

A aM]a`gabXA Yh`cXg

A. aYi []bcgW`hj ah]cb abXWYa]Vg

M. aeruginosa FACHB-915 was pre-cultured in sterile BG11 medium at $25 \pm 1^\circ\text{C}$ and a 12 h diurnal-nocturnal alternation with cool white lights at $25 \mu\text{mol photons/m}^2\cdot\text{s}$. 20 days pre-cultivation, *M. aeruginosa* cells reaching exponential growth phase were collected by centrifugation ($4000 \text{ g } 4^\circ\text{C}, 10 \text{ min}$), and used as inoculums for downstream experiments.

Kaempferol and luteolin were purchased from Energy Chemical, Inc. (Shanghai, China). Individual stock solution were prepared in dimethyl sulfoxide (DMSO) and stored at 4°C before use. Other chemicals used in this study were analytical grade except as stated by kit.

9l dY]a Yba`gyi d

All Erlenmeyer flasks were autoclaved at 121°C for 20 min before experimental use. For each exposure test group, individual 250 mL flask containing 100 mL sterile BG11 medium was spiked with certain dose of kaempferol or luteolin at test beginning without any replenishment along test period. Concentrations of kaempferol or luteolin were 0.5, 1, 4, 16 or 32 mg/L in each 100 mL medium, with DMSO amount below 0.1% (v/v).

the stimulated APC on day 4 and enhanced PE content along the test probably as adaptive strategy to 32 mg/L luteolin stress, the inhibited

was generally correlated to SOD/CAT activities at each kaempferol or luteolin level: i) Exposed to 16-32 mg/L kaempferol and 32 mg/L luteolin, MDA content on each day indicated *M. aeruginosa* cell damage, causing inhibited *M. aeruginosa* growth. Noteworthy, most SOD and CAT activity stimulation on day 8 could subsequently alleviate oxidative damage on day 14, according to decreasing MDA content from day 8 to 14 (down to 0.09, 0.13 and 0.12 nmol/10⁸ cells, respectively) (Figure 5). To some extent, such relatively-alleviated oxidative stress on day 14 further inactivated SOD/CAT, leading to lower SOD/CAT activities on day 14 (SOD down to 0.80, 0.98 and 1.49 U/10⁸ cells, CAT down to 5.67, 4.88, 3.38 U/10⁸ cells) (Figures 5A-5D).

ii) At 16 mg/L luteolin, SOD and CAT activities were absent on day 4 but both stimulated on day 8.

