Liposome-induced exfoliation of graphite to few-layer graphene dispersion with antibacterial activity

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1. Stability Measurements



Figure S1. Stability measurements for graphene-loaded liposome prepared in water (grey) or in PBS (black) over 48 h: (A) Dimensional variation expressed as mean diameter (intensity); (B) homogeneity of the suspension expressed as polydispersity index;^{S1} (C) changes of graphene concentration.



FigureS2. Variation over time of the ζ -potential of graphene-loaded liposomes prepared in water.



FigureS3. Stability measurements for graphene-loaded liposome in PBS as obtained after preparation (grey) and after 10-fold dilution (black): (A) dimensional variation expressed as mean diameter (intensity); (B) homogeneity of the suspension expressed as polydispersity index.^{S1}

2. Raman Fitting

In order to evaluate the degree of exfoliation of graphene, the 2D band at ~2700 cm⁻¹ was fitted as previously reported.^{S2-S4}

The fitting obtained from 2 different experiments and 6-8 different spots for each sample are reported below.



Figure S4. Fitting of the 2D band obtained from 8 different spots (A-H).



Figure S5. Fitting of the 2D band obtained from 6 different spots (A-F).

3. Precipitation of graphene embedded into liposomes



Figure S6. A) and C) Lipo-G in milli-Q water. B) Lipo-G aqueous suspensions upon addition of 10 μ L milli-Q water and D) Lipo-G aqueous suspensions upon addition of 10 μ L of aqueous Triton X-100 ([Triton X-100]_{final} = 3.5 mM). The white arrow indicates the formation of black precipitate.

4. Gram staining images





Figure S7. Representative Gram staining images of *Staphylococcus aureus* ATCC 29231 and *Escherichia coli* ATCC8739 grown for 2 h in the presence of GO and LIPO-G compared to the controls (SF and LIPO). The absence of a significant bacterial aggregation and a reduction in the total number of bacterial cells are clearly displayed in GO and LIPO-G treated samples. Images from optical microscope at 1000×.

References

- S1. J. Pereira-Lachataignerais R. Pons, P. Panizza, L. Courbin, J rouch, O López, *Chem. Phys. Lipids*, 2006, **140**, 88.
- S2. A. C. Ferrari, J. C. Meyer, V. Scardaci, C. Casiraghi, M. Lazzeri, F. Mauri, S. Piscanec, D. Jiang, K.
 S. Novoselov, S. Roth, A. K. Geim, *Phys. Rev. Lett.*, 2006, **97**, 187401.
- S3. A. C: Ferrari, D. M. Basko, Nat. Nanotechnol., 2013, 8, 235.
- S4. I. Calizo, D. Teweldebrhan, W. Bao, F. Miao, C. N. Lau, A. A. Balandin, *J. Phys.: Conf. Ser.*, 2008, 109, 012008.