

Supplemental reagents

Ethanol, trichloroacetic acid, anthracene, sulfuric acid, α -naphthol, iodine, potassium iodide, indamine, glycine, and sodium chloride were all analytically pure grades. Papain, Dextran T70, Dextran T40, Dextran T10, 4% paraformaldehyde, and Dextran Blue 2000 were purchased from Solarbio Technology Co., Ltd. (Beijing, China). Minimum Eagle's medium was purchased from Sigma-Aldrich Company (Darmstadt, Germany). Fetal bovine serum (FBS) was purchased from Biological Industries (Kibbutz Beit Haemek, Israel). The Thermo Scientific™ GeneJET Viral DNA/RNA Purification kit was purchased from Thermo Fisher Scientific (Waltham, MA, USA). The EZNA® Gel Extraction Kit, PCR purification kit, and Plasmid extraction kit were purchased from OMEGA (Connecticut, USA). The pMD™18-T Vector Cloning Kit and RNAiso Plus were purchased from Takara Bio-medical Technology Co., Ltd. (Beijing, China).

Table S1. Monosaccharide composition.

Name	CAS Number (Sigma)	Molecular For- mula	Content ($\mu\text{g}/\text{mg}$)
Fucose (Fuc)	2438-80-4	C ₆ H ₁₂ O ₅	4.4807
Arabinose (Ara)	5328-37-0	C ₅ H ₁₀ O ₅	1.3101
Rhamnose (Rha)	10030-85-0	C ₆ H ₁₄ O ₆	2.1440
Galactose (Gal)	26566-61-0	C ₆ H ₁₂ O ₆	4.5202
Glucose (Glc)	50-99-7	C ₆ H ₁₂ O ₆	9.4481
Xylose (Xyl)	58-86-6	C ₅ H ₁₀ O ₅	3.9688
Mannose (Man)	3458-28-4	C ₆ H ₁₄ O ₆	3.5177
Galacturonic Acid (Gal-UA)	14982-50-4	C ₆ H ₁₀ O ₇	7.3167
Guluronic Acid (Gul-UA)	15769-56-9	C ₆ H ₁₀ O ₇	59.4252
Glucuronic Acid (Glc-UA)	6556-12-3	C ₆ H ₁₀ O ₇	2.6896
Mannuronic Acid (Man-UA)	6814-36-4	C ₆ H ₁₀ O ₇	222.3577

