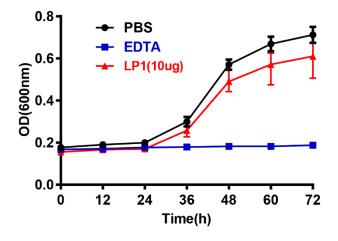


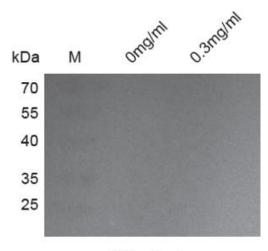
**Figure S1.** Recombinant 30K proteins were purified by NTA agarose affinity chromatography with a stepwise imidazole gradient. M: molecular weight marker; SP: soluble protein; FT: flow-through fraction.

## Figure S2



**Figure S2**. Antifungal activity of BmLP1 was performed by detecting the growth curve. 96-well plates were filled with  $300\mu$ L of potato liquid medium containing  $10\mu$ g BmLP1 and Candida albicans at a final concentration of 1 × 105 cells. 96-well plates were incubated at 30 °C, and fungal growth was observed

by monitoring the absorbance at 600 nm after culturing for 0, 12, 24, 36, and 48 h. Vertical bars represent the mean  $\pm$  SE (n = 3).



LP1 antibodies

**Figure S3**. Binding of BmLP1 to Hela cells revealed using western blotting. 0.3mg/ml concentrations of purified proteins in DMEM medium were incubated with Hela cells for 2 h at room temperature, washed four times with PBS after trypsin digestion, and Hela cell proteins were extracted using RIPA. BmLP1 was detected using BmLP1 specific polyclonal antibodies.

**Table S1.** Primers used in this study. F: Forward primer; R: Reverse primer; The BmlpX-F/R for protein expression; The BmlpX qPCR-F/R for RT-qPCR.

Primers	Sequence (5'-3')
Bmlp1-F	TACTTCCAATCCATGACACTTGCACCAAGAACTGA
Bmlp1-R	TATCCACCTTTACTGTTAGTAGGGGACGATGTACC
Bmlp2-F	TACTTCCAATCCATGGGCGTCGTGGAACTATCC
Bmlp2-R	TATCCACCTTTACTGTTAGAAAGGTGTAATGAACC
Bmlp3-F	TACTTCCAATCCATGGACGTCCCTAACGACATTCT
Bmlp3-R	TATCCACCTTTACTGTTAGAAAGCCTTTATACCCC
Bmlp4-F	TACTTCCAATCCATGGGCGTCACTGAAATGTCCG
Bmlp4-R	TATCCACCTTTACTGCTAGAAAGGTGTAACGAACC
Bmlp7-F	TACTTCCAATCCATGGCAGATTCCGACGTCCCT
Bmlp7-R	TATCCACCTTTACTGTTAAAATGCCTTAACACCCC
Bmlp1 qPCR-F	TGTTATCGAGCTTCAGGT
Bmlp1 qPCR-R	CCCCATCCAGTTTAGAG
Bmlp2 qPCR-F	AGCACCAGGGAGCAAT
Bmlp2 qPCR-R	TCACCGTCGTGTCCAA
Bmlp3 qPCR-F	GAGGAGAAGAAGAGCGAAGT
Bmlp3 qPCR-R	GCCCTGGAGCCAAAGT
Bmlp4 qPCR-F	ACACCACCAGGGAGCA
Bmlp4 qPCR-R	TGTAACGAACCAGGAGAAG
Bmlp7 qPCR-F	GGAGAAGAAGAGCGAAGT
Bmlp7 qPCR-R	TGGAGCCAAAGTTGATAG