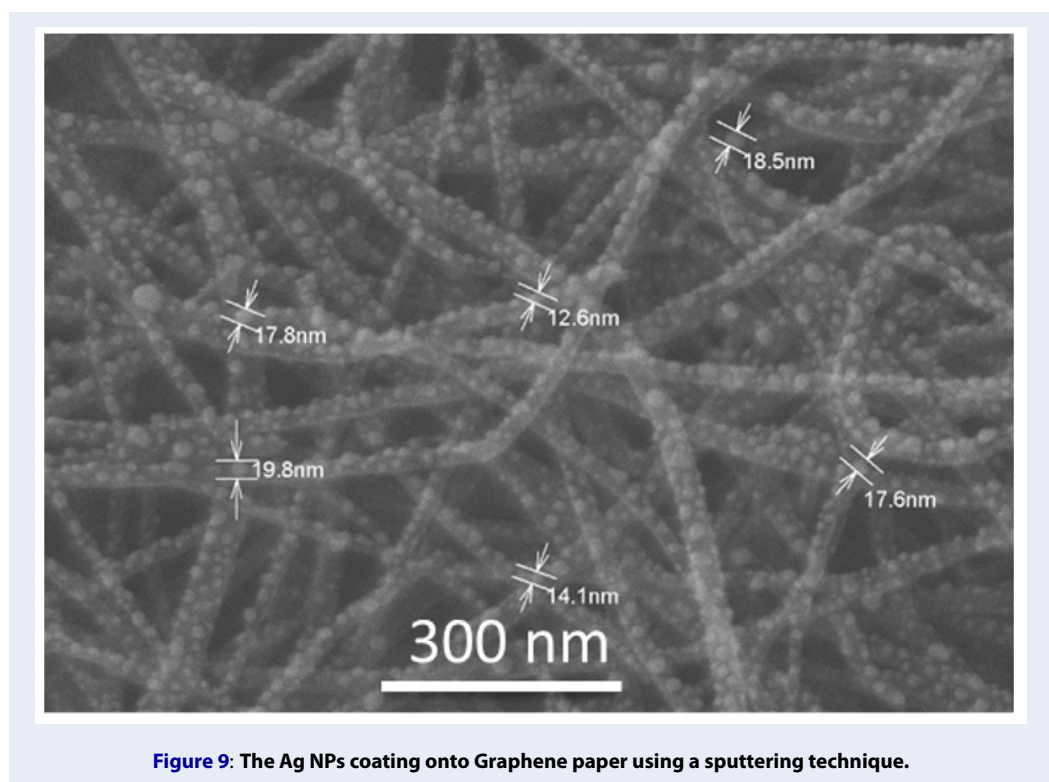


Figure 9: The Ag NPs coating onto Graphene paper using a sputtering technique.

the β - γ alkenes (3, Figure 11 b), and they activated easily with permanganate. When the process continued, the buttressing-induced strain on the β - γ alkenes increased (4, Figure 11 b), and it broadens the space for carbonyl. As a result, the strain is induced by the enlarging hole on the CNTs. Finally, the unzipping process of CNTs was initiated to open to the graphene nanoribbons (5, Figure 11 b), and it was described in Figure 11.

Furthermore, Tour et al.²² suggested the optimal weight ratio between MWCNTs/KMnO₄ = 1/5 and stirring time of 1 hour at 55-70 °C in the reaction and could obtain 100% yield of nanoribbon structures. While Dimiev et al.²³ suggested that the optimal weight ratio between MWCNTs/KMnO₄ = 1/3 could achieve the highest GNRs formation. However, the time or temperature of the reaction was not recorded. In this study, the stirring time from 15 to 60 mins at

