

Electronic Supporting Information

A direct method for the preparation of glycolipid-metal nanoparticle conjugates: sophorolipids as reducing and capping agents for the synthesis of water redispersible silver nanoparticles and their antibacterial activity

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ESI-1: Preparation and characterization of acid sophorolipid: Was prepared according to the reported procedures.¹

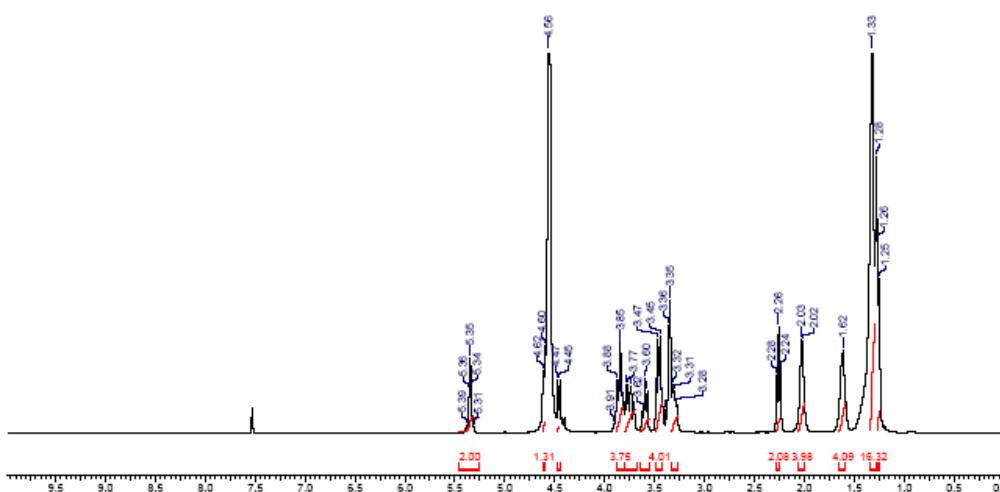
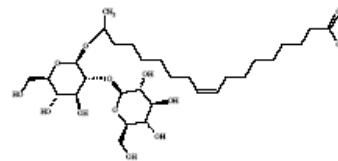
$[\alpha]_D^{25} = -12.5$ ($c = 0.1$ g/mL) lit.^{1b}: $[\alpha]_D^{25} = -12.8$ ($c = 0.0104$ g/mL). ^1H NMR (400 MHz, CDCl_3 - CD_3OD): δ 1.25 (d, $J = 6.2$ Hz, 3H), 1.28–1.46 (m, 16H), 1.57–1.64 (m, 4H), 2.0–2.04 (m, 4H), 2.26 (t, $J = 7.5$ Hz, 1H), 3.28–3.28 (m, 1H), 3.28–3.49 (m, 5H), 3.56–3.62 (m, 2H), 3.70–3.78 (m, 3H), 3.81–3.88 (m, 3H), 4.46 (d, $J = 7.8$ Hz, 1H), 4.61 (d, $J = 7.8$ Hz, 1H), 5.31–5.39 (m, 2H). ^{13}C NMR (100 MHz, CDCl_3 - CD_3OD): δ 22.1 (q), 26.4 (t), 26.7 (t), 28.3 (t), 30.3 (t), 30.4 (t), 30.5 (t), 30.7 (t), 30.8 (t), 30.9 (t), 36.7 (t), 37.8 (t), 62.8 (t), 63.1 (t), 71.4 (d), 71.7 (d), 75.6 (d), 77.5 (d), 77.6 (d), 77.9 (d), 78.1 (d), 78.9 (d), 82.1 (d), 102.5 (d), 104.7 (d), 130.9 (2d), 179.9 (s). MS (ESI): $m/z = 645.47$ (100%, $[\text{M}+\text{Na}]^+$); 661.45 (52%, $[\text{M}+\text{K}]^+$). Anal. Calcd for $\text{C}_{30}\text{H}_{54}\text{O}_{13}$: C, 57.86; H, 8.74. Found: C, 57.60; H, 9.05.

1. a) *Preparation*: Zhou, S.; Xu, C.; Wang, J.; Gao, W.; Akhverdiyeva, R.; Shah, V.; Gross, R. *Langmuir*, **2004**, *20*, 7926.
b) *Hydrolysis*: Rau, U.; Heckmann, R.; Wray, V.; Lang, S. *Biotechnol. Lett.* **1999**, *21*, 973.
c) *Detailed spectral and analytical data*: Azim, A.; Shah, V.; Doncel, G. F.; Peterson, N.; Gao, W.; Gross, R. *Bioconj. Chem.* **2006**, *17*, 1523.

