

Supplementary Information

Annealing effect on UV-illuminated recovery in gas response of graphene-based NO₂ sensors

Chia-Ming Yang^{*a,b,c,d}, *Tsung-Cheng Chen*^a, *Yu-Cheng Yang*^{a,b}, *M. Meyyappan*^e

a Department of Electronic Engineering, Chang-Gung University, Taoyuan 333, Taiwan

b Institute of Electro-Optical Engineering, Chang Gung University, Taoyuan 333, Taiwan

c Biosensor Group, Biomedical Engineering Research Center, Chang Gung University, Taoyuan 333, Taiwan

d Department of General Surgery, Chang Gung Memorial Hospital, Linkou 333, Taiwan

e Center for Nanotechnology, NASA Ames Research Center, Moffett Field, CA 94035, USA

Corresponding author: cmyang@mail.cgu.edu.tw; Tel.: +886-3-2118800#5960

C.-M. Yang and T.-C. Chen contributed equally to this work.

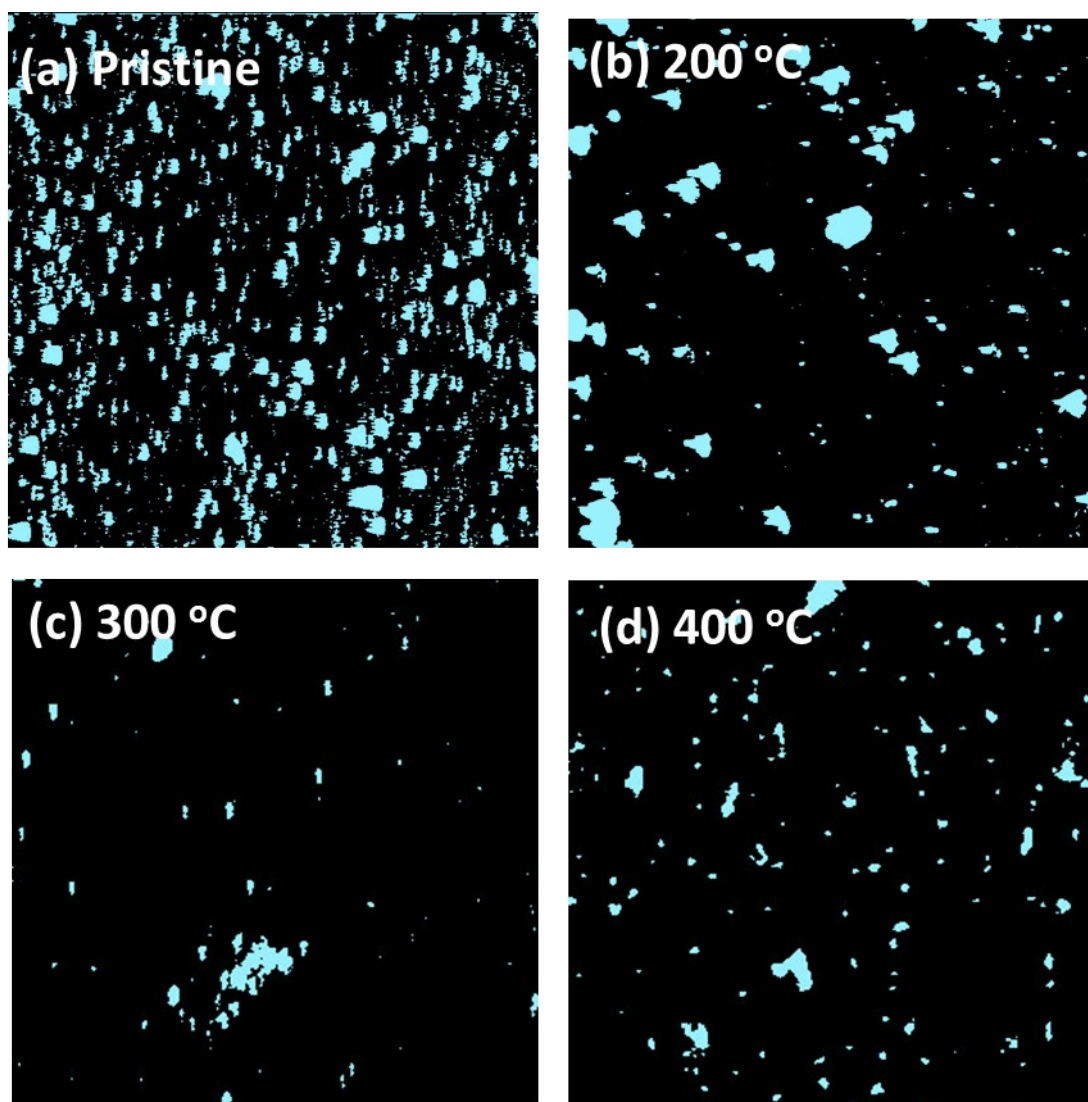


Fig. S1 AFM images analyzed in imageJ software: (a) pristine graphene and with annealing at (b) 200 °C, (c) 300 °C and (d) 400 °C. The red area could be referred to PMMA residues with a threshold value setting.

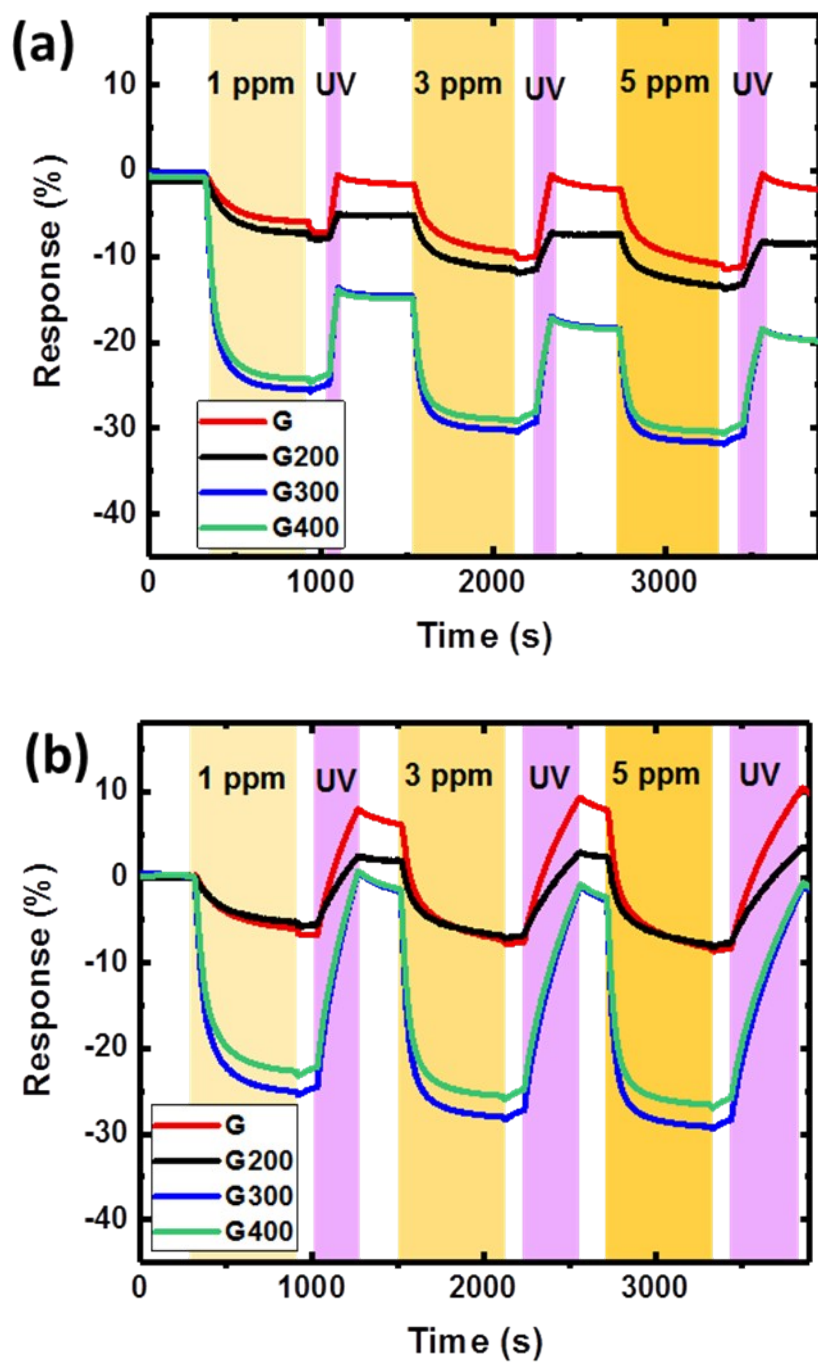


Fig. S2 Dynamic response versus time for the device treated with RTA at different temperatures and exposure to various concentrations of NO₂ with UV illumination during the recovery time. The illumination time is decided by the full recovery of the response of (a) the “G” device and (b) the “G300” device.