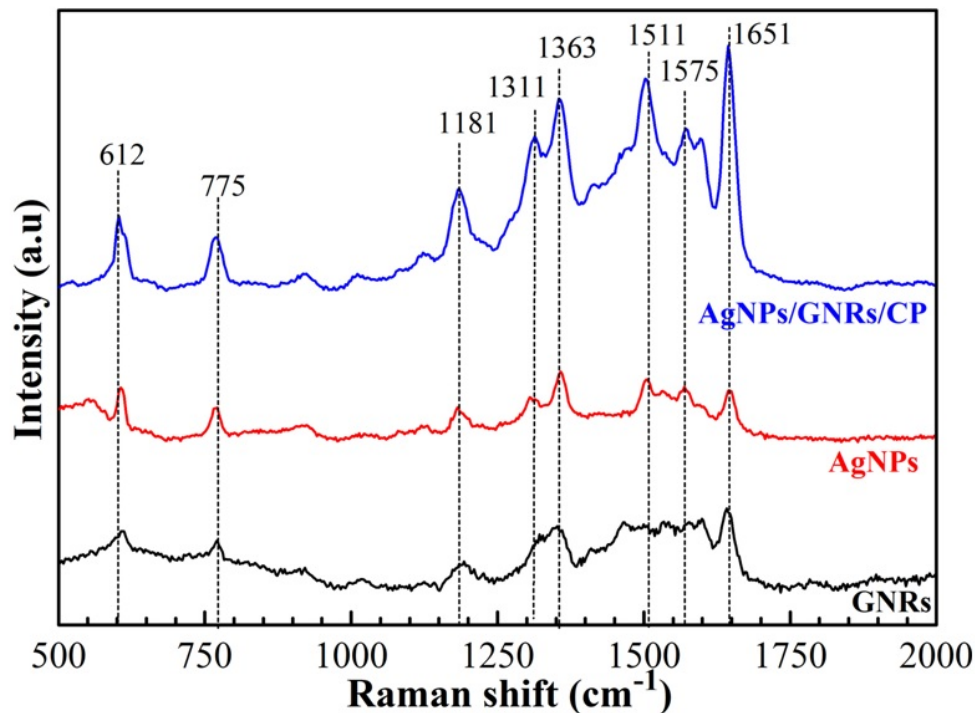


Figure 10: The SERS behavior spectra of the R6G molecules on the GNRs, Ag NPs, and Ag NPs/GNRs substrates.

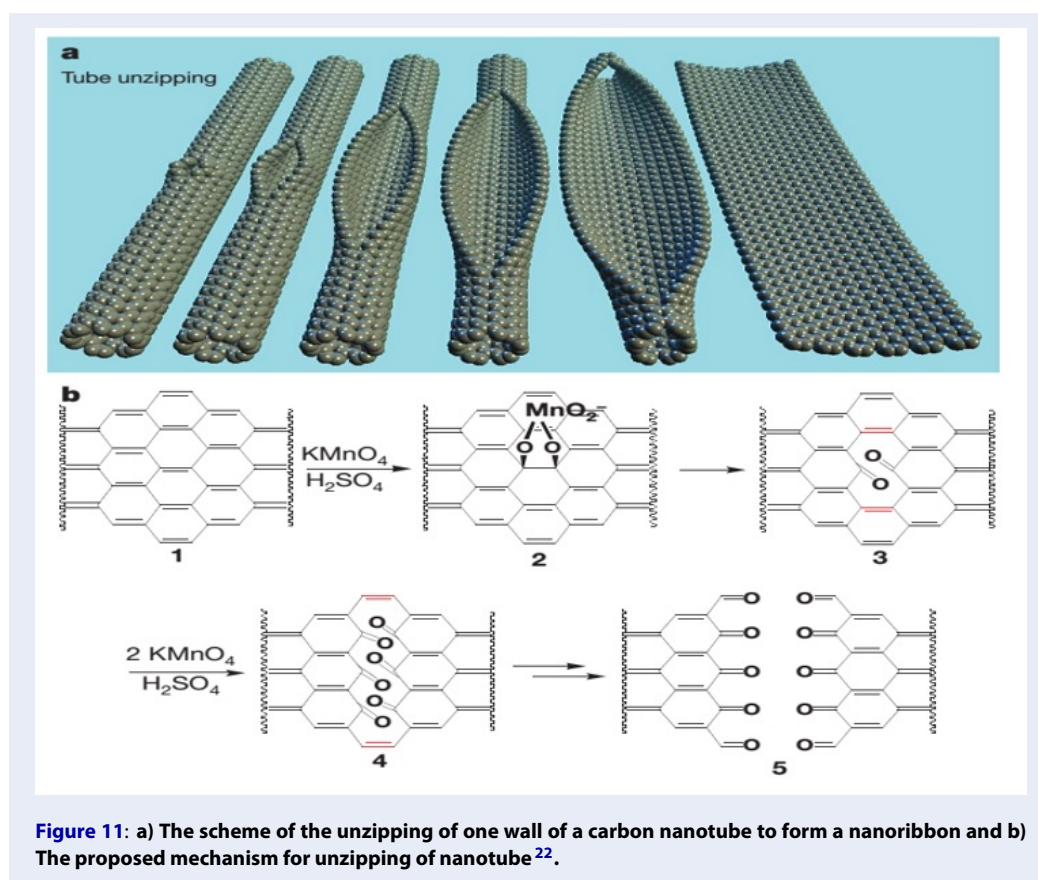
a temperature of 100°C, we can get the highest GNRs formation at the stirring time of 45 mins and a temperature of 100°C. (Figure 8 b). Therefore, this will affect the quality of both Graphene paper and SERS performance as well.