1H NMR OF SOPHOROLIPID

21 Sep 2007

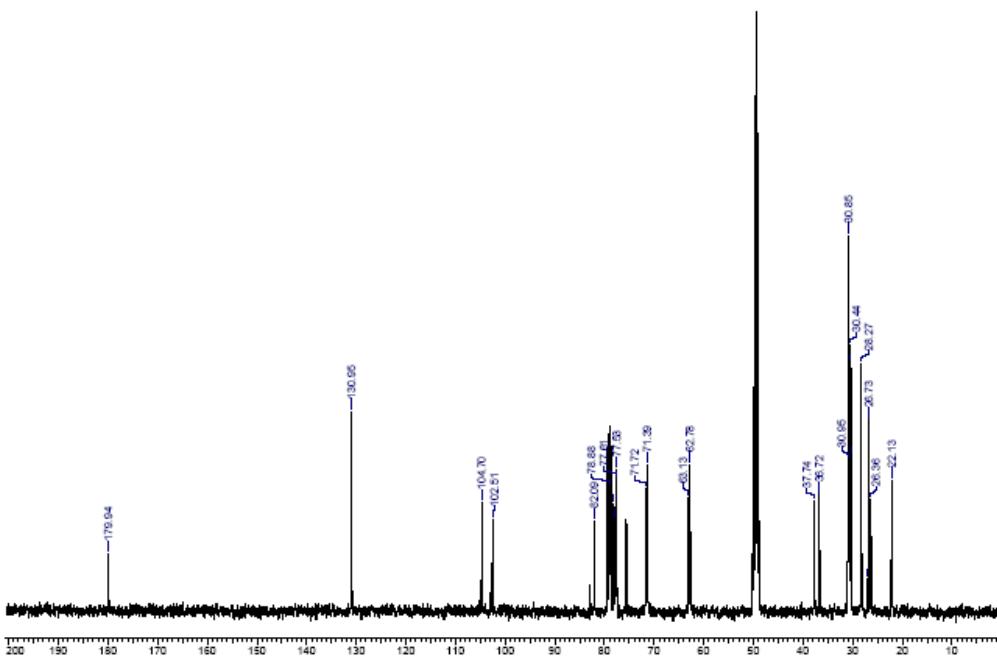
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13C NMR OF SOPHOROLIPID

