

Supplementary data

Table

Table 1: Designed specific primers of L-asparaginase genes from four strains of lactic acid bacteria.

Primers	Sequences
<i>St</i> -ASNase F	5' AGTAGTAC <u>CATATG</u> ATTAAAAAATCCTAG 3'
<i>St</i> -ASNase R	5' TTTCTCGAGCCCTTCAATATAATC 3'
<i>Lp</i> -ASNase F	5' GAGGAGGAGTAC <u>CATATG</u> AAGAAGATTTTGG 3'
<i>Lp</i> -ASNase R	5' CCACTCGAGTGAATTGGCAGTCGC 3'
<i>La</i> -ASNase F	5' GGAAAT <u>CATATG</u> AAGAAATTATTATTATG 3'
<i>La</i> -ASNase R	5' AAGCTCGAGTTTAAGAGTTACTTC 3'
<i>Ls</i> -ASNase F	5' GGA <u>CTGACAC</u> ATATGAAAAAATCCTCGTC 3'
<i>Ls</i> -ASNase R	5' TTTTAGCTCGAGTTTGAGACGGCGTTGC 3'

The underlines of sequences indicate the restriction sites, NdeI and XhoI, respectively.

Supplementary data

Figure

kDa 1 2 3

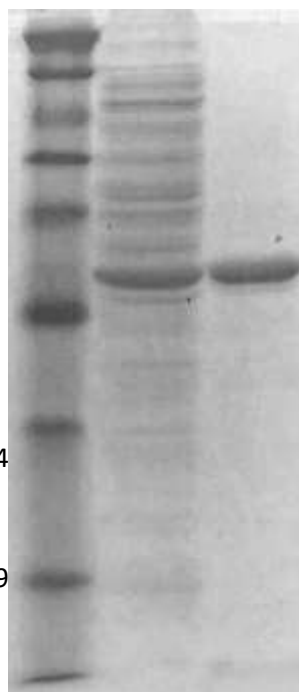
44.3

29.0

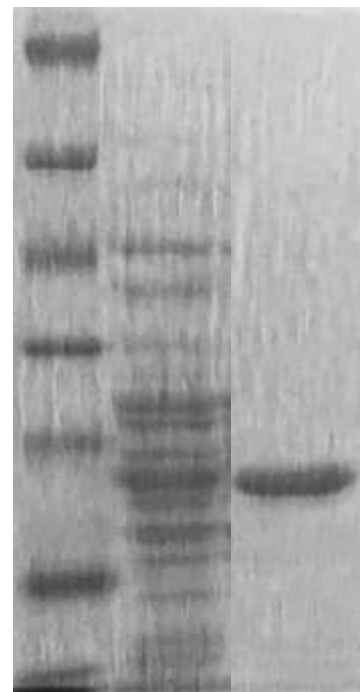
kDa 1 2 3

44

29



(A)



(B)

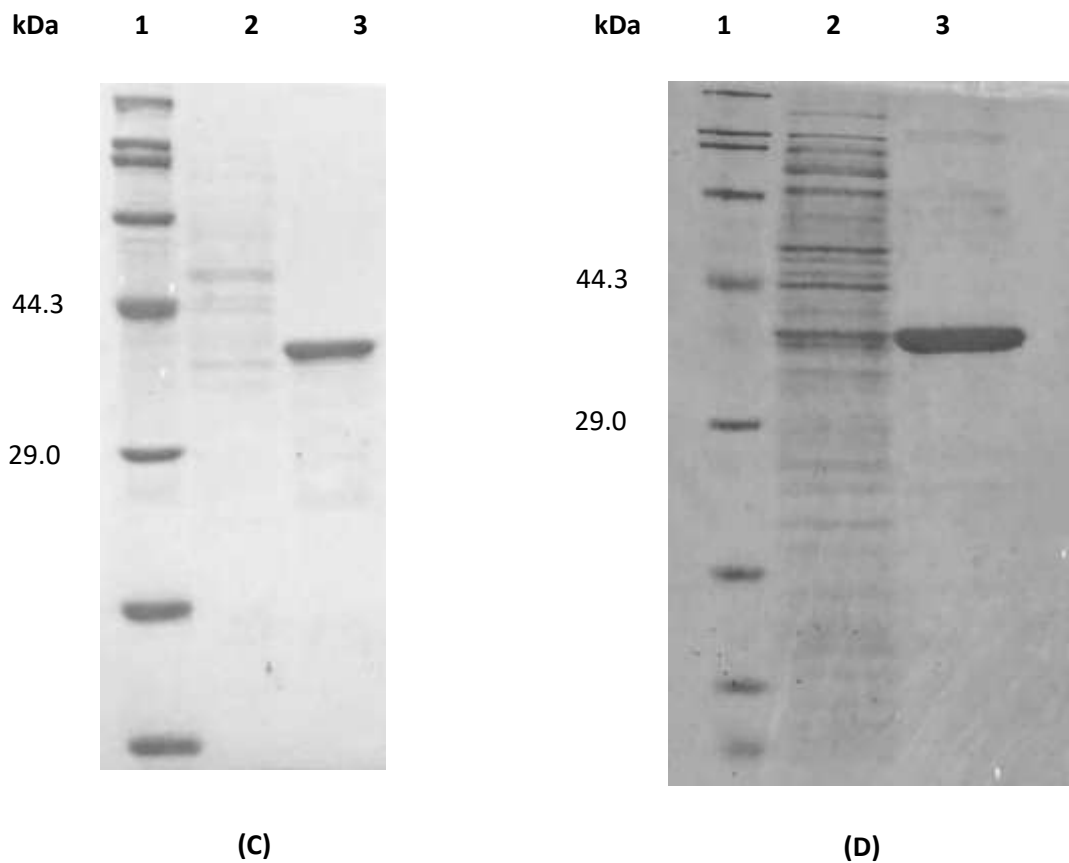


Figure 1: SDS-PAGE analyses of recombinant L-asparaginases from four strains of lactic acid bacteria

(A) *St*-ASNase, (B) *Lp*-ASNase, (C) *Ls*-ASNase and (D) *La*-ASNase; Lane1: Marker (Myosin, 200.0 kDa; β -galactosidase 116.0 kDa; Phosphorylase B 97.2kDa; Serum Albumin 66.4 kDa; Ovalbumin 44.3 kDa; Carbonic anhydrase 29.0 kDa; Trypsin Inhibitor 20.1 kDa; Lysozyme 14.3 kDa; Aprotinin 6.5 kDa), Lane2: Cell-free extract, Lane3: Purified recombinant L-asparaginase by using Ni SepharoseMT 6 Fast Flow column chromatography (pH 8.0)