Goul *et al.*¹⁵ showed that graphene has fabricated using a chemical vapor deposition (CVD) method and incorporated with gold nanoparticles (AuNPs) by e-beam evaporation method. This SERS substrate could detect R6G with a low concentration of about 8×10^{-7} M. He reported that the graphene contributed to the enhancement of SERS performance from based on a charge-transfer mechanism related to the EM and CM mechanism. Moreover, Zhou *et al.*¹⁶ revealed that the EF of rhodamine 6G molecules only obtained 5.6×10^7 due to the presence of the Ag NPs aggregates in this SERS structure. Hsu *et al.*¹⁷ reported that Ag/reduced graphene oxide substrate was prepared by the microwave-assisted green synthesis. The Ag/reduced graphene oxide substrate was used to analyze the 4-aminothiophenol (4-ATP) molecules with limitation detection about 10^{-10} M, and EF value was 1.27×10^{10} . All the above studies talked about the enhancement of the Raman scattering signals with the contribution of graphene via a charge-transfer phenomenon and surface plasmon resonance of noble

metal nanoparticles (Ag, Au NPs). In our study, the SERS structure contains both graphene nanoribbons and silver nanoparticles. The target is to detect R6G molecules at low concentrations. The SERS performance of Ag/GNRs/CP substrate exhibits a remarkably high enhanced Raman intensity compared with that of AgNPs and GNRs substrate. It could be noticed that combine the properties of GNRs and AgNPs plays an important role in the enhancement of the Raman scattering signal.

CONCLUSION

A simple pathway to fabricate the Silver nanoparticles/Graphene Nanoribbons/ Cellulose paper toward Surface-Enhanced Raman Spectroscopy application is presented in this work. The process for Graphene Nanoribbons formation is investigated in detail, including the disperse MWCNTs in acid, unzipping process at room temperature, high temperature at various times. We suggest that the suitable conditions to create the Graphene Nanoribbons using MWCNTs for SERS platform were stirring in 45 mins at 100°C. Moreover, the SERS performance of each AgNPs, GNRs, and Ag/GNRs platform was also reported. The SERS substrate with a uniform diameter of Ag NPs



and no agglomeration exhibited a significant Raman intensity enhancement of the R6G molecule detection with the Ag NPs/GNRs substrate. This work presents a facile route to fabricate the SERS platform with high flexibility and sensitivity. The SERS substrate could offer a new approach for the detection of trace species in biochemical research and environmental monitoring.

ABBREVIATIONS

GNRs: Graphene Nanoribbons
MWCNTs: Multi-walled carbon nanotubes
Ag NPs: Silver nanoparticles
SERS: Surface-enhanced Raman spectroscopy
EM: Electromagnetic mechanism
CM: Chemical mechanism
LSPR: Localized surface plasmon resonance
R6G: Rhodamine 6G

COMPETING INTERESTS

The author(s) declare that they have no competing interests.

AUTHOR'S CONTRIBUTION

Tieu Tu Doanh has conceived of the present idea, carried out and written the manuscript with support from Vu Thi Hanh Tu, Nguyen Van Cattien.

Tieu Tu Doanh, Thai Duong, Nguyen Cong Danh and Ton Nu Quynh Trang carried out the experiments in group.

Ton Nu Quynh Trang and Ngo Vo Ke Thanh have supported the analysis techniques.

ACKNOWLEDGEMENTS

This research is funded by the Saigon Hitech Park Labs, under grant number NVTXTCN 7/2020.