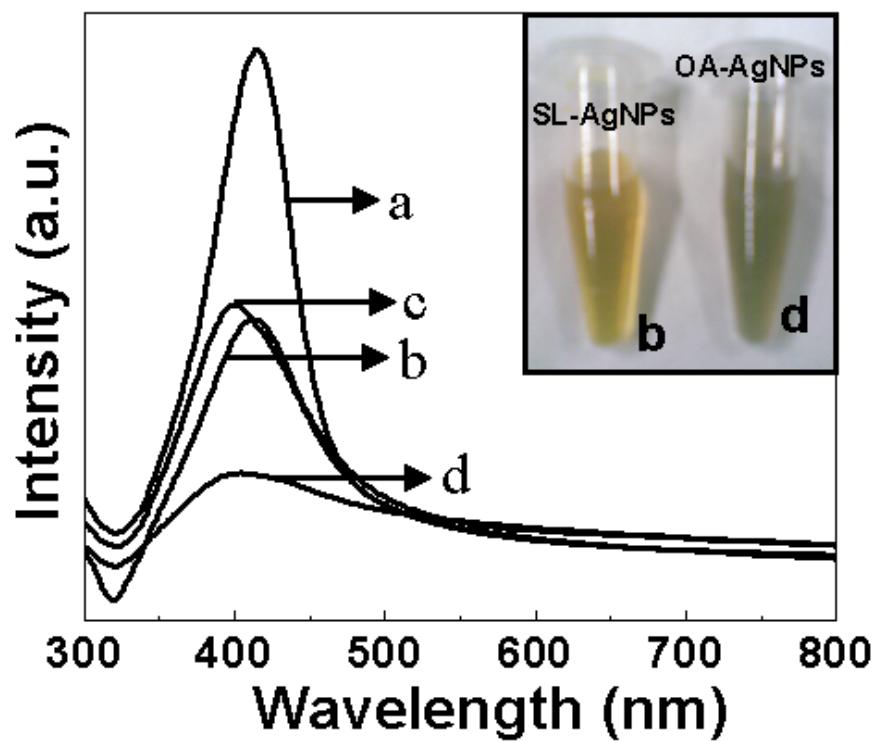
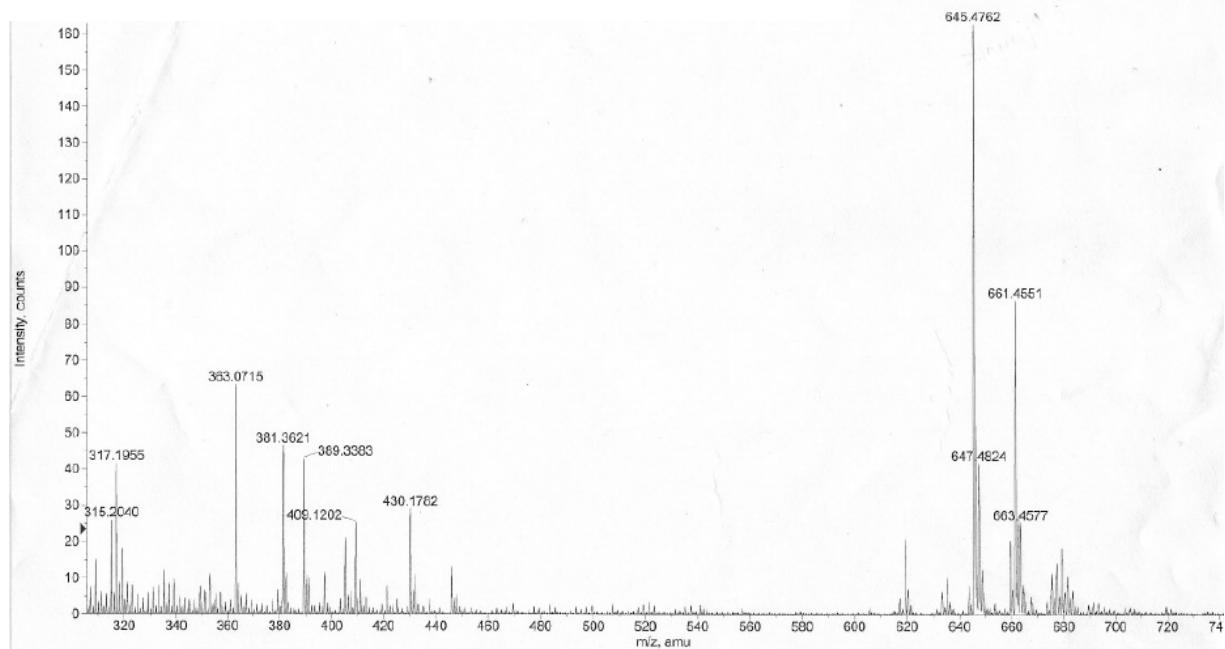
21 Sep 2007

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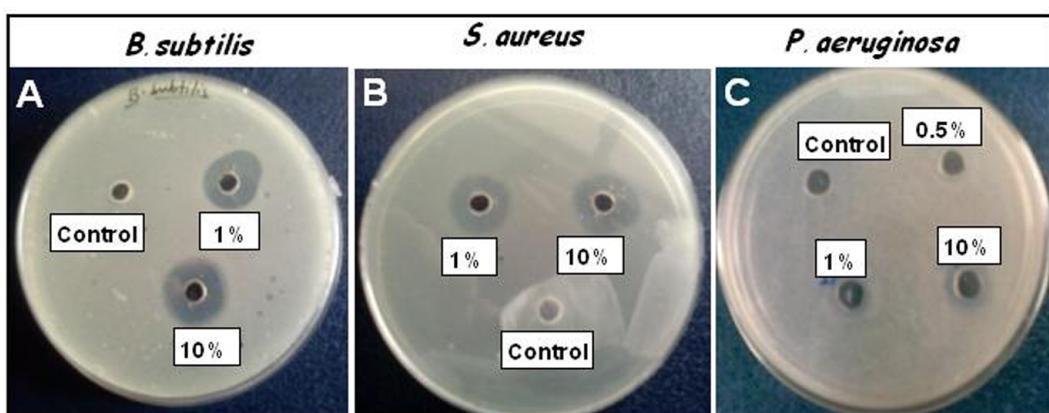


