Isolation, Characterization, and Imaging Analysis of Human Mediator Complexes Using HaloTag Technology.

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Abstract and introduction

The understanding of protein complex assembly and mapping of protein interactions has rapidly grown in recent years due to significant advances in mass spectrometry. As experiments turn toward characterization of the human proteome and the complexity it presents, a need remains to efficiently capture complexes intact, particularly weak or transient interactors as well as large multiprotein complexes. Here we present a new technology based upon the use of a protein fusion tag, termed HaloTag, which allows for highly specific and covalent immobilization of proteins complexes, as well as the ability to do correlative cellular localization studies using fluorescent ligands. In studies presented here, this technology was used to further understanding of the multi-protein Mediator complex assemblies and their cellular function.

HaloTag technology

HaloTag (HT) is a genetically encoded 33kDa protein fusion tag.
Engineered to covalently bind various ligands, imparting multi-functionality.

Mediator Transcriptional Co-activator and Design

Mediator exists in two forms: Core Mediator and Mediator+Kinase Module
Place HaloTag on Core component MED26, and Kinase Module component, MED13

Expression and pulldown
Halo-Mediator fusions in HEK293

Mediator isolation using MED26-Halo and Halo-MED13

More Spectrometry Analysis
- Run pulldown samples on gel
- Excise into 10 segments per sample
- Trypsin digestion
- Nano LC-MS/MS (LTQ Orbitrap Velos)
- Database search (Mascot)
- Database processing (Scaffold)
- Data Analysis (Spectral Counting and Normalized Spectral Abundance Factors (NSAFs))

- Significant enrichment over control of proteins from both HaloTag constructs.
- Successful expression and capture of MED13 (239kDa) as a HaloTag fusion.

NSAF plot of Mediator subunits from MED26 Isolations

Excellent enrichment and capture of core Mediator subunits in biological triplicates.
No Mediator subunits identified in HaloTag alone control.

NSAF plot of Mediator subunits from MED13 Isolations

Excellent enrichment and capture of core Mediator subunits is biological duplicates.
Additional capture of CDK9 Module subunits (CDK8, Cyclin C, and MED12)

Summary

HaloTag technology allows for broad study of protein function both in vitro and in vivo, including complex isolation and cellular imaging.
Rapid and covalent isolation of complexes promotes maintenance of complex integrity, particularly macromolecular complexes.
Efficient isolation of core Mediator and Mediator bound to it is kinase module using MED26-HaloTag and Halo-MED13, respectively.
Mass spectrometry analysis showed distinct complexes associated with different Mediator subunits.
Cellular imaging revealed discrete nuclear localization of MED26 and MED13.

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