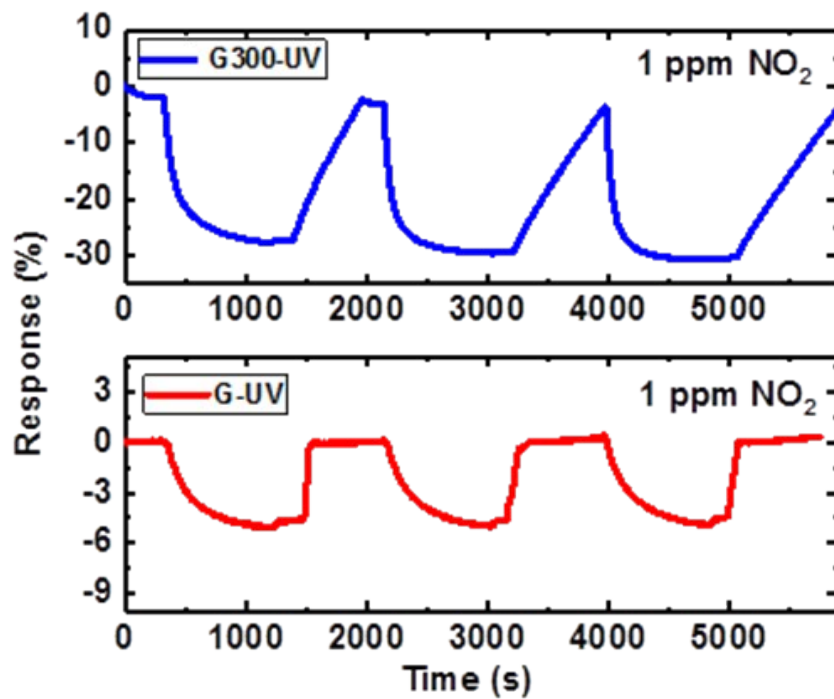


Fig. S3 Repeatability and stability studies of the graphene sensors exposed to 1 ppm of NO₂.

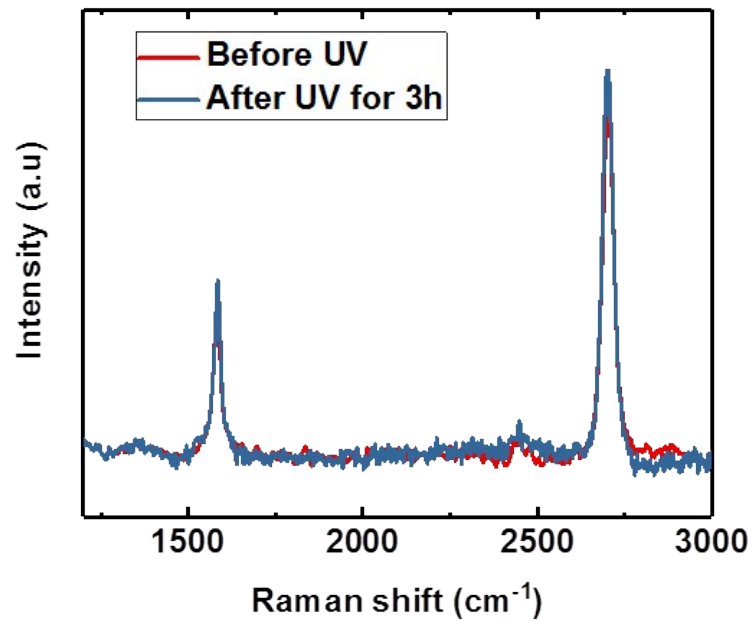


Fig. S4 Raman spectra of the pristine monolayer graphene before and after UV illumination for 3h.