Mass spectra of acidic sophorolipid

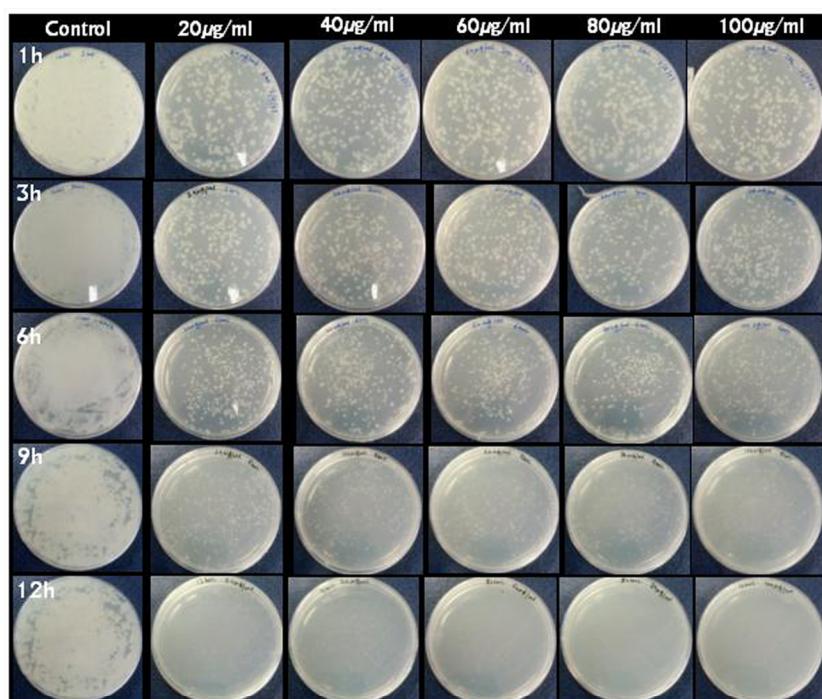
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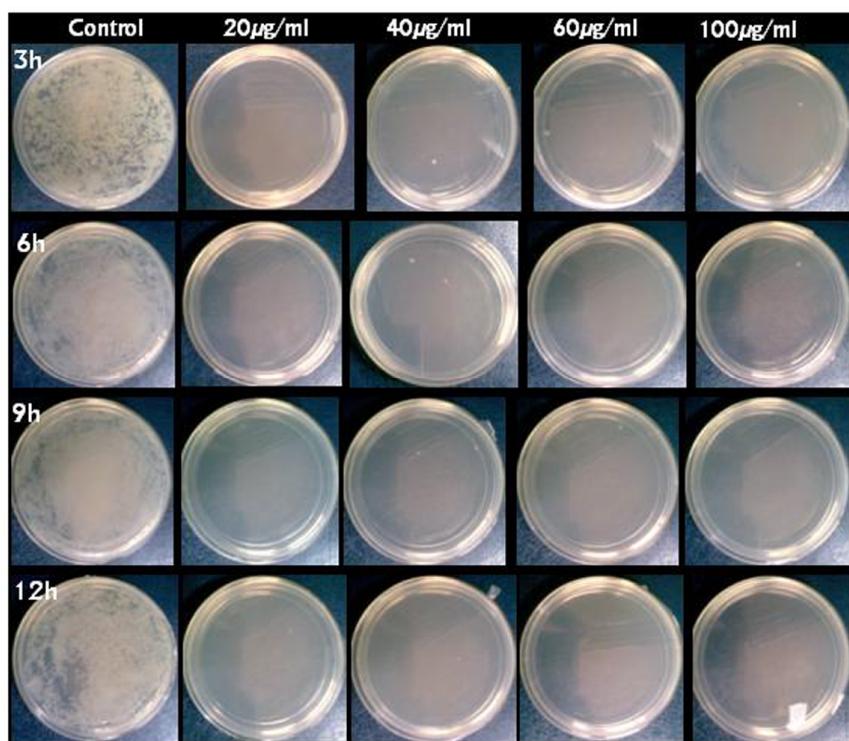
ESI-Fig-1: UV-vis spectra of SL-AgNPs (curve-a), OA-AgNPs (curve-b), redispersred SL-AgNPs (curve-c) and redispersred OA-AgNPs (curve-d). Inset shows the corresponding colour of redispersred SL-AgNPs (vial-c) and OA-AgNPs (vial-d).



