Rapid purification system for His-tagged protein in a cell-free protein synthesis system

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We present a rapid protein expression and purification method in a cell-free system. Overall process is a single step that is isolated His-tagged proteins during cell-free synthesis from a batch reaction. However, a higher proportion of other proteins may also bind to immobilized metal-ion affinity chromatography (IMAC) resin. We described the use of pre-treated S30 extract with same resin and washing buffer with a low concentration of imidazole(30 mM) for the removal of contaminants. The results document that it should be possible to purify with a higher purity and to overcome the above problems. Our method may allow a faster and more convenient approach to study of High-throughput technology compared to classical in vivo system.