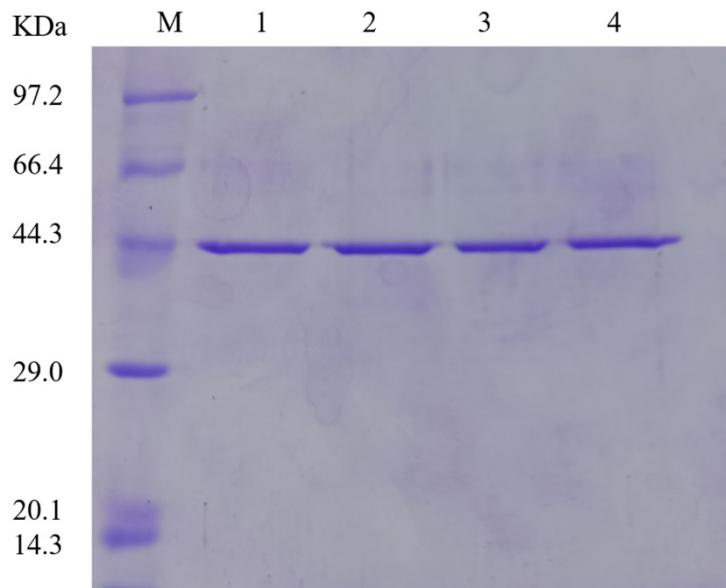


**Figure S1.** The state of the purified rAlys1 in solution. (a) The purified rAlys1 was gel-filtered through a Superdex peptide 200 10/300 column and monitored at a wavelength of 280 nm. (b) SDS-PAGE analysis of the state of the purified rAlys1. M, protein marker; column 1, the elution of second peak; column 2, the elution of first peak.



**Figure S2.** SDS-PAGE analysis of alginate lyases rAlys1 and its mutants.

M, protein marker; column 1, rAlys1; column 2, Q127A; column 3, H129A; column 4, Y279A.

**Table S1.** Primers used for mutations of rAlys1

| Mutagenesis | Primers Sequences   |
|-------------|---|
| Q127A       | 5'-CTTTTCAACCGTCGTGGAG <u>CAATT</u> CATAGTGATGAAGGAC-3'     |
|             | 5'-GTCCTTCATCACTATGAATT <u>GCT</u> CCAACGACGGTTGAAAAAG-3'   |
| H129A       | 5'-CCGTCGTTGGACAAATT <u>GCT</u> AGTGATGAAGGACACG-3'         |
|             | 5'-CGTGTCTTCATCACT <u>AGC</u> AATTGTCCAACGACGG-3'           |
| Y279A       | 5'-CAATATTTAACAAAGGGG <u>GCT</u> CCAACCAATCGAATGG-3'        |
|             | 5'-CCATTGATTGG <u>GTG</u> GCAGCCC <u>TTG</u> TTAAAATATTG-3' |