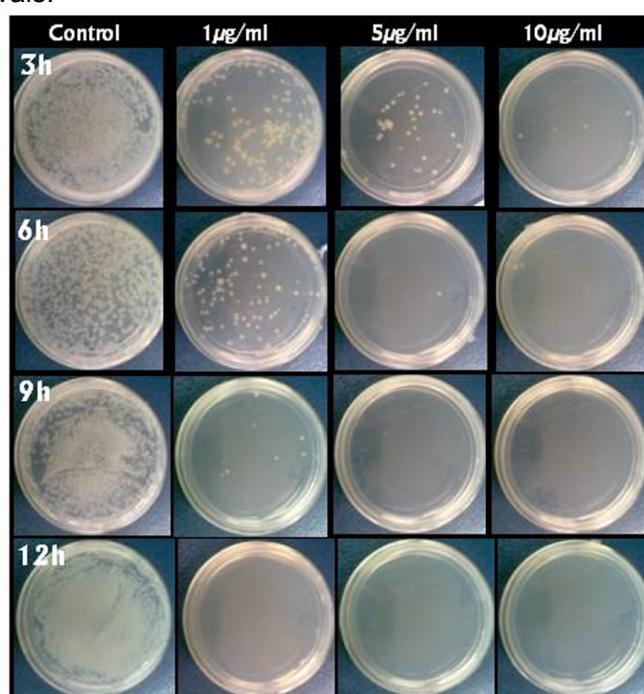
ESI-Fig-2: Luria-Agar plates showing antibacterial activity of different concentrations of pure sophorolipid against *B. subtilis* (plate-A), *S. aureus* (plate-B) and *P. aeruginosa* (plate-C).



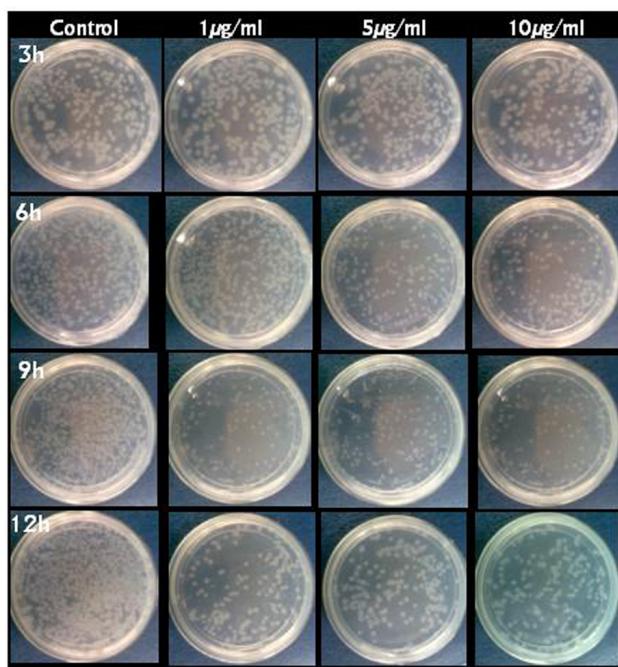
ESI-Fig-3: Luria-Agar plates showing antibacterial activity of different concentrations of SL-AgNPs (20 µg/mL, 40 µg/mL, 60 µg/mL, 80 µg/mL and 100 µg/mL) against *B. subtilis* at different time intervals.



