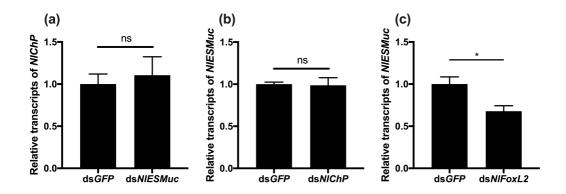
**Table S1**. Primers used in this work.

Primer usage	Primer name	Primer sequence (5'- 3')	Concentration
double stranded RNA synthesis	T7NlESMuc-F	TAATACGACTCACTATAGGGCAAACAGTAC AACTGCACCTCC	10mM
	T7NlESMuc-R	TAATACGACTCACTATAGGGGCTTGGCGAT ACTGTGCCAGG	10mM
	T7GFP-F	TAATACGACTCACTATAGGGAGAATGAGTA AAGGAGAAGAACTTTTC	10mM
	T7GFP-R	TAATACGACTCACTATAGGGAGATTTGTAT AGTTCATCCATGCCATG	10mM
quantitative real-time PCR	qNlESMuc-F	AGTGGCATTCGTTCTTAAAGG	10mM
	qNlESMuc-R	TGTGGTTGTGGCTTGTGG	10mM
	RPS11-F	CCGATCGTGTGGCGTTGAAGGG	10mM
	RPS11-R	ATGGCCGACATTCTTCCAGGTCC	10mM
	18S-F	GTAACCCGCTGAACCTCCT	10mM
	18S-R	TCCGAAGACCTCACTAAATC	10mM



**Figure S1**. (a) The expression level of a known eggshell protein gene NlChP in dsNlESMuc-injected female BPHs at 72 h post emergence. There was no significant difference. (b) The expression level of NlESMuc in dsNlChP-injected female BPHs at 72 h post emergence. There was no significant difference. (c) The expression level of NlESMuc after the RNAi of NlFoxL2, which regulates eggshell formation in BPHs, at 72 h post emergence. The mRNA level of NlESMuc decreased by 32.24% (p = 0.04).