Supplementary Information

Electrochemical 4-chlorophenol sensing properties of plasma-treated multilayer graphene modified photolithography patterned platinum electrode

Padmanathan Karthick Kannan^{1,*}, Rogerio V. Gelamo², Hywel Morgan³, Palaniswamy Suresh⁴, Chandra Sekhar Rout^{5,*}

¹School of Chemical Engineering, Sungkyunkwan University, Suwon 440-746, Republic of Korea

²Instituto de Ciências Tecnológicas e Exatas, UFTM, Uberaba, Minas Gerais 38064-200, Brazil

³Electronics and Computer Science, University of Southampton, Southampton SO17 1BJ, United Kingdom

⁴Department of Natural Products Chemistry, School of Chemistry, Madurai Kamaraj University, Madurai 625021, Tamil Nadu, India

⁵School of Basic Sciences, Indian Institute of Technology Bhubaneswar, Bhubaneswar, 751013, Odisha, India

E-mail: <u>pkk.matsci@gmail.com</u>, <u>csrout@gmail.com</u>, <u>csrout@iitbbs.ac.in</u>

Figure Captions

Fig. S1 Pictorial representation of the fabrication process for patterning Pt electrodes on a glass

substrate by photolithography

Fig. S2 Estimation of Surface area for PMLG sample from AFM image

Fig. S3 Estimation of Surface area for OMLG sample from AFM image

Fig. S4 Stability data measured of OMLG/Pt electrode measured in 0.1 M PBS with 10 μ M of

4-CP

 Table S1 Detection of 4-CP in tap water samples

Fabrication of Patterned Platinum Electrodes

A schematic diagram of the electrode fabrication is shown in Figure S1. A thin metal film (25 nm titanium (Ti) and 300 nm platinum (Pt)) was deposited by sputtering (Helios, Leybold Optics, USA) onto glass wafers (SCHOTT AG, UK), cleaned with fuming nitric acid (FNA) and dried at 230 °C. The platinum coated glass wafers were then cleaned with FNA, followed by Isopropyl Alcohol (IPA) (Fisher, UK) and dried in nitrogen (N₂). They were dehydrated at 230 °C for 2 hours. Positive photoresist S1813 (Chestech, Warwickshire, UK) was spin-coated on the wafer and baked at 95°C for 1 minute. An acetate mask (Micro Lithography Services Ltd, Essex, UK) was used to define the electrode features. After exposure of the resist, the wafer was developed, rinsed and dried in N₂. Ion beam milling (IBM) (Ion Fab, Oxford Instruments, UK) was used to etch the platinum and the remaining resist removed with N-Methyl-2-pyrrolidone (NMP; Sigma-Aldrich, UK) with ultra-sonication at 60°C for 20 minutes. After IPA cleaning and N₂ drying, the wafer was transferred to RIE80+ (Oxford Instruments, UK) for O₂ descum cycle. The wafer was then cleaned with FNA, followed by SU8-5 (Chestech, Warwickshire, UK) was used to insulate the contact pads.



Fig. S1

Pristine MLG data 5.txt	- Bloco de notas — 🗆 🗙		-10
<u>Arquivo</u> <u>Editar</u> <u>Formata</u>	ır E <u>x</u> ibir Aj <u>u</u> da		
Statistical Quanti	ties		_
File:	C:\Users\Rogério\Documents\Resultados UFT		80
Data channel:	Value (max)		
Selected area:	165 x 406 at (11, 263) px		60
	0.646 × 1.589 at (0.043, 1.031) um		
Mask in use:	No		
Average value:	0.89 nm		40
Minimum:	-21.94 nm		
Maximum:	28.95 nm		
Median:	3.71 nm		20
Ra:	8.24 nm		
Rms:	9.91 nm		
Rms (grain-wise):	9.91 nm		
Skew:	-0.217		0
Kurtosis:	-0.434		
Surface area:	0.349102 µm^2		_
Projected area:	0.340576 µm^2		-7
Variation:	61.475 10 ⁻¹⁵ m ²		
Inclination θ :	2.84 deg	E00 nm	
Inclination ϕ :	-65.32 deg	Suu nm	
		~	
<	>		

Fig. S2

O2 treated MLG data.t	t - Bloco de notas — 🗆 🗙	
<u>Arquivo Editar Formata</u>	ar E <u>x</u> ibir Aj <u>u</u> da	
Statistical Quanti	ties	
File:	C:\Users\Rogério\Documents\Resultados UFT	Contraction of
Data channel:	Value (max)	
Selected area:	512 × 512 at (0, 0) px	X
	2.000 × 2.000 at (0.000, 0.000) µm	
Mask in use:	No	
Average value:	3.22 nm	
Minimum:	-62.62 nm	
Maximum:	92.89 nm	
Median:	2.04 nm	
Ra:	8.85 nm	
Rms:	14.03 nm	
Rms (grain-wise):	14.03 nm	
Skew:	1.33	
Kurtosis:	8.84	
Surface area:	4.267198 μm ²	
Projected area:	4.000000 µm^2	
Variation:	982.44 10^-15 m^2	
Inclination θ :	0.19 deg	and the second
Inclination ϕ :	60.55 deg	





Fig. S4

Sample	Added	Founded	RSD	Recovery
	(mM)	(mM)	(%)	(%)
	4	3.8	2.0	95
Tap water	8	7.3	1.5	91

Table S1