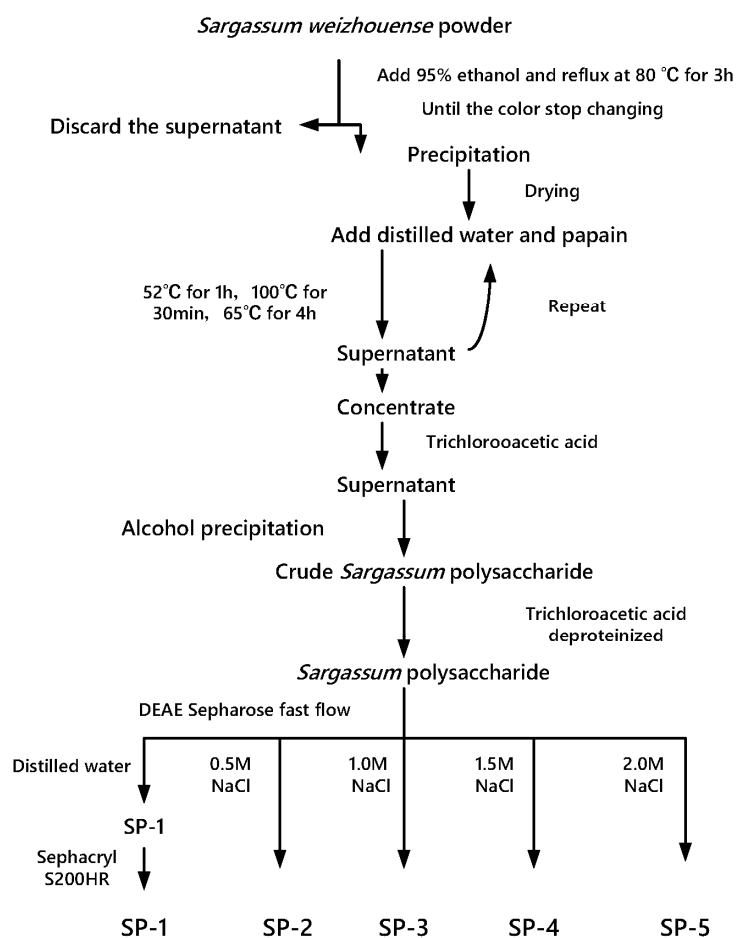


Figure S1. The process of extraction, isolation, and purification of *Sargassum weizhouense*.

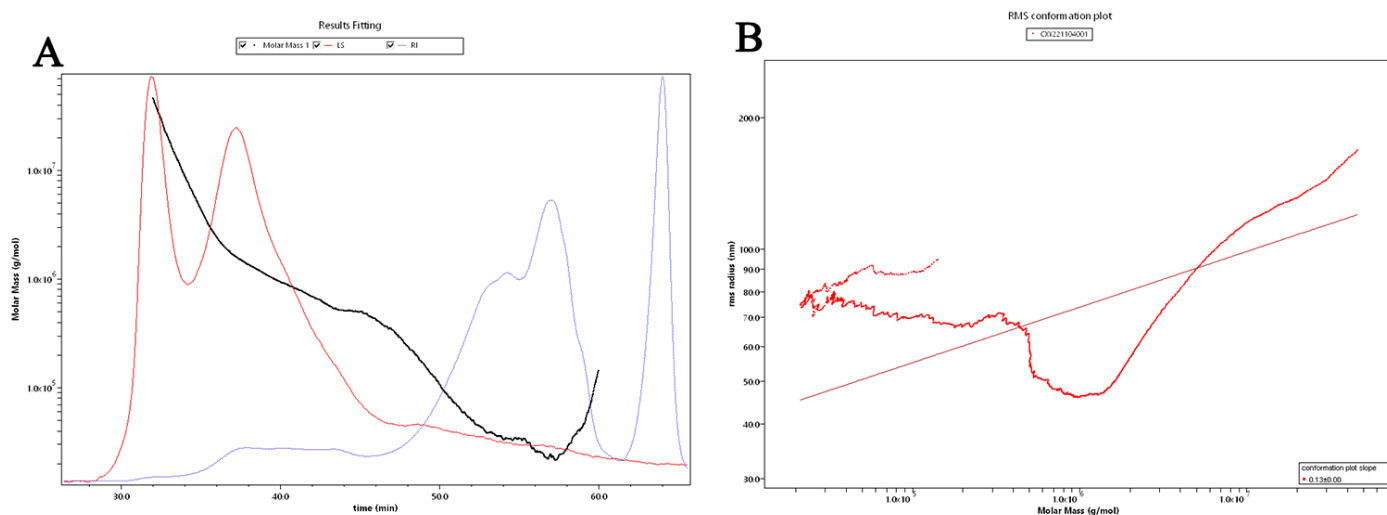


Figure S2. (A) Plot of absolute molecular weight analysis. Red line: the multi-angle laser light scattering signal (LS, unit: V). Blue line: the differential signal (RI, unit: RIU). Black line: the molecular weight fitted from the two signals. (B) The molecular configuration analysis of SP-1.

