

sgRNA design and *in vitro* nucleolytic analysis of the Cas9-RNP complex for transgene-free genome editing of the *eIF4E1* gene from *Capsicum annuum* L.

Josefanny Tham¹, Alfred Patisenah¹, Tommy Octavianus Soetrisno Tjia², Santiago Signorelli³, Intan Taufik¹, Karlia Meitha^{1*}

Supplementary Figure

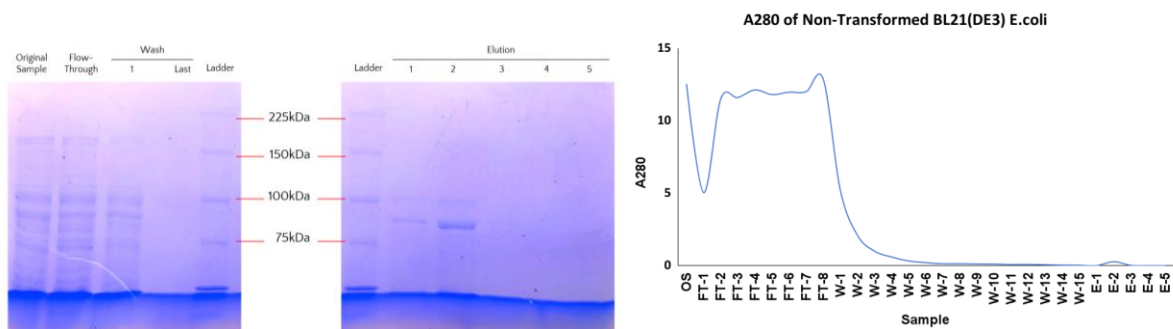


FIGURE S1 a SDS-PAGE of cytoplasmic fraction of non-transformed *E. coli* BL21(DE3) showed the presence background protein up to the second of eluted fraction (Broad range protein ladder (Thermo Fisher, USA); Separating 8%). **b** A₂₈₀ (mg/mL) chromatogram of cytoplasmic fraction of non-transformed *E. coli* BL21(DE3) purification process confirmed the presence of background protein (OS: Original sample, FT: Flow-through, W: Wash, E: Elution)