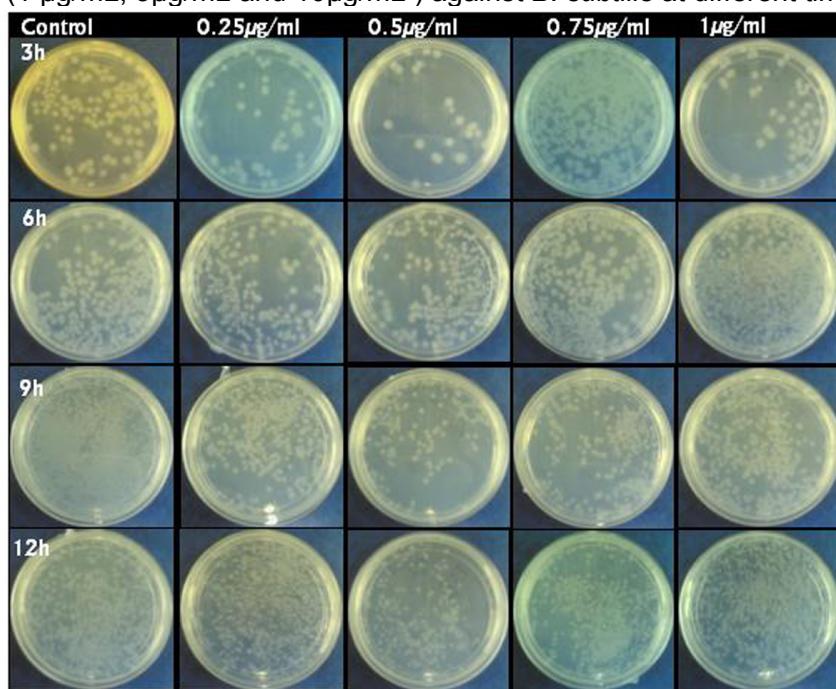
ESI-Fig-4: Luria-Agar plates showing antibacterial activity of different concentrations of SL-AgNPs (20 µg/mL, 40 µg/mL, 60 µg/mL and 100 µg/mL) against *P. aeruginosa* at different time intervals.



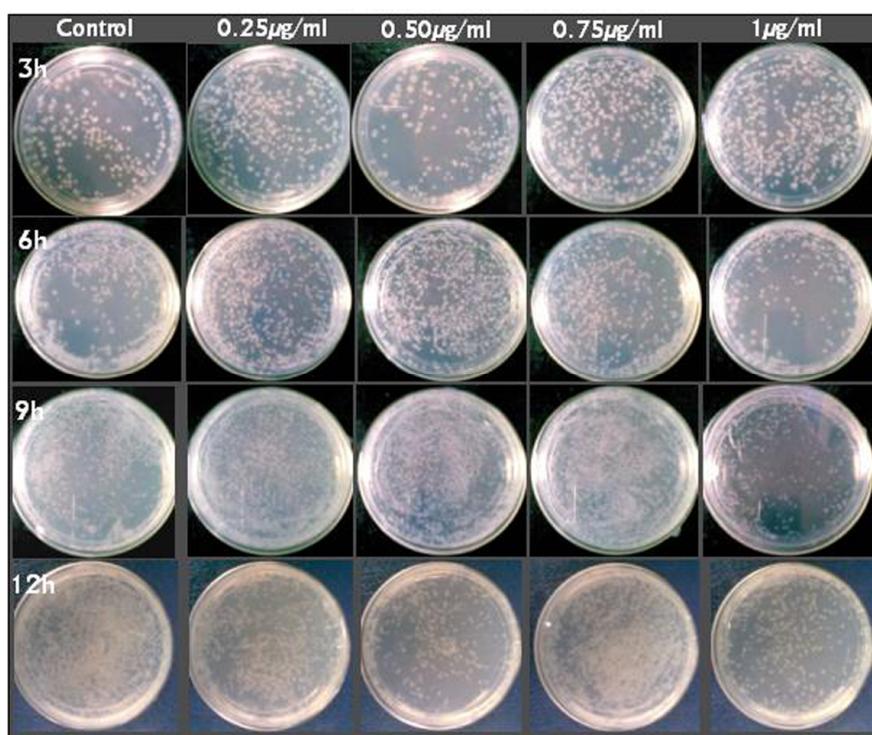
ESI-Fig-5: Luria-Agar plates showing antibacterial activity of different concentrations of SL-AgNPs (1 µg/mL, 5 µg/mL and 10 µg/mL) against *P. aeruginosa* at different time intervals.



ESI-Fig-6: Luria-Agar plates showing antibacterial activity of different concentrations of SL-AgNPs ($1 \mu\text{g}/\text{mL}$, $5\mu\text{g}/\text{mL}$ and $10\mu\text{g}/\text{mL}$) against *B. subtilis* at different time intervals.



ESI-Fig-7: Luria-Agar plates showing antibacterial activity of different concentrations of SL ($0.25\mu\text{g}/\text{mL}$, $0.50\mu\text{g}/\text{mL}$, $0.75\mu\text{g}/\text{mL}$ and $1.0\mu\text{g}/\text{mL}$) against *B. subtilis* at different time intervals.



ESI-Fig-8: Luria-Agar plates showing antibacterial activity of different concentrations of SL (0.25µg/mL, 0.50µg/mL, 0.75µg/mL and 1.0µg/mL) against *P. aeruginosa* at different time intervals.