REFERENCES

1. Fleischmann M, Hendra PJ, McQuillan AJ. Raman spectra of pyridine adsorbed at a silver electrode. *Chemical Physics Letters*. 1974;26(2). Available from: [https://doi.org/10.1016/0009-2614\(74\)85388-1](https://doi.org/10.1016/0009-2614(74)85388-1).
2. Albrecht MG, Creighton JA. Anomalously intense Raman spectra of pyridine at a silver electrode. *Journal of the American Chemical Society*. 1977;99(15):5215–5217. Available from: <https://doi.org/10.1021/ja00457a071>.
3. Persson BNJ, Zhao K, Zhang Z. Chemical Contribution to Surface-Enhanced Raman Scattering. *Physics Review Letters*. 2006;96(20):207401. PMID: 16803204. Available from: <https://doi.org/10.1103/PhysRevLett.96.207401>.

4. Wu DY, Liu XM, Duan S, Xu X, Ren B, Lin SH, et al. Chemical Enhancement Effects in SERS Spectra: A Quantum Chemical Study of Pyridine Interacting with Copper, Silver, Gold and Platinum Metals. *The Journal of Physical Chemistry*. 2008;112(11):4195–4204. Available from: <https://doi.org/10.1021/jp0760962>.
5. Nie S, Emory SR. Probing Single Molecules and Single Nanoparticles by Surface-Enhanced Raman Scattering. *Science*. 1997;275(5303):1102–1106. PMID: 9027306. Available from: <https://doi.org/10.1126/science.275.5303.1102>.
6. Kneipp K, Wang Y, Kneipp H, Itzkan I, Dasari RR, Feld MS. Population Pumping of Excited Vibrational States by Spontaneous Surface-Enhanced Raman Scattering. *Physics Review Letters*. 1996;76(14):2444–2447. PMID: 10060701. Available from: <https://doi.org/10.1103/PhysRevLett.76.2444>.
7. Kneipp K, Wang Y, Kneipp H, Perelman LT, Itzkan I, Dasari RR, et al. Single Molecule Detection Using Surface-Enhanced Raman Scattering (SERS). *Physics Review Letters*. 1997;78:1667. Available from: <https://doi.org/10.1103/PhysRevLett.78.1667>.
8. Dieringer JA, Nd LR, Scheidt KA, Duynes RPV. Ultrastable Substrates for Surface-Enhanced Raman Spectroscopy Fabricated by Atomic Layer Deposition: Improved Anthrax Biomarker Detection. *Journal of the American Chemical Society*. 2006;128(31):10304–10309. PMID: 16881662. Available from: <https://doi.org/10.1021/ja0638760>.
9. Wang H, Zhou Y, Jiang X, Sun B, Zhu Y, Wang H, et al. Simultaneous Capture Detection and Inactivation of Bacteria as Enabled by a Surface - Enhanced Raman Scattering Multifunctional Chip. *Angewandte Chemie*. 2015;54(17):5132–5136. PMID: 25820791. Available from: <https://doi.org/10.1002/anie.201412294>.
10. Zhan M, Yu H, Li L, Nguyen DT, Chen W. Detection of Hexavalent Chromium by Copper Sulfide Nanocomposite. *Analytical Chemistry*. 2019;91(3):2058–2065. PMID: 30621389. Available from: <https://doi.org/10.1021/acs.analchem.8b04501>.
11. Stoller D, Park S, Zhu Y, An J, Ruoff RS. Graphene-Based Ultracapacitors. *Nano Letters*. 2008;8(10):3498–3502. PMID: 18788793. Available from: <https://doi.org/10.1021/nl802558y>.
12. Talirz L, Ruffieux P, Fasel R. On-surface synthesis of atomically precise graphene nanoribbons. *Advanced Materials*. 2016;28(29):6222–6231. PMID: 26867990. Available from: <https://doi.org/10.1002/adma.201505738>.
13. Lai H, Zhang FXY, Wang L. Recent Progress on Graphene-Based Substrates for Surface enhanced Raman Scattering Application. *Journal of Materials Chemistry B*. 2018;6:4008–4028. PMID: 32255147. Available from: <https://doi.org/10.1039/C8TB00902C>.
14. Ling X, et al. Can graphene be used as a Substrate for Raman Enhancement? *Nano Letters*. 2010;10(2):553–561. PMID: 20039694. Available from: <https://doi.org/10.1021/nl903414x>.
