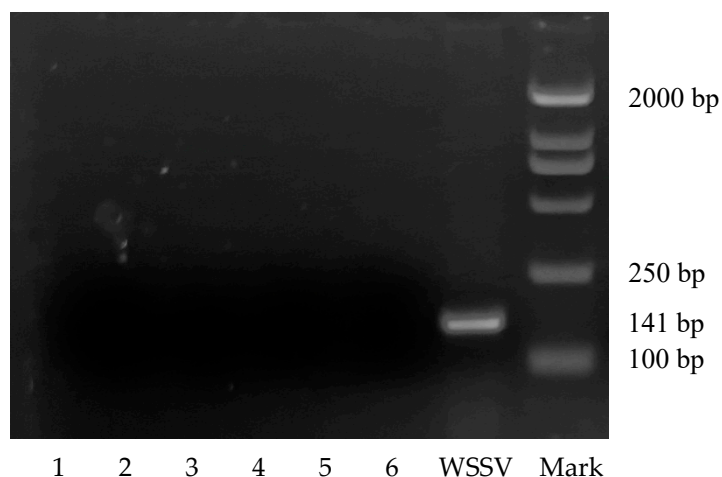


Table S1. Specific information on medicinal plants used in this study, including plant names, medicinal parts, extracting solvents, safe concentrations, experimental concentrations.

Plants	Parts	Solvent	Safe con. (mg/kg)	Experimental con. (mg/kg)
<i>Alpinia officinarum</i> Hance	Rhizome	Methanol	>150	100
<i>Amomum villosum</i> Lour.	Fruit	Methanol	>150	100
<i>Chelidonium majus</i> Linn.	Herb	Methanol	>150	100
<i>Fibraurea recisa</i> Pierre	Stem and leaves	Methanol	>150	100
<i>Iris dichotoma</i> Pall.	Herb and roots	Methanol	>150	100
<i>Leonurus japonicus</i> Houttuyn	Herb	Methanol	>80	50
<i>Linum usitatissimum</i> Linn.	Seeds	Methanol	>200	100
<i>Litsea mollis</i> Hemsl.	Fruit and roots	Methanol	>150	100
<i>Oxalis corniculata</i> Linn.	Herb	Methanol	>200	100
<i>Paederia foetida</i> Linn.	Herb and roots	Methanol	>150	100
<i>Pinellia ternata</i> (Thunb.) Breit.	Tubers	Methanol	>150	100
<i>Solanum nigrum</i> Linne.	Herb	Methanol	>80	50
<i>Humulus scandens</i> (Lour.) Merr.	Herb	Methanol	>150	100
<i>Urtica fissa</i> E. Pritz.	Leaves	Methanol	>150	100
<i>Xanthium strumarium</i> Linne.	Fruit	Methanol	>80	50

Figure S1. Six crayfish were randomly selected for PCR detection (VP28, 141 bp), and the WSSV virus solution was used as a positive control. The results showed that these crayfish were in normal state.



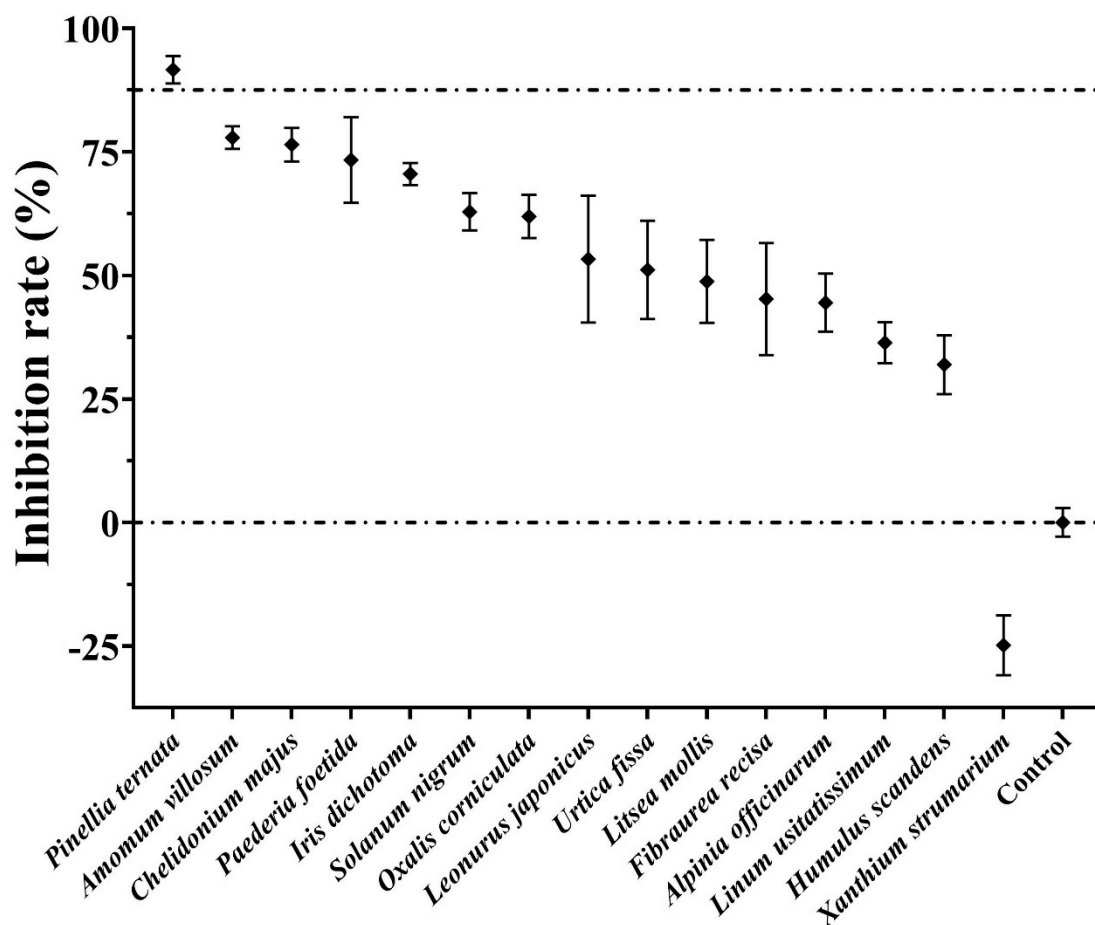


Figure S2. The inhibition rates of 15 different Chinese herbal extracts against WSSV were evaluated in vivo (100 mg/kg, 24hpi). The calculation formula was: inhibition rate = (control group - treatment group) / control group * 100 %. The data was shown as Mean \pm SD (n=5).

Figure S3. The schematic depicting the experimental procedures in Figures 1 and 2.

