

**Crystallization and preliminary X-ray crystallographic studies of
Tpa1p from *Saccharomyces cerevisiae***

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The yeast Tpa1 (*Termination and PolyAdenylation*) protein contains a prolyl and lysyl hydroxylase domain. Tpa1p is ubiquitously distributed throughout the eukaryotic kingdom, from *C. elegans* to humans. Recently, it was proposed that Tpa1 is part of a messenger ribonucleoprotein (mRNP) complex at the 3'-UTR of mRNAs, associates specifically with components of the translation termination complex, and is involved in both translation termination and regulation of normal mRNA decay through translation termination-coupled poly(A) shortening. To gain insight into the Tpa1 function at the molecular level, we have overexpressed and crystallized Tpa1p. X-ray diffraction data were collected to 2.5 Å using synchrotron X-rays. The native crystal belongs to the monoclinic space group C2, with unit cell parameters of $a = 236 \text{ Å}$, $b = 136 \text{ Å}$, $c = 83 \text{ Å}$, and $\beta = 90.0$. The asymmetric unit contains three monomers, giving a crystal volume per protein mass (V_m) of $3.0 \text{ Å}^3 \text{ Da}^{-1}$ and a solvent content of 59%. Single wavelength anomalous diffraction data were collected at 100 K from a crystal of the SeMet-substituted protein. Model building is in progress.