15. Goul R, Das S, Liu QF, Xin M, Lu RT, Hui R, et al. Quantitative analysis of surface enhanced Raman spectroscopy of Rhodamine 6G using a composite graphene and plasmonic Au nanoparticle substrate. *Carbon*. 2017;111:386–392. Available from: <https://doi.org/10.1016/j.carbon.2016.10.019>.
16. Zhou Y, Yang J, Ma S, Zhao N, Cheng X, Zhong T. Fabrication of free-standing and flexible silver nanoparticle-graphene films and their surface-enhanced Raman scattering activity. *Chemical Monthly*. 2014;145:11–17. Available from: <https://doi.org/10.1007/s00706-013-0987-6>.
17. Hsu KC, Chen DH. Microwave-assisted green synthesis of Ag/reduced graphene oxide nanocomposite as a surface-enhanced Raman scattering substrate with high uniformity. *Nanoscale Research Letters*. 2014;9(1):193. PMID: 24808800. Available from: <https://doi.org/10.1186/1556-276X-9-193>.
18. Caires AJ, Alves DCB, Fantini C, Ferlauto AS, Ladeira LO. One-pot in situ photochemical synthesis of graphene oxide/gold nanorod nanocomposites for surface-enhanced Raman spectroscopy. *The Royal Society of Chemistry Advance*. 2015;5:46552–46557. Available from: <https://doi.org/10.1039/C4RA17207H>.
19. Liang X, Liang BL, Pan ZH, Lang XF, Zhang YG, Wang GS, et al. Tuning plasmonic and chemical enhancement for SERS detection on graphene-based Au hybrids. *Nanoscale*. 2015;7:20188–20196. PMID: 26575834. Available from: <https://doi.org/10.1039/C5NR06010A>.
20. Leem JY, Wang MC, Kang P, Nam SW. Mechanically Self-Assembled, Three-Dimensional Graphene-G7684-7690 old Hybrid Nanostructures for Advanced Nanoplasmonic Sensors. *Nano Letters*. 2015;15:11. PMID: 26501429. Available from: <https://doi.org/10.1021/acs.nanolett.5b03672>.
21. Tieu DT, Trang TNQ, Hung LVT, Thu VTH. Assembly engineering of Ag@ZnO hierarchical nanorod arrays as a pathway for highly reproducible surface-enhanced Raman spectroscopy applications. *Journal of Alloys and Compounds*. 2019;808:151735. Available from: <https://doi.org/10.1016/j.jallcom.2019.151735>.
22. Kosynkin D, Dimiev AM, Sinitskii A, Tour JM. Longitudinal Unzipping of Carbon Nanotubes to Form Graphene Nanoribbons. *Nature*. 2009;458:872–876. PMID: 19370030. Available from: <https://doi.org/10.1038/nature07872>.
23. Dimiev AM, Khannanov A, Vakhitov I, Kiiamov A, Shukhina K, Tour JM. Revisiting the Mechanism of Oxidative Unzipping of Multiwall Carbon Nanotubes to Graphene Nanoribbons. *American Chemical Society Nano*. 2018;12(4):3985–3993. PMID: 29578700. Available from: <https://doi.org/10.1021/acsnano.8b01617>.
24. Tieu DT, Nguyen MK, Nguyen TTH, Ngo TVK, Do QH. Study to fabricate the large scale Buckypaper based on Carbon Nanotubes. *Communication in Physics*. 2018;28(3). Available from: <https://doi.org/10.15625/0868-3166/28/3/10758>.
25. Zhou Q, Meng G, Huang Q, Zhu C, Tang H, Qian Y, et al. Ag-nanoparticles-decorated NiO-nanoflakes grafted Ni-nanorod arrays stuck out of porous AAO as effective SERS substrates. *Physical Chemistry Chemical Physics*. 2014;16(8). PMID: 24419246. Available from: <https://doi.org/10.1039/c3cp54119c>.
26. Lee J, Hua B, Park S, Ha M, Lee Y, Fan Z, et al. Tailoring surface plasmons of high-density gold nanostar assemblies on metal films for surface-enhanced Raman spectroscopy. *Nanoscale*. 2014;6:616–623. PMID: 24247586. Available from: <https://doi.org/10.1039/C3NR04752K>.