

High-level production of spider silk protein by fed-batch cultivation of recombinant *Escherichia coli* coexpressing *E. coli* dead gene and its purification

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Silk proteins from *Nephila clavipes* are fibrous proteins containing repetitive sequences with both crystalline and amorphous domains. The synthetic genes had contiguous units of the consensus repeat sequence of the silk protein were constructed in a tandem multimeric form. Silk proteins were expressed in *Escherichia coli* BL21(DE3) under the strong inducible T7 promoter. For efficient production of silk proteins with large multimers, coexpression of DeaD-box protein was also carried out to increase stability of long mRNA transcripts by synchronized transcription and translation. In order to obtain high-level production of silk protein in large amounts, pH-stat fed-batch cultures were carried out. Finally silk protein was simply purified using IMAC method.