

# K29 Tetra-Ubiquitin

Cat. No. SBB-UP0078  
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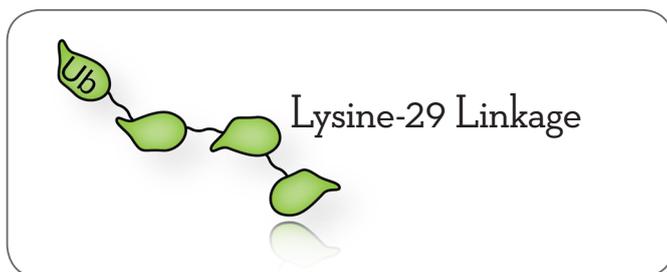


# South Bay Bio

## K29 Tetra-Ubiquitin

The array of cellular processes initiated and regulated by ubiquitin has been partially explained by the structural diversity of differently linked ubiquitin chains. In a ubiquitin chain, ubiquitin moieties can be conjugated through one of their lysine residues (K6, K11, K27, K29, K33, K48 and K63) or the N-terminal methionine residue (M1), offering countless possibilities to assemble a specific polymer. Ubiquitin molecules can also be modified by other post-translational modifications, including acetylation and phosphorylation, adding another layer of ubiquitin signal regulation and diversification.

K29 linked polyubiquitination has been shown to be an inhibitor of Wnt signaling, which plays an important role in embryogenesis, and its deregulation has been shown to play a role in tumorigenesis. This K29 linked tetra-ubiquitin was enzymatically conjugated, and purified via liquid chromatography.



## References

- 1) Dikic, I., Wakatsuki, S., & Walters, K. J. (2009). Ubiquitin-binding domains – from structures to functions. *Nature Reviews Molecular Cell Biology*, 10(10), 659-671. <https://doi.org/10.1038/nrm2767>
- 2) Akutsu, M., Dikic, I., & Bremm, A. (2016). Ubiquitin chain diversity at a glance. *Journal of Cell Science*, 129(5), 875-880. <https://doi.org/10.1242/jcs.183954>

## Product Information

**Quantity:** 25 µg      **Molecular Weight:** 34 kDa

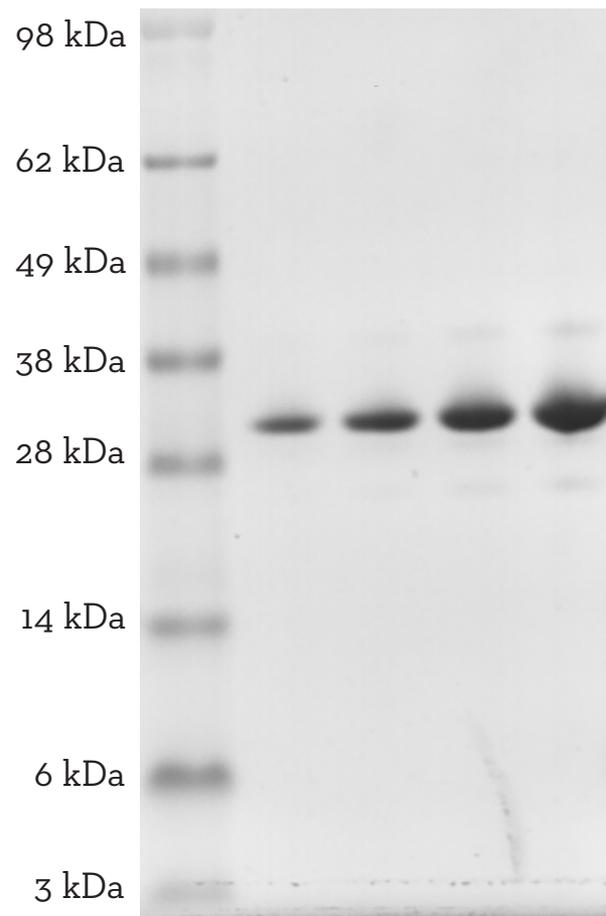
**Concentration:** 29 µM, 1 mg/mL

**Purity:** >95% by SDS-PAGE

**Storage Buffer:** 50 mM HEPES pH 7.5

**Storage:** -80C, Avoid multiple freeze / thaw

## Quality Control and Performance Data



**K29 linked Tetra-Ubiquitin SDS-PAGE.** From left to right, increasing amounts of tetra-ubiquitin were loaded onto a 10-20% SDS-PAGE gel, stained with Coomassie brilliant blue. Purity is > 95%.

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