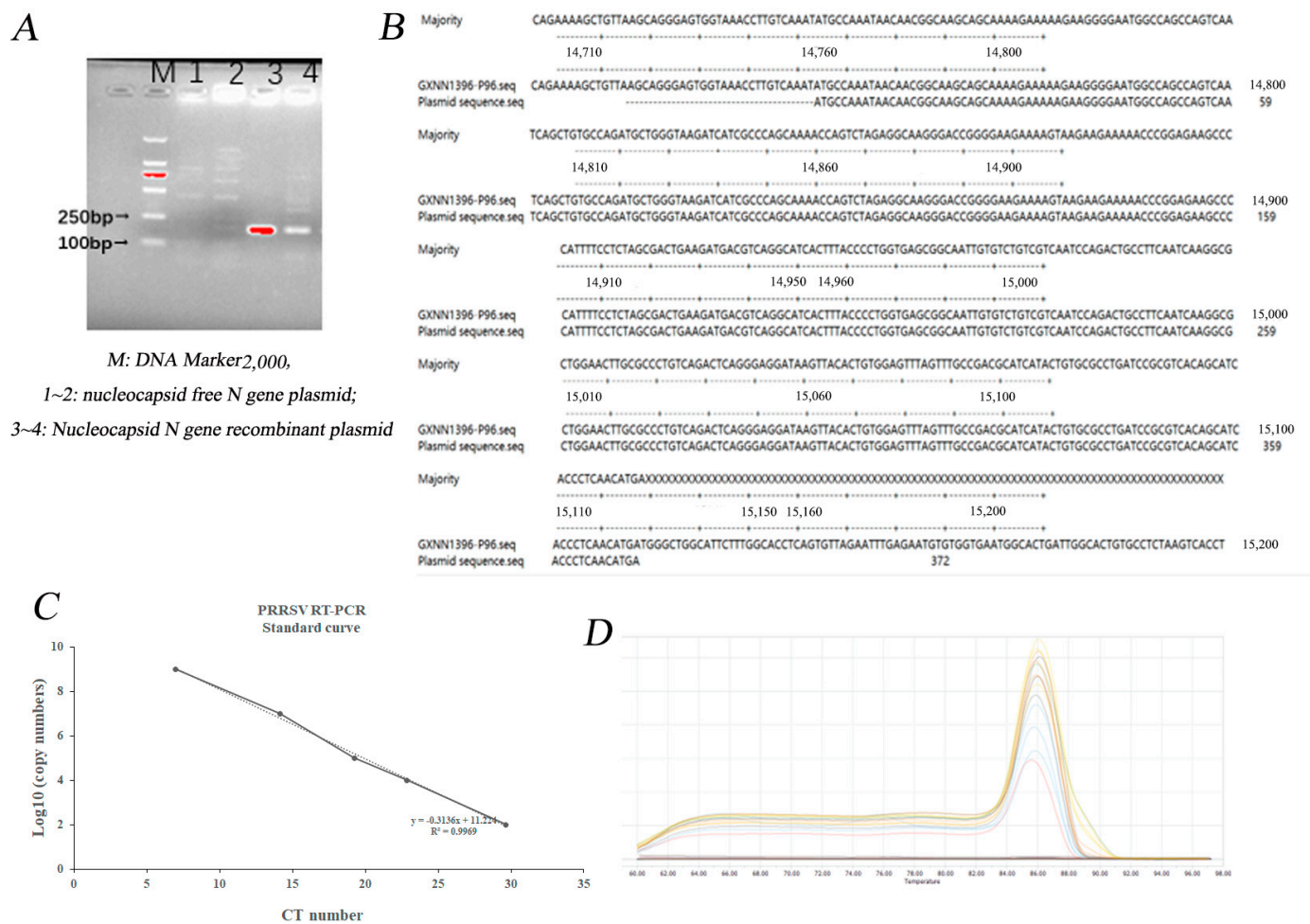


Figure S3. Construction of plasmid and establishment of qRT-PCR standard curve. (A) Gel electrophoresis detection of N gene recombinant plasmid. (B) Recombinant plasmid and PRRSV sequence alignment results. Note—the upper part is the sequence of plasmid N gene, whereas the lower part is PRRSV-GXNN1396. (C) Standard curve line of PRRSV qRT-PCR (CT-X). Note—The corresponding template concentrations from right to left in the above figure are: 8.396×10^9 , 8.396×10^7 , 8.396×10^5 , 8.396×10^4 , and 8.396×10^2 copies/ μ L. (D) Establishment of qRT